1aoz
Evolutionary trace report by report_maker
June 30, 2010

CONTENTS

1 Introduction

2 Chain 1aozA
2.1 P37064 overview
2.2 Multiple sequence alignment for 1aozA
2.3 Residue ranking in 1aozA
2.4 Top ranking residues in 1aozA and their position on the structure
   2.4.1 Clustering of residues at 25% coverage.
   2.4.2 Overlap with known functional surfaces at 25% coverage.
   2.4.3 Possible novel functional surfaces at 25% coverage.

3 Notes on using trace results
3.1 Coverage
3.2 Known substitutions
3.3 Surface
3.4 Number of contacts
3.5 Annotation
3.6 Mutation suggestions

4 Appendix
4.1 File formats
4.2 Color schemes used
4.3 Credits

1 INTRODUCTION

From the original Protein Data Bank entry (PDB id 1aoz):
Title: Refined crystal structure of ascorbate oxidase at 1.9 angstroms resolution
Compound: Mol id: 1; molecule: ascorbate oxidase; chain: a, b; ec: 1.10.3.3; engineered: yes
Organism, scientific name: Cucurbita Pepo Var. Melopepo
1aoz contains a single unique chain 1aozA (552 residues long) and its homologue 1aozB.

2 CHAIN 1AOZA

2.1 P37064 overview

From SwissProt, id P37064, 100% identical to 1aozA:
Description: L-ascorbate oxidase (EC 1.10.3.3) (Ascorbase) (ASO).
Organism, scientific name: Cucurbita pepo var. melopepo (Zucchini).
Taxonomy: Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons; rosids; eudicotyledon
Function: May be involved in a redox system involving ascorbic acid.
Catalytic activity: 2 L-ascorbate + O(2) = 2 dehydroascorbate + 2 H(2)O.
Cofactor: This protein belongs to the multicopper oxidases which contain three distinct Cu centers known as type 1 or blue, type 2 or normal, and type 3 or coupled binuclear.
Similarity: Belongs to the multicopper oxidase family.
Similarity: Contains 3 plastocyanin-like domains.
About: This Swiss-Prot entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use as long as its content is in no way modified and this statement is not removed.
2.2 Multiple sequence alignment for 1aozA

For the chain 1aozA, the alignment 1aozA.msf (attached) with 494 sequences was used. The alignment was downloaded from the HSSP database, and fragments shorter than 75% of the query as well as duplicate sequences were removed. It can be found in the attachment to this report, under the name of 1aozA.msf. Its statistics, from the alistat program are the following:

- Format: MSF
- Number of sequences: 494
- Total number of residues: 230661
- Smallest: 208
- Largest: 552
- Average length: 466.9
- Alignment length: 552
- Average identity: 29%
- Most related pair: 99%
- Most unrelated pair: 0%
- Most distant seq: 30%

Furthermore, <1% of residues show as conserved in this alignment.

The alignment consists of 33% eukaryotic (1% arthropoda, 22% fungi, 10% plantae), and <1% prokaryotic sequences. (Descriptions of some sequences were not readily available.) The file containing the sequence descriptions can be found in the attachment, under the name 1aozA.descr.

2.3 Residue ranking in 1aozA

The 1aozA sequence is shown in Figs. 1–2, with each residue colored according to its estimated importance. The full listing of residues in 1aozA can be found in the file called 1aozA.ranks_sorted in the attachment.

2.4 Top ranking residues in 1aozA and their position on the structure

In the following we consider residues ranking among top 25% of residues in the protein. Figure 3 shows residues in 1aozA colored by their importance: bright red and yellow indicate more conserved/important residues (see Appendix for the coloring scheme). A Pymol script for producing this figure can be found in the attachment.

2.4.1 Clustering of residues at 25% coverage.

Fig. 4 shows the top 25% of all residues, this time colored according to clusters they belong to. The clusters in Fig. 4 are composed of the residues listed in Table 1.
Fig. 4. Residues in 1a0zA, colored according to the cluster they belong to: red, followed by blue and yellow are the largest clusters (see Appendix for the coloring scheme). Clockwise: front, back, top and bottom views. The corresponding Pymol script is attached.

Table 1. Clusters of top ranking residues in 1a0zA.

