Crystal Structure of Human L-Isoaspartyl Methyltransferase*

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The enzyme L-isooaspartyl methyltransferase initiates the repair of damaged proteins by recognizing and methylating isomerized and racemized aspartyl residues in aging proteins. The crystal structure of the human enzyme containing a bound S-adenosyl-L-homocysteine cofactor is reported here at a resolution of 2.1 Å. A comparison of the human enzyme to homologs from two other species reveals several significant differences among otherwise similar structures. In all three structures, we find that three conserved charged residues are buried in the protein interior near the active site. Electrostatics calculations suggest that these buried charges might make significant contributions to the energetics of binding the charged S-adenosyl-L-methionine cofactor and to catalysis. We suggest a possible structural explanation for the observed differences in reactivity toward the structurally similar L-isooaspartyl and D-aspartyl residues in the human, archaeal, and eu bacterial enzymes. Finally, the human structure reveals that the known genetic polymorphism at residue 119 (Val/Ile) maps to an exposed region away from the active site.

The loss of biological functions in the aging process can be attributed in part to nonenzymatic chemical reactions that degrade biomolecules including DNA, proteins, and small molecules. We have been interested in enzymatic mechanisms from which organisms have evolved to limit the accumulation of isomerized and racemized aspartyl residues, which form spontaneously from normal L-aspartic acid and L-asparagine residues within proteins as they age. Much attention has been directed at understanding the action of the L-isooaspartate(L-aspartate) O-methyltransferase or protein L-isooaspartyl methyltransferase (PIMT) (EC 2.1.1.77). PIMT is an enzyme that recognizes altered aspartyl residues in proteins and polypeptides and drives their eventual conversion to normal L-aspartyl residues (1–3). In the best-characterized pathway, a methyl group is transferred from S-adenosyl-L-methionine (AdoMet) to form a methyl ester on the α-carboxyl group of an L-isooaspartyl residue. Subsequent nonenzymatic reactions result in rapid L-succinimide formation and hydrolysis, which generates some “repaired” L-aspartyl residues as well as some L-isooaspartyl residues, which can then enter the cycle again for eventual conversion to the normal peptide linkage. PIMTs have been highly conserved during evolution and are found in almost all eucaryotic cells as well as in most archaeabacteria and most Gram-negative eu bacteria (4, 5).

Recently, three-dimensional structures have been determined for PIMT from the hyperthermophilic eubacterium Thermotoga maritima (6) and the thermophilic archaebacterium Pyrococcus furiosus (7). With the exception of a C-terminal domain apparently unique to the T. maritima enzyme, these enzymes have similar amino acid sequences and a similar three-dimensional fold (5). For the P. furiosus enzyme, the crystal structures of AdoMet-ligated and isooaspartyl peptide-ligated forms have revealed how the enzyme can recognize a wide variety of damaged L-isooaspartyl and D-aspartyl residues while excluding normal L-aspartyl residues (7). However, despite their sequence similarities, the kinetic properties and substrate specificities of the enzymes from various species are distinct. For example, the T. maritima enzyme is unable to methylate polypeptides containing D-aspartyl residues (5), whereas the P. furiosus enzyme is able to recognize such racemized residues (8).

With an interest in mammalian aging, efforts have been focused on understanding the three-dimensional structure and substrate specificity of the human form of the enzyme. A particularly interesting puzzle is why the human and P. furiosus enzymes have the ability to also repair methylated D-aspartyl residues and why these enzymes also appear to have a considerably higher affinity for L-isooaspartyl residues than do many plant and eu bacterial enzymes (9). The human enzyme is produced in two alternatively spliced forms that have distinct C termini (10, 11). There is also a widespread genetic polymorphism that results in the presence of either a valine or isoleucine residue at position 119 (12).

EXPERIMENTAL PROCEDURES

Data Collection and Processing—The human PIMT was cloned, expressed, and purified (13), and subsequent crystallization screenings were performed in the presence of an acidic solution that was saturated with respect to dithiothreitol and AdoHcy as described previously (14–16). The best crystals were grown from a solution of 0.2 M MgAc, 0.1 M cacydatate, pH 6.5, and 20% polyethylene glycol 8000. Data were collected in 1° oscillations using one cryo-cooled crystal at beamline X8C at the National Synchrotron Light Source. A data set was collected to 2.1-Å resolution and processed using DENZO and SCALEPACK (17). The crystal belongs to the monoclinic space group P21, with a = 48.0 Å, b = 53.7 Å, c = 49.0 Å, β = 115.6°, and one molecule in the asymmetric unit. In the initial data processing, an ice-ring caused many reflections to be discarded, resulting in an acceptable value for Rmerge (5.3%) but poor

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‡ The abbreviations used are: PIMT, protein L-isooaspartyl methyltransferase; AdoHcy, S-adenosyl-L-homocysteine; AdoMet, S-adenosyl-L-methionine; K_B, Boltzman constant times temperature.

