RESEARCH ARTICLE

The Crystal Structure of D-Threonine Aldolase from *Alcaligenes xylosoxidans* Provides Insight into a Metal Ion Assisted PLP-Dependent Mechanism

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Abstract

Threonine aldolases catalyze the pyridoxal phosphate (PLP) dependent cleavage of threonine into glycine and acetaldehyde and play a major role in the degradation of this amino acid. In nature, L- as well as D-specific enzymes have been identified, but the exact physiological function of D-threonine aldolases (DTAs) is still largely unknown. Both types of enantio-complementary enzymes have a considerable potential in biocatalysis for the stereospecific synthesis of various β-hydroxy amino acids, which are valuable building blocks for the production of pharmaceuticals. While several structures of L-threonine aldolases (LTAs) have already been determined, no structure of a DTA is available to date. Here, we report on the determination of the crystal structure of the DTA from *Alcaligenes xylosoxidans* (AxDTA) at 1.5 Å resolution. Our results underline the close relationship of DTAs and alanine racemases and allow the identification of a metal binding site close to the PLP cofactor in the active site of the enzyme which is consistent with the previous observation that divalent cations are essential for DTA activity. Modeling of AxDTA substrate complexes provides a rationale for this metal dependence and indicates that binding of the β-hydroxy group of the substrate to the metal ion very likely activates this group and facilitates its deprotonation by His193. An equivalent involvement of a metal ion has been implicated in the mechanism of a serine dehydratase, which harbors a metal ion binding site in the vicinity of the PLP cofactor at the same position as in DTA. The structure of AxDTA is completely different to available structures of LTAs. The enantio-complementarity of DTAs and LTAs can be explained by an approximate mirror symmetry of crucial active site residues relative to the PLP cofactor.
**Introduction**

Threonine aldolases are pyridoxal phosphate (PLP)-dependent enzymes which catalyze the reversible cleavage of β-hydroxy amino acids (e.g., threonine) into glycine and the corresponding aldehyde (e.g., acetaldehyde) [1]. Threonine aldolases show a high selectivity for the absolute configuration at the Cα atom, but generally possess only moderate stereoselectivity at the β-carbon (Fig 1). Thus, they can be classified into D-threonine aldolases (DTAs, EC 4.1.2.42) and L-threonine aldolases (LTAs, EC 4.1.2.5).

While their exact physiological function is still under discussion, they appear to play a major role in threonine degradation. For instance, Liu et al. report that TA catalyzed cleavage of threonine is not the major source of cellular glycine in wild-type *E. coli*, but opens an alternative pathway to glycine, if the major pathway via serine hydroxymethyltransferase is somehow blocked [2]. In a different study Kim et al. found that 4-phosphoerythronate dehydrogenase deficient *E. coli* strains need L-threonine aldolase to synthesize pyridoxal phosphate. LTA is one of several enzymes participating in this alternative PLP synthesis pathway, where it catalyzes the condensation of glycolaldehyde with glycine to form L-4-hydroxythreonine. The phosphorylation of this compound produces L-4-phosphohydroxythreonine which is an intermediate in the PLP synthesis pathway [3]. Even less is known about the role of D-specific threonine aldolases in nature.

The reverse reaction, i.e., the condensation of glycine and an aldehyde species, possesses a considerable potential in biocatalysis. Since TAs accept a relatively large variety of aldehyde electrophiles, they allow the biocatalytic synthesis of various β-hydroxy-α-amino acids [1]. On the other hand, it was long believed that only glycine is accepted as the donor molecule. The work of the group of Griengl, however, showed that specific aldolases—the LTA from *Aeromonas jandaei* and the DTA from *Pseudomonas sp.*—also accept alanine instead of glycine as donor which paves the way to α,α'-dialkyl amino acids [4].

A number of crystal structures of LTAs have been determined in the recent past. These enzymes came from different organisms (including *E. coli*, *Pseudomonas putida* and *Thermotoga maritima*) but all share the same overall fold, which is similar to the fold of aspartate-amionotransferases [5]. The proposed catalytic mechanism of these enzymes involves the formation of the external aldimine intermediate—typical for PLP-dependent enzymes—and the deprotonation of the β-hydroxy group of this species by an active site histidine residue [5].