<table>
<thead>
<tr>
<th>cluster color</th>
<th>size</th>
<th>member residues</th>
</tr>
</thead>
<tbody>
<tr>
<td>red</td>
<td>133</td>
<td>17,18,19,32,33,34,48,50,52</td>
</tr>
<tr>
<td></td>
<td></td>
<td>58,59,60,61,62,63,66,71,72</td>
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<td></td>
<td></td>
<td>73,74,75,76,78,79,80,81,82</td>
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<td></td>
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<td>83,85,86,89,91,93,98,99,100</td>
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<td></td>
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<td>101,102,103,104,105,106,110</td>
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<td></td>
<td>111,113,114,115,116,117,120</td>
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<td>121,123,133,141,143,144,173</td>
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<td></td>
<td>174,208,213,214,215,216,217</td>
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<td>220,222,224,225,226,227,228</td>
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<tr>
<td></td>
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<td>229,231,232,234,238,239,240</td>
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<td>451,452,454,456,458,460,462</td>
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<td></td>
<td></td>
<td>464,477,480,481,482,485,489</td>
</tr>
<tr>
<td></td>
<td></td>
<td>490,492,494,495,497,498,499</td>
</tr>
<tr>
<td></td>
<td></td>
<td>500,501,502,503,504,506,507</td>
</tr>
<tr>
<td></td>
<td></td>
<td>508,509,510,512,516,517</td>
</tr>
<tr>
<td>blue</td>
<td>2</td>
<td>41,42</td>
</tr>
<tr>
<td>yellow</td>
<td>2</td>
<td>28,29</td>
</tr>
</tbody>
</table>

Table 2. The top 25% of residues in 1a0zA at the interface with NAG. (Field names: res: residue number in the PDB entry; type: amino acid type; subst’s: substitutions seen in the alignment; with the percentage of each type in the bracket; noc/bb: number of contacts with the ligand, with the number of contacts realized through backbone atoms given in the bracket; dist: distance of closest approach to the ligand.)

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>subst’s (%)</th>
<th>cvg</th>
<th>noc/bb</th>
<th>dist (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>71</td>
<td>W</td>
<td>W (65) Y (6) H (3) Q (4) E (3) DM F (2) SGV A (1) LI</td>
<td>0.19</td>
<td>13/0</td>
<td>3.42</td>
</tr>
</tbody>
</table>

Table 3. List of disruptive mutations for the top 25% of residues in 1a0zA, that are at the interface with NAG.

Figure 5 shows residues in 1a0zA colored by their importance, at the interface with 1aozANAG601.

Cu-o-cu linkage binding site. Table 4 lists the top 25% of residues at the interface with 1aozAC20702 (cu-o-cu linkage). The following table (Table 5) suggests possible disruptive replacements for these residues (see Section 3.6).

Table 4. The top 25% of residues at the interface with 1aozAC20702 (cu-o-cu linkage).

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>subst’s (%)</th>
<th>cvg</th>
<th>noc/bb</th>
<th>dist (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>106</td>
<td>H</td>
<td>H (90) S (5) . (2) KIP AMQT</td>
<td>0.01</td>
<td>10/0</td>
<td>2.16</td>
</tr>
<tr>
<td>506</td>
<td>H</td>
<td>H (89) R (6) . (3) GF</td>
<td>0.01</td>
<td>11/0</td>
<td>2.07</td>
</tr>
<tr>
<td>104</td>
<td>H</td>
<td>H (89) F (6) . (2) NVG WKQAS</td>
<td>0.02</td>
<td>12/0</td>
<td>2.19</td>
</tr>
<tr>
<td>448</td>
<td>H</td>
<td>H (95) . (3) TEY</td>
<td>0.02</td>
<td>11/0</td>
<td>3.57</td>
</tr>
</tbody>
</table>

2.4.2 Overlap with known functional surfaces at 25% coverage.

The name of the ligand is composed of the source PDB identifier and the heteroatom name used in that file.
Fig. 5. Residues in 1aozA, at the interface with NAG, colored by their relative importance. The ligand (NAG) is colored green. Atoms further than 30 Å away from the geometric center of the ligand, as well as on the line of sight to the ligand were removed. (See Appendix for the coloring scheme for the protein chain 1aozA.)

Table 4. The top 25% of residues in 1aozA at the interface with cu-o-cu linkage. (Field names: res: residue number in the PDB entry; type: amino acid type; subst's: substitutions seen in the alignment; with the percentage of each type in the bracket; noc/bb: number of contacts with the ligand, with the number of contacts realized through backbone atoms given in the bracket; dist: distance of closest approach to the ligand.)