§ The atomic coordinates and structure factors (code 1KR5) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics, Rutgers University, New Brunswick, NJ (http://www.rcsb.org/).

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coverage (completeness = 82.5%). This incompleteness hindered structure determination. By re-integrating the data with a higher threshold for background noise, the number of reflections increased from 10,970 to 12,512, and the completeness and \( R_{	ext{merge}} \) were increased to 94.7 and 7.4% overall. In the outermost resolution shell, the corresponding values were 75.0 and 25.6%.

**Structure Solution and Refinement**—The structure was solved by molecular replacement with the program EPMR (18) using the structure of PIMT from *P. furiosus* (7) as a search model (Protein Data Bank accession number 1JG1). In the resulting \( 2F_o - F_i \) electron density map, an atomic model was easily built and adjusted using the program O (19). The structure was refined by simulated annealing and B-factor refinement as implemented in the program CNS (20). After three rounds of model building and refinement, water molecules were inserted into the model, and the refinement was converged after an additional three cycles. The final model contains residues 7–224, S-adenosyl-L-homocysteine, and 64 water molecules. The stereochemistry was verified with the programs PROCHECK (21) and ERRAT (22). The final \( R \) value is 22.3% (\( R_{	ext{merge}} = 26.4\% \)) (Table I).

**Electrostatics Calculations**—The electrostatic potential in the region of space encompassing the protein molecule was calculated by solving the Poisson-Boltzmann equation on a three-dimensional grid as described by Warwicker and Watson (23) and implemented by others (24, 25) using locally written software. The grid spacing was 0.75 Å. The solvent and protein regions were assigned dielectric values of 78 and 4, respectively, and the Debye constant was taken to be 0.128 Å\(^{-1}\). Asp, Glu, Arg, and Lys residues were assigned their full charges. The linear finite difference equations were solved by iteration using simultaneous overrelaxation and Chebyshev acceleration (26). The values for the resulting electrostatic potential at particular points of interest were obtained by trilinear interpolation on the grid.

**RESULTS AND DISCUSSION**

**Protein Fold**—The human PIMT enzyme folds as a doubly wound \( \alpha/\beta/\alpha \) sandwich structure as expected for this superfamily of enzymes (27, 28). As recently reported in the structures from *T. maritima* (6) and *P. furiosus* (7), \( \beta \)-strands 6 and 7 are rearranged relative to the classical Rossman fold in what is now a familiar trademark of the PIMT enzymes. The labeling of secondary structural elements is shown in Fig. 1A illustrating the central \( \alpha/\beta/\alpha \) motif along with additional N-terminal \( \alpha \)-hel-
ices and one C-terminal α-helix. It has been reported that the bovine enzyme contains a disulfide bond linking Cys-42 to Cys-94 (29). In the structure reported here, those residues are 20 Å apart, suggesting that no disulfide bond exists in the native enzyme, and that the formation of such a disulfide may occur during the processing of tryptic peptides for analysis.

**Comparison with Other PIMT Structures**—The structure of human PIMT is shown in the same orientation in Fig. 1B along with the aligned PIMT structures from *T. maritima* (1DL5) and *P. furiosus* (1JG3) with root mean square differences for the aligned C-α atoms of 1.4 and 0.9 Å, respectively. There are a few notable features that distinguish the human PIMT struc-
tured from those structures reported recently from other organisms. In the human enzyme, the C-terminal part of the chain takes a different course. Also in the human enzyme, the loop following α-helix B contains a short α-helix not found in the two previously reported structures (Fig. 2C). Additional minor variations are apparent in the loop regions before and after β-strand 6.

Electrostatic Effects of Buried Charges Near the Active Site—An inspection of the crystal structure surprisingly revealed three charged amino acid residues (Asp-83, Arg-36, and Asp-109) essentially buried in the protein interior. Buried charges are relatively rare because of the energetic cost of desolvating a charge and moving it into the protein interior where the dielectric value is much lower than it is in the aqueous solvent. When buried charges do occur, the low dielectric value can produce significant electrostatic effects. The occurrence in this enzyme of three buried charges seems especially unusual. These charged residues are also relatively well conserved across the PIMT family of enzymes, further implicating them in some functional role. Although the charged residues are also present in the other two recently reported PIMT structures (Asp-97, Arg-48, and Glu-121 in P. furiosus and Glu-81, Arg-27, and Glu-107 in T. maritima), the electrostatic effects of these residues were not discussed in earlier analyses.