First D-threonine aldolase encoding genes were discovered in 1997 [6] and Liu et al. showed that DTAs are PLP-dependent and additionally need divalent ions to be catalytically active [7]. Based on homology models, these enzymes were classified as belonging to the alanine racemase family (fold-type III) [8] indicating a different evolutionary origin compared to LTAs. No crystal structure of a DTA is available to date. The alanine-racemase family includes bacterial racemases, eukaryotic ornithine decarboxylases as well as D-serine deaminases. The structures of these proteins consist of a typical alanine-racemase-like domain (eight-stranded α/β-barrel) together with a region mainly composed of β-strands. The work of Seebeck and Hilvert underlined the relation between racemases and aldolases. They were able to redesign an alanine...
racemase from *Geobacillus stearothermophilus* into a D-threonine aldolase by replacing a single active site tyrosine (Tyr265) by alanine [9,10].

Here, we report on the determination of the crystal structure of the DTA from *Alcaligenes xylosoxidans* (AxDTA) at 1.5 Å resolution. Our results allow the identification of a metal binding site close to the PLP-cofactor in the active site of the enzyme. Modeling of AxDTA substrate complexes provides a rationale for the metal dependence of DTAs. The results indicate that binding of the β-hydroxy group of the substrate to the metal ion may activate this group and thereby facilitate its deprotonation. Based on our structure, the enantio-complementarity of DTAs and LTAs can be explained by an approximate mirror symmetry of crucial active site residues relative to the PLP-cofactor.

**Materials and Methods**

**Reagents**

All chemicals were of the highest grade commercially available from Sigma-Aldrich (St. Louis, MO, U.S.A.), Fluka (Buchs, Switzerland) or Merck (Darmstadt, Germany). Crystallization equipment and materials were from Hampton Research (Aliso Viejo, CA), Molecular Dimensions (Newmarket, UK) and Douglas Instruments (Hungerford, UK).

**Cloning of the D-threonine aldolase gene**

Genomic DNA from *Alcaligenes xylosoxidans* IFO 12669 was isolated using Qiagen Genomic-tip 100/G (Qiagen, Hilden, Germany). PCR amplifications were performed using the genomic DNA as template. The oligonucleotides used as primers were forward (5′–3′): CACCATGTC CCAGGAAGTCATACGCGGC and reverse (5′–3′): TCAGCGCGARAARCCSCGC. The forward primer contained an ATG start codon and the reverse primer contained TCA nucleotides complementary to a TGA stop codon (in bold in the sequence). A four nucleotides CACC 5′-overhang (underlined in the sequence) was added to the forward primer to allow cloning into the pET101/D-TOPO vector (Invitrogen). PCR reactions were carried out in 50 μl *Pfx* amplification buffer (Invitrogen), 0.3 mM dNTP, 1 mM MgSO4, 15 pmol of each primer (45 pmol for a degenerated primer), 1 μg of genomic DNA, and 1.25 units of Platinum *Pfx* DNA polymerase (Invitrogen). Temperature cycling was as follows: (1) 96°C for 4 min; (2) 96°C for 1 min, 64°C for 1 min, and 68°C for 1.5 min during 30 cycles.

The amplified PCR products were analyzed by agarose gel electrophoresis. The fragments with the correct size were excised from the gel and purified with a Gel extraction kit (Qiagen). The restriction pattern of the purified PCR fragments was checked. PCR fragments with the expected restriction pattern were inserted in the pET101/D-TOPO vector according to the protocol described by Invitrogen. The resulting construct pET101/DTA was used to transform *Escherichia coli* TOP 10 cells and these cells were grown on selective medium (LB with 100 μg/ml carbenicillin). Plasmid DNA of recombinant TOP10 clones was isolated and checked by restriction analysis. Plasmid DNA with the expected size and restriction pattern was used to transform *E. coli* BL21 Star (DE3) cells.