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>subst's (%)</th>
<th>cvg</th>
<th>noc/bb</th>
<th>dist (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>450</td>
<td>H</td>
<td>H(89)</td>
<td>0.03</td>
<td>8/0</td>
<td>2.06</td>
</tr>
<tr>
<td></td>
<td></td>
<td>D(6)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>.(3)TYA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>SRN</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>60</td>
<td>H</td>
<td>H(88)</td>
<td>0.04</td>
<td>8/0</td>
<td>4.13</td>
</tr>
<tr>
<td></td>
<td></td>
<td>T(4)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>.(4)C</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>S(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>62</td>
<td>H</td>
<td>H(89)</td>
<td>0.04</td>
<td>10/1</td>
<td>1.98</td>
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<td>N(4)</td>
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<td>.(3)</td>
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<td>12/0</td>
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<td>A(1)</td>
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<td></td>
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</tr>
<tr>
<td></td>
<td></td>
<td>E(4)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>.(3)P</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Q(1)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>102</td>
<td>F</td>
<td>W(68)</td>
<td>0.10</td>
<td>2/0</td>
<td>4.83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>F(19)S</td>
<td></td>
<td></td>
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</tr>
<tr>
<td></td>
<td></td>
<td>.(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>M(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>L(5)ACY</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>V(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 5. List of disruptive mutations for the top 25% of residues in 1aozA, that are at the interface with cu-o-cu linkage.

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>disruptive mutations</th>
</tr>
</thead>
<tbody>
<tr>
<td>106</td>
<td>H</td>
<td>(E) (T) (D) (Q)</td>
</tr>
<tr>
<td>506</td>
<td>H</td>
<td>(E) (D) (TQM) (SNLPI)</td>
</tr>
<tr>
<td>104</td>
<td>H</td>
<td>(E) (D) (T) (Q)</td>
</tr>
<tr>
<td>448</td>
<td>H</td>
<td>(Q) (M) (E) (K)</td>
</tr>
<tr>
<td>450</td>
<td>H</td>
<td>(E) (M) (Q) (D)</td>
</tr>
<tr>
<td>60</td>
<td>H</td>
<td>(E) (Q) (M) (K)</td>
</tr>
<tr>
<td>62</td>
<td>H</td>
<td>(E) (T) (QM) (CDG)</td>
</tr>
<tr>
<td>508</td>
<td>H</td>
<td>(TE) (D) (CG) (Q)</td>
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<tr>
<td>102</td>
<td>F</td>
<td>(K) (E) (Q) (R)</td>
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</table>

Table 5. Continued

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<tr>
<th>res</th>
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<th>subst's (%)</th>
<th>cvg</th>
<th>noc/bb</th>
<th>dist (Å)</th>
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</thead>
<tbody>
<tr>
<td>106</td>
<td>H</td>
<td>H(89)</td>
<td>0.03</td>
<td>8/0</td>
<td>2.06</td>
</tr>
<tr>
<td></td>
<td></td>
<td>D(6)</td>
<td></td>
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</tr>
<tr>
<td></td>
<td></td>
<td>.(3)TYA</td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>SRN</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>506</td>
<td>H</td>
<td>H(88)</td>
<td>0.04</td>
<td>8/0</td>
<td>4.13</td>
</tr>
<tr>
<td></td>
<td></td>
<td>T(4)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>.(4)C</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>S(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>104</td>
<td>H</td>
<td>H(89)</td>
<td>0.04</td>
<td>10/1</td>
<td>1.98</td>
</tr>
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<td></td>
<td></td>
<td>N(4)</td>
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<td>.(3)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
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<td></td>
<td>D(1)SP</td>
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<tr>
<td>448</td>
<td>H</td>
<td>H(89)</td>
<td>0.03</td>
<td>12/0</td>
<td>2.14</td>
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<td>A(1)</td>
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<td>E(4)</td>
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<td>.(3)P</td>
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<tr>
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<td>Q(1)</td>
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</tr>
<tr>
<td>450</td>
<td>H</td>
<td>W(68)</td>
<td>0.10</td>
<td>2/0</td>
<td>4.83</td>
</tr>
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<td>F(19)S</td>
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<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>.(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>M(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
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<td>L(5)ACY</td>
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<td></td>
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</tr>
<tr>
<td></td>
<td></td>
<td>V(2)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 6. Residues in 1aozA, at the interface with cu-o-cu linkage, colored by their relative importance. The ligand (cu-o-cu linkage) is colored green. Atoms further than 30 Å away from the geometric center of the ligand, as well as on the line of sight to the ligand were removed. (See Appendix for the coloring scheme for the protein chain 1aozA.)