The buried protein charges are especially interesting in view of the charged nature of the reactive groups; the cofactor bears a sulfonium, and the polypeptide substrates bear an attacking carboxylate. It may also be relevant that the reaction is accompanied by a neutralization of the charges on both reactive groups. Because all three buried charges lie in the vicinity of the active site and the cofactor binding site, they might make important contributions to the energetics of binding or catalysis.

The electrostatic potential was calculated (see “Experimental Procedures”) and examined in detail at the positions that would be occupied by the sulfonium and the isoaspartyl substrate carboxylate in a productive complex. The contributions to the electrostatic potential were included from the protein as well as from charged groups on the AdoMet cofactor excluding the sulfonium charge itself. The protein charge closest to the reactive groups is Asp-109 (5.6 Å). As a result, the protein makes a significant net negative contribution to the potential experienced by the charged reactive groups in the active site. In units of k_BT per elementary charge, the protein produces a very high electrostatic potential of −18 k_BT at the sulfonium and −6 k_BT at the carboxylate.

Mutagenesis experiments could help clarify the possible roles of buried charges in binding and catalysis, but the electrostatic calculations already provide some clues. It is noteworthy that the electrostatic potential difference at the two reactive charged groups does not appear to specifically promote the chemical step (methyl transfer). In fact, such an electrostatic driving force would probably be unproductive at this point in the reaction mechanism, because the reaction is energetically favorable as it is even in the unbound state. Rather, the electrostatic contributions from the protein may be serving to help bind the charged AdoMet form of the cofactor by partially compensating for the significantly unfavorable energy associated with binding and desolvating its charged sulfonium in the buried active site cleft. Given the extraordinary burial of the cofactor, it seems likely that the rate-limiting steps in the reaction cycle could involve cofactor binding or release. The strong electrostatic interactions between buried charges on the protein and the cofactor could play important roles in those steps of the reaction cycle.

Role of Proline 45 in Substrate Specificity—A proline residue conserved in the active site of archael and eucaryotic enzymes may explain the ability of these enzymes to catalyze methylation of d-Asp more readily than the eubacterial enzymes. One of the recently reported structures (7) from P. furiosus was determined in the presence of a polypeptide substrate, VYP’(isoAsp)pHA. This structural complex provides a framework for understanding the substrate specificities of the enzymes from various sources. L-isoAsp residues can be methylated by PIMT from all three organisms, whereas the d-Asp amino acid can be methylated by PIMT from human and P. furiosus (8) but not from T. maritima (5). PIMT from human and P. furiosus both have a proline in a key position (Pro-49/Pro-65) at which PIMT from T. maritima is replaced by a valine (Val-45). When the structure of the l-isoAsp polypeptide substrate visualized in the P. furiosus structure is overlaid onto the other two protein structures, no steric clashes are seen with either the human or the T. maritima enzyme (Fig. 2A). Contrasting results are obtained when a similar analysis is performed with a D-Asp-containing polypeptide, which was modeled earlier on the basis of the l-isoAsp structure (7). The analysis suggests that the carbonyl group of the D-Asp would clash with CG2 in Val-45 in the structure from T. maritima (Fig. 2B) but not with the equivalently positioned proline in human and P. furiosus. The idea that residue 45 plays a key role in substrate specificity could be tested by mutagenesis. Counter to this argument, a proline residue is present in the nematode enzyme, which does not recognize d-aspartyl residues. Thus, other factors may also be important for the ability of the human and P. furiosus enzymes to recognize d-aspartyl residues with high efficiency.

Polymorphism—Because of a genetic polymorphism in Cau-
asian populations, two alternative alleles of the isoaspartyl methyltransferase PCMT1 gene (on chromosome 6) lead to the production of enzymes containing either a valine or isoleucine residue at position 119 (12). The presence of the isoleucine residue is associated with increased thermal stability of the enzyme, whereas the presence of the valine residue is associated with higher apparent affinity for protein, but not peptide, methyl-accepting substrates (12). In the human methyltransferase structure reported here, Val-119 is found to be almost completely exposed. The residue is located on the opposite side of the molecule from the methylation active site, so it is not immediately clear how the change from a valine to an isoleucine residue can affect the affinity for protein substrates. However, the larger isoleucine residue would be expected to form a hydrophobic interaction with Leu-130 located in α-helix 7 and might thereby increase the stability of the protein (Fig. 2C).

In addition to the genetic polymorphism, an alternative splicing of the human PCMT1 gene transcript in most tissues leads to two electrophoretic variants of the enzyme (10, 11). These products differ at the C terminus by the presence of WK or DEI as the terminal amino acid sequence. In the structure seen here, clear electron density is not observed for the C-terminal residues after 224, and it appears that these residues probably project from the enzyme surface.

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REFERENCES
Structure of Human L-Isoaspartyl Methyltransferase