**Expression and purification**

The purification was based on the procedure published by Liu et al. [7]. The DEAE Sepharose Fast Flow FPLC purification was the only step carried out at room temperature. All other purification operations were performed at 0 to 4°C. An aqueous solution of 50 mM Tris/HCl (pH 7.5), 20 μM PLP and 1.3 mM DTT was used as buffer.
E. coli BL21 Star (DE3) cells harboring the pET101/DTA plasmid were grown aerobically at 28°C in 2x1 liter LB-medium containing 100 μg/ml carbenicillin. Protein expression was induced with 0.002% (w/w) L-arabinose at an OD_{620} of approximately 0.6. After overnight incubation at 28°C the cells were centrifuged for 15 min at 12000 g, washed with buffer and re-centrifuged for 10 min at 8300 g. The pellet was stored at -20°C until the sonication procedure took place. After resuspension in a buffer mass corresponding to five times the mass of the cell wet weight and addition of 100 μM MnCl₂, the cells were disrupted by ultrasonic oscillation on an ice/acetone mixture for 3 x 4 min. The cell debris was removed by centrifugation at 50000 g for 20 min. The supernatant was stored at -20°C over night.

An additional centrifugation at 20000 g for 10 min was necessary to clear the milky solution after thawing. The clear supernatant solution was brought stepwise to 25% saturation with ammonium sulfate, incubated overnight with shaking and centrifuged at 25000 g for 30 min. The precipitate collected by centrifugation was dissolved overnight in the MnCl₂ containing buffer. Due to the incomplete dissolution of the precipitate the solution was re-centrifuged at 25000 g for 10 min before application on the DEAE Sepharose column.

The supernatant was applied to a DEAE Sepharose Fast Flow column (XK 16, 1.6 cm diameter with 25 ml bed volume) equilibrated with the buffer (MnCl₂). After the column had been washed thoroughly with the buffer, a linear gradient elution was performed with the buffer supplemented by NaCl thereby increasing the concentration from 0 to 300 mM within 20 column volumes. The flow rate was maintained at 2.5 ml/min. The peak fractions were tested for threonine aldolase activity and the active fractions were analyzed using SDS-PAGE.

The purified fractions were pooled, applied to a Centricon Plus-20 ultrafiltration unit with a 30000 Dalton molecular weight cut off (Amicon Bioseparations) and desalted by applying 4 times 10 mM Na-Hepes (pH 7), 100 μM PLP, 100 μM MnCl₂ and 1 mM DTT. The retentate was diluted with the same buffer to a final volume of approximately 1.5 ml. The sample was aliquoted and stored at -80°C.

Crystallization

Crystallization drops containing purified AxDTA at a concentration of 20 mg/ml in 10 mM Hepes, pH 7.0, 0.1 mM MnCl₂, 1 mM DTT were set up with commercially available screens using the micro batch as well as the vapor batch method on an Oryx-7 crystallization robot (Douglas Instruments Ltd.). Although these trials yielded initial crystals in several conditions, we failed to identify crystals which diffracted sufficiently for data collection. Hence, after many attempts to manually optimize the crystallization conditions, we applied the method of selective lysine methylation, following the protocol of Rayment et al. [11]. AxDTA samples were dialyzed against 200 mM sodium borate buffer pH 8.5, diluted to a concentration of 1 mg/ml and treated with formaldehyde and sodium borohydride to selectively methylate lysine residues. The reaction was quenched by adding an excess of glycine (3-fold amount compared to the added formaldehyde). Before setting up crystallization trials the solution was dialyzed against the original buffer (10 mM Hepes, pH 7.0, 0.1 mM MnCl₂, 1 mM DTT) and concentrated to ~10 mg/ml.

Crystallization trials with methylated AxDTA were again set up with different commercially available crystallization screens using the micro batch method. Crystallization conditions were manually optimized using the vapor batch setup by mixing equal amounts (1.0 μL) of protein and precipitant. Crystals of AxDTA grew within one day after streak seeding from a solution containing 200 mM sodium chloride, 100 mM Tris/HCl, pH 8.1 and 13% (w/v) PEG-3350. For diffraction data collection, crystals were harvested from their mother liquor with Cryo-Loops (Hampton Research) and were cryo-protected by soaking in mother liquor containing...
20% (v/v) 2-methylpentane-2,4-diole (MPD) for a few seconds prior to flash-cooling in liquid nitrogen.