Figure 6 shows residues in 1aozA colored by their importance, at the interface with 1aozAC2O702.

Copper (ii) ion binding site. Table 6 lists the top 25% of residues at the interface with 1aozACU701 (copper (ii) ion). The following table (Table 7) suggests possible disruptive replacements for these residues (see Section 3.6).
Table 6. The top 25% of residues in 1aozA at the interface with copper (ii) ion. (Field names: res: residue number in the PDB entry; type: amino acid type; subst’s: substitutions seen in the alignment; with the percentage of each type in the bracket; noc/bb: number of contacts with the ligand, with the number of contacts realized through backbone atoms given in the bracket; dist: distance of closest approach to the ligand.)

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>subst’s (%)</th>
<th>cvg</th>
<th>noc/bb</th>
<th>dist (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>445</td>
<td>H</td>
<td>H(88) N(1) Q(5) .(4) DT .(3) TVA</td>
<td>0.03</td>
<td>7/1</td>
<td>2.10</td>
</tr>
<tr>
<td>507</td>
<td>C</td>
<td>C(89) S(5)</td>
<td>0.03</td>
<td>3/1</td>
<td>2.13</td>
</tr>
<tr>
<td>512</td>
<td>H</td>
<td>H(88) R(3) S(1) .(4) G N(1) QTA</td>
<td>0.05</td>
<td>7/1</td>
<td>2.05</td>
</tr>
<tr>
<td>517</td>
<td>M</td>
<td>L(41) M(32) Q(6) .(5) F(11) KI VGE</td>
<td>0.12</td>
<td>4/0</td>
<td>2.90</td>
</tr>
<tr>
<td>509</td>
<td>I</td>
<td>I(48) F(7) V(10) M(2) N(6) .(3) K(1) HTY</td>
<td>0.21</td>
<td>5/1</td>
<td>4.16</td>
</tr>
</tbody>
</table>

Table 7. List of disruptive mutations for the top 25% of residues in 1aozA, that are at the interface with copper (ii) ion.

<table>
<thead>
<tr>
<th>res</th>
<th>type</th>
<th>disruptive mutations</th>
</tr>
</thead>
<tbody>
<tr>
<td>445</td>
<td>H</td>
<td>(E) (T) (M) (VA)</td>
</tr>
<tr>
<td>507</td>
<td>C</td>
<td>(KR) (E) (Q) (H)</td>
</tr>
<tr>
<td>512</td>
<td>H</td>
<td>(E) (D) (M) (Q)</td>
</tr>
<tr>
<td>517</td>
<td>M</td>
<td>(Y) (H) (T) (R)</td>
</tr>
<tr>
<td>509</td>
<td>I</td>
<td>(Y) (R) (T) (H)</td>
</tr>
</tbody>
</table>

Figure 7 shows residues in 1aozA colored by their importance, at the interface with 1aozACU701.

Cu-o linkage binding site. Table 8 lists the top 25% of residues at the interface with 1aozAC1O703 (cu-o linkage). The following table (Table 9) suggests possible disruptive replacements for these residues (see Section 3.6).

Table 8. The top 25% of residues at the interface with 1aozAC1O703 (cu-o linkage). (Field names: res: residue number in the PDB entry; type: amino acid type; subst’s: substitutions seen in the alignment; with the percentage of each type in the bracket; noc/bb: number of contacts with the ligand, with the number of contacts realized through backbone atoms given in the bracket; dist: distance of closest approach to the ligand.)

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Table 8. The top 25% of residues in 1aozA at the interface with cu-o linkage. (Field names: res: residue number in the PDB entry; type: amino acid type; subst’s: substitutions seen in the alignment; with the percentage of each type in the bracket; noc/bb: number of contacts with the ligand, with the number of contacts realized through backbone atoms given in the bracket; dist: distance of closest approach to the ligand.)

Table 9. List of disruptive mutations for the top 25% of residues in 1aozA, that are at the interface with cu-o linkage.

Table 9. continued

Table 10. continued

Table 10. List of disruptive mutations for the top 25% of residues in 1aozA, that are at the interface with cu-o linkage.

2.4.3 Possible novel functional surfaces at 25% coverage. One group of residues is conserved on the 1aozA surface, away from (or substantially larger than) other functional sites and interfaces recognizable in PDB entry 1aoz. It is shown in Fig. 9. The right panel shows (in blue) the rest of the larger cluster this surface belongs to. The residues belonging to this surface “patch” are listed in Table 10, while Table 11 suggests possible disruptive replacements for these residues (see Section 3.6).

Fig. 8. Residues in 1aozA, at the interface with cu-o linkage, colored by their relative importance. The ligand (cu-o linkage) is colored green. Atoms further than 30 Å away from the geometric center of the ligand, as well as on the line of sight to the ligand were removed. (See Appendix for the coloring scheme for the protein chain 1aozA.)

Fig. 9. A possible active surface on the chain 1aozA. The larger cluster it belongs to is shown in blue.
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Table 10. Residues forming surface “patch” in 1aozA.

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3 NOTES ON USING TRACE RESULTS

3.1 Coverage

Trace results are commonly expressed in terms of coverage: the residue is important if its “coverage” is small - that is if it belongs to some small top percentage of residues [100% is all of the residues in a chain], according to trace. The ET results are presented in the form of a table, usually limited to top 25% percent of residues (or to some nearby percentage), sorted by the strength of the presumed evolutionary pressure. (i.e., the smaller the coverage, the stronger the pressure on the residue.) Starting from the top of that list, mutating a couple of residues should affect the protein somehow, with the exact effects to be determined experimentally.

3.2 Known substitutions

One of the table columns is “substitutions” - other amino acid types seen at the same position in the alignment. These amino acid types may be interchangeable at that position in the protein, so if one wants to affect the protein by a point mutation, they should be avoided. For example if the substitutions are “RVK” and the original protein has an R at that position, it is advisable to try anything, but RVK. Conversely, when looking for substitutions which will not affect the protein, one may try replacing, R with K, or (perhaps more surprisingly), with V. The percentage of times the substitution appears in the alignment is given in the immediately following bracket. No percentage is given when the substitution appears in the alignment is given in the immediately following bracket. No percentage is given in the cases when it is smaller than 1%. This is meant to be a rough guide - due to rounding errors these percentages often do not add up to 100%.

3.3 Surface

To detect candidates for novel functional interfaces, first we look for residues that are solvent accessible (according to DSSP program) by at least 10 Å², which is roughly the area needed for one water molecule to come in the contact with the residue. Furthermore, we require that these residues form a “cluster” of residues which have neighbor within 5 Å from any of their heavy atoms.

Note, however, that, if our picture of protein evolution is correct, the neighboring residues which are not surface accessible might be equally important in maintaining the interaction specificity - they should not be automatically dropped from consideration when choosing the set for mutagenesis. (Especially if they form a cluster with the surface residues.)

3.4 Number of contacts

Another column worth noting is denoted “noc/bb”; it tells the number of contacts heavy atoms of the residue in question make across the interface, as well as how many of them are realized through the backbone atoms (if all or most contacts are through the backbone, mutation presumably won’t have strong impact). Two heavy atoms are considered to be “in contact” if their centers are closer than 5 Å.

3.5 Annotation

If the residue annotation is available (either from the pdb file or from other sources), another column, with the header “annotation” appears. Annotations carried over from PDB are the following: site (indicating existence of related site record in PDB ), S-S (disulfide bond forming residue), hb (hydrogen bond forming residue), jb (james bond forming residue), and sb (for salt bridge forming residue).

3.6 Mutation suggestions

Mutation suggestions are completely heuristic and based on complementarity with the substitutions found in the alignment. Note that they are meant to be disruptive to the interaction of the protein with its ligand. The attempt is made to complement the following properties: small (AVGSTC], medium (LPNQDEMIK], large (WFYHR], hydrophobic (LPVAMWF], polar (GTCY]; positively [KHR], or negatively [DE] charged, aromatic (WFYH], long aliphatic chain (EKROQM], OH-group possession (SDET], and NH2 group possession (NQKR]. The suggestions are listed according to how different they appear to be from the original amino acid, and they are grouped in round brackets if they appear equally disruptive. From left to right, each bracketed group of amino acid types resembles more strongly the original (i.e. is, presumably, less disruptive) These suggestions are tentative - they might prove disruptive to the fold rather than to the interaction. Many researcher will choose, however, the straightforward alanine mutations, especially in the beginning stages of their investigation.