**Structure determination and refinement**

A complete diffraction dataset was collected from a single crystal at the Swiss Light Source (SLS) of the Paul Scherrer Institute in Villingen, Switzerland (beamline X06DA). A full dataset was collected from a monoclinic crystal (space group $P_2_1$) up to 1.5 Å resolution and processed with the program AUTOMAR (marScale version 3.09–6, MarResearch). The calculated Matthews coefficient [12,13] indicated a 99% probability for the presence of two AxDTA molecules per asymmetric unit. The structure was determined by following the protocol of DiMaio et al. [14] employing the programs PHASER [15] and ROSETTA [16]. Suitable template structures were identified using the web server HHpred [17]. The highest-ranking models (PDB-entries 3GWQ and 3LLX) were used as initial molecular replacement templates for PHASER [15]. For the top five PHASER-solutions initial density maps were calculated and used as additional input for ROSETTA. Applying a 10% energy cutoff, ROSETTA built 2000 new models for each PHASER solution. To evaluate and score these models in terms of structural relevance, PHASER runs in MR_RNP mode (molecular replacement, refinement and phasing) were performed. The best scoring models (based on log-likelihood statistics) were further used as input for the automated chain-tracing/rebuilding programs AutoBuild [18] and ArpWarp [19]. Both programs independently produced almost complete structures of both chains.

Structure refinement and model rebuilding were carried out with the programs PHENIX [18] and COOT [20] by alternating real-space fitting against $\sigma$A-weighted $2F_O-F_C$ and $F_O-F_C$ electron density maps and least square optimizations. $R_{\text{free}}$ values were computed from 5% randomly chosen reflections, which were not used during refinement [21]. Clear residual electron density was assigned to the PLP-cofactor, one sodium ion and one manganese ion in each protomer. No electron density was observed for amino acids 1–6 in protomer A as well as for residues 1–7 in protomer B. Water molecules were placed into the difference electron density map and accepted or rejected according to geometry criteria as well as refined B-factors. In the later stages of the refinement, two TLS groups per protomer (including the respective PLP-cofactor) were defined based on an analysis using the TLSMD web server [22]. In addition, anisotropic atomic displacement parameters were refined for the four cations (2 Mn$^{2+}$, 2 Na$^+$). The final model was refined to $R = 15\%$ and $R_{\text{free}} = 18\%$. Validation of the structure was carried out with the program MOLPROBITY [23] yielding a Ramachandran plot with 98.4% of the residues in favored, 1.5% in allowed and 0.1% in disallowed regions. Details pertaining to data statistics and structure refinement are listed in Table 1.

**Substrate docking**

To model putative binding modes of different substrates in the active site of AxDTA the program AutoDock 4.0 [24] was used as implemented in YASARA Structure [25]. Molecular models of the two diastereomers of D-phenylserine, (2R,3S)-phenylserine and (2R,3R)-phenylserine, covalently linked to the PLP cofactor (thereby representing the external aldimine) were generated and optimized in YASARA. The total net charge of each ligand was -2. Lys59 which binds PLP in the enzyme’s resting state was treated as uncharged. Position and orientation as well as torsion angles—except those involving atoms of the delocalized $\pi$-system of PLP—were allowed to vary. The docking was restricted to a 28x28x23 Å box around the center of the active site. Twenty independent simulation runs were performed on each ligand employing a genetic algorithm (population size 150, number of generations 22000). Structures with the lowest energy in each independent run were
clustered with a root-mean-square-deviation tolerance of 1.5 Å and subsequently filtered based on the binding mode of the PLP-part of the ligand compared to the crystal structure. Molecular mechanics optimization was done within YASARA using the YASARA2 force field and employing the standard optimization protocol [26–28].

Results and Discussion
Crystallization and structure determination

Although purified AxDTA crystallized under many different conditions from various commercially available screens, those crystals did not produce diffraction data suitable for structure determination and we obtained only weak diffraction to a maximum resolution of 3.5 Å. Hence,

| Table 1. Data collection and refinement statistics. |
|-----------------------------------------------|
| **Data collection**                              |
| X-ray source                              | SLS-X06DA                      |
| Wavelength (Å)                             | 0.9537                         |
| Temperature                                | 100 K                           |
| Space group                                | P2_1                           |
| a, b, c (Å)                                 | 62.27, 84.28, 72.85             |
| β (°)                                       | 111.9                          |
| Resolution (Å)                              | 47.65–1.50 (1.55–1.50)         |
| Reflections                                 | 1422127                         |
| Unique reflections                         | 111114                          |
| Multiplicity (β)                            | 7.4 (7.4)                      |
| Completeness (%) (β)                       | 99.6 (100)                     |
| R_{P.I.M.} (β)                              | 0.035 (0.132)                  |
| R_{merge} (β)                               | 0.084 (0.333)                  |
| <I/σ(I) (β)                                 | 6.3 (1.6)                      |
| C_C1/2 (β)                                 | 0.999 (0.969)                  |
| C_C* (β)                                   | 1.000 (0.992)                  |
| **Refinement**                              |
| Resolution (Å)                              | 35.76–1.50                     |
| R_{work} / R_{free}                         | 0.1484 / 0.1767                |
| **No. of atoms**                            |
| Protein                                     | 5798                           |
| Cofactor/substrate                          | 36                             |
| Water                                       | 955                            |
| **Mean B-factors (Å²)**                     |
| Protein                                     | 17.10                          |
| Cofactor/substrate                          | 13.60                          |
| Water                                       | 33.80                          |
| Other entities                              | 17.13                          |
| All atoms                                   | 19.50                          |
| **Root-mean-square-deviations**             |
| Bond lengths (Å)                            | 0.019                          |
| Bond angles (°)                             | 1.356                          |
| Ramachandran outliers (%)                  | 0.1                            |

Values in parentheses are for highest-resolution shell.

doi:10.1371/journal.pone.0124056.t001
we methylated AxDTA in order to improve crystal quality [11]. During methylation we observed a noticeable color change of the reaction solution from yellowish to colorless. Although, isoelectric focusing experiments failed to demonstrate significant differences between methylated and untreated samples, crystallization trials with methylated AxDTA yielded crystals diffracting well beyond 2 Å resolution. During structure refinement, however, we could not detect any signs of lysine methylation in the electron density. It appears that the improvement of crystal quality was rather a stabilizing side effect due to the added chemicals during methylation or due to the additional purification and dialysis step, than a consequence of the methylation itself.

We determined the structure of AxDTA by molecular replacement at a crystallographic resolution of 1.5 Å and refined it to final R-values of R = 15% and Rfree = 18%. The crystal contained two molecules per asymmetric unit connected by a non-crystallographic two-fold axis. For detailed data and refinement statistics see Table 1.

Three-dimensional structure of AxDTA

Each protomer comprises an alanine-racemase-like domain with an eight-stranded α/β-barrel (residues 32–274) together with a β-strand domain consisting of residues from the N-terminus (residues 1–31) and the C-terminus (residue 275–379). This β-domain can be subdivided into two β-stranded motifs composed of a closed, antiparallel 5-stranded β-barrel (residue 275–347) and a 3-stranded β-sheet built up by N- and C-terminal residues (Fig 2A). To classify the β-barrel domain we did a CATH database search employing the CATHEDRAL algorithm [29]. The best hit with an SSAP score of 79.9 and a root-mean-square-deviation (r.m.s.d.) of 3.1 Å

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Fig 2. Schematic representation of the structure of AxDTA. (A) Cartoon representation of the AxDTA protomer showing the alanine-racemase-like domain in red and the β-domain in blue. The PLP-cofactor bound to Lys59 is shown in yellow. The manganese ion is depicted as a magenta and the sodium ion as a green sphere. Residues coordinating the Mn- and Na-ion are shown as gray sticks, metal bound water molecules are shown as small red spheres. Metal coordination is indicated by light blue, dashed lines. (B) Cartoon representation of the AxDTA dimer with one protomer shown in darker red/blue (domain coloring as in A) and the other in lighter red/blue.

doi:10.1371/journal.pone.0124056.g002
belongs to the CATH superfamily 2.40.10.230, annotated as a probable tRNA pseudouridine synthase domain.

Previous studies indicated that AxDTA was active as a monomer [7]. For our crystal structure, however, a PISA analysis [30] predicted a stable dimer in solution. The structure of the AxDTA dimer is similar to dimers of alanine racemases and involves non-covalent contacts between both subdomains in a head-to-tail arrangement (Fig 2B). A superposition of the two crystallographically independent chains resulted in an r.m.s.d. of 0.12 Å for 302 out of 357 aligned Ca atoms. A superposition with the template structure used for molecular replacement (PDB-entry 3LLX) resulted in an r.m.s.d. of 1.27 Å (228 out of 375 aligned Ca atoms).

The active site of AxDTA is mainly built up from residues of the alanine racemase-like domain (Fig 3A). The smaller β-domain of the second protomer contributes residues from two loop regions (residues 295–298 and 319–323). Clear, residual electron density in the active site of both protomers was modeled by a molecule of pyridoxal phosphate (PLP) in two alternate conformations: covalently linked to Lys59 (occupancy ~40%) and non-covalently bound (occupancy ~60%). In general, PLP binds covalently to an active-site lysine in the resting state of the enzyme (referred to as the internal aldimine) [1]. In both cases, the pyridine ring of PLP forms π-stacking interactions with Tyr187 (distance ~3.7 Å). The pyridine nitrogen atom forms a hydrogen bond to Gln249. The phosphate group of the cofactor interacts with the hydroxyl groups of Thr233, Ser252 and Tyr260 as well as with main chain amide groups of Thr233 and Gly251. The hydroxyl group at the pyridine ring of PLP forms hydrogen bonds with the side chain amide of Gln81 and the guanidinium group of Arg157.

Liu et al. showed that divalent ions are crucial for the activity of D-threonine aldolases from Arthrobacter sp. [31] as well as from Alcaligenes xylosoxidans [7]. In their experiments, manganese ions had the biggest effect. We indeed observed an ion binding site (large difference electron density) relatively close to the PLP cofactor. Since manganese was a buffer component in our crystallization trials (0.1 mM MnCl2, see Materials and Methods), we modeled this density by a Mn2+ ion. This ion fitted the density very well in both protomers and the coordination geometry was also consistent with a manganese bound at this position. The ion is coordinated only by two amino acid residues, His347 (via the Ne atom, 2.2 Å) and Asp349 (via the carboxylate, 2.2 Å), both from the alanine-racemase-like domain. Four water molecules (distances

![Fig 3. Structure of the active site and the metal binding sites of AxDTA.](https://example.com/fig3.png)
from 1.8 to 2.3 Å) complete a distorted octahedral coordination sphere. The metal binding site is located at approximately 5 Å distance from the aldehyde group of PLP (Fig 3A) and the manganese ion is nearly coplanar with the pyridine ring (0.4 Å out of the plane).

A second metal binding site was observed at the C-terminal end of an α-helix consisting of residues Asp162 to Pro177 (Fig 3B). Unassigned electron density at this position was modeled by a sodium ion, which was also a buffer component. The residues Ala173, Arg174, Leu176 and Leu179 at the C-terminal end of the helix provide carbonyl groups as ligands to the Na+ ion. Its positive charge is further stabilized by the helix dipole. The coordination sphere is completed by two water molecules. This metal binding site is located at the surface of the enzyme approximately 30 Å away from the active site.

Modeling of substrate complexes

In order to identify potentially crucial residues in the active site and to get an idea about the possible involvement of the manganese ion in the catalyzed reaction, we docked the two diastereomers of D-phenylserine, (2R,3S)-phenylserine (syn-D-phenylserine, D-threo-phenylserine) and (2R,3R)-phenylserine (anti-D-phenylserine, D-erythro-phenylserine), as external aldimine adducts to PLP into the active site of AxDTA. Phenylserine has been described as a good substrate for this enzyme [1,7,32]. A number of calculated, low-energy docking poses were discarded, because the PLP-part of the ligand did not bind in the same way as in the crystal structure as judged by the position and orientation of the pyridine ring and the position of the phosphate group. The remaining binding modes were analyzed with respect to chemical and mechanistic plausibility, especially regarding the fulfillment of the Dunathan stereo-electronic requirements [33]. According to the Dunathan hypothesis, the Cα-Cβ bond is supposed to be approximately perpendicular to the plane of the pyridine ring of PLP in order to facilitate C-C bond breaking in the threonine aldolase reaction. This filtering process yielded a single binding mode for the external aldimine of (2R,3S)-phenylserine which fulfilled all constraints, whereas no suitable binding mode of the external aldimine of (2R,3R)-phenylserine was observed. This is qualitatively in line with the known syn-diastereospecificity of the enzyme [1,32].

The resulting complex structure was further geometry optimized and is shown in Fig 4. In this structure the carboxylate of the substrate forms a salt bridge with the guanidinium group of Arg157 and is hydrogen bonded to the main-chain amide group of Asp321. The
Cβ-hydroxy-group is coordinating