4 APPENDIX

4.1 File formats

Files with extension “ranks_sorted” are the actual trace results. The fields in the table in this file:

- alignment# number of the position in the alignment
- residue# residue number in the PDB file
- type amino acid type
- rank rank of the position according to older version of ET
- variability has two subfields:
  1. number of different amino acids appearing in in this column of the alignment
  2. their type
- rho ET score - the smaller this value, the lesser variability of this position across the branches of the tree (and, presumably, the greater the importance for the protein)
- cvg coverage - percentage of the residues on the structure which have this rho or smaller
- gaps percentage of gaps in this column

4.2 Color schemes used

The following color scheme is used in figures with residues colored by cluster size: black is a single-residue cluster; clusters composed of more than one residue colored according to this hierarchy (ordered

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</table>
4.3 Credits

4.3.1 Alistat *alistat* reads a multiple sequence alignment from the file and shows a number of simple statistics about it. These statistics include the format, the number of sequences, the total number of residues, the average and range of the sequence lengths, and the alignment length (e.g. including gap characters). Also shown are some percent identities. A percent pairwise alignment identity is defined as *(idents / MIN(len1, len2))* where idents is the number of exact identities and len1, len2 are the unaligned lengths of the two sequences. The "average percent identity", "most related pair", and "most unrelated pair" of the alignment are the average, maximum, and minimum of all *(N)(N-1)/2* pairs, respectively. The "most distant seq" is calculated by finding the maximum pairwise identity (best relative) for all *N* sequences, then finding the minimum of these *N* (relative) for all *N* sequences, then finding the minimum of all *(N)(N-1)/2* pairs, respectively. The "most distant sequence alignment at *Importance*" is copyrighted by W. Kabsch, C. Sander and MPI-MF, 1983, 1985, 1988, 1994-1995, CMBI version.

4.3.2 CE To map ligand binding sites from different source structures, *report_maker* uses the CE program: http://cl.sdsc.edu/. Shindyalov IN, Bourne PE (1998) “Protein structure alignment by incremental combinatorial extension (CE) of the optimal path”. Protein Engineering 11(9) 739-747.

4.3.3 DSSP In this work a residue is considered solvent accessible if the DSSP program finds it exposed to water by at least 10Å², which is roughly the area needed for one water molecule to come in the contact with the residue. DSSP is copyrighted by W. Kabsch, C. Sander and MPI-MF, 1983, 1985, 1988, 1994-1995, CMBI version by Elmar.Krieger@cmbi.kun.nl November 18,2002.


http://swift.cmbi.kun.nl/swift/hssp/

4.3.5 LaTeX The text for this report was processed using *LaTeX*: Leslie Lamport, “*LaTeX*: A Document Preparation System Addison-Wesley,” Reading, Mass. (1986).


http://www.drive5.com/muscle/

4.3.7 Pymol The figures in this report were produced using Pymol. The scripts can be found in the attachment. Pymol is an open-source application copyrighted by DeLano Scientific LLC (2005). For more information about Pymol see http://pymol.sourceforge.net/. (Note for Windows users: the attached package needs to be unzipped for Pymol to read the scripts and launch the viewer.)

4.4 Note about ET Viewer

Dan Morgan from the Lichtarge lab has developed a visualization tool specifically for viewing trace results. If you are interested, please visit:

http://mammoth.bcm.tmc.edu/traceview/

The viewer is self-unpacking and self-installing. Input files to be used with ETV (extension .etvx) can be found in the attachment to the main report.

4.5 Citing this work


4.6 About report_maker

*report_maker* was written in 2006 by Ivana Mihalek. The 1D ranking visualization program was written by Ivica Reš. *report_maker* is copyrighted by Lichtarge Lab, Baylor College of Medicine, Houston.

4.7 Attachments

The following files should accompany this report:

- 1aozA.complex.pdb - coordinates of 1aozA with all of its interacting partners
- 1aozA.etvx - ET viewer input file for 1aozA
- 1aozA.cluster_report.summary - Cluster report summary for 1aozA
- 1aozA.ranks - Ranks file in sequence order for 1aozA
- 1aozA.clusters - Cluster descriptions for 1aozA
- 1aozA.msf - the multiple sequence alignment used for the chain 1aozA
- 1aozA.descr - description of sequences used in 1aozA msf
- 1aozA.ranks_sorted - full listing of residues and their ranking for 1aozA
- 1aozA.1aozANAG601.if.pml - Pymol script for Figure 5
- 1aozA.cbcvg - used by other 1aozA-related pymol scripts
- 1aozA.1aozAC2O702.if.pml - Pymol script for Figure 6
- 1aozA.1aozACU701.if.pml - Pymol script for Figure 7
- 1aozA.1aozAC1O703.if.pml - Pymol script for Figure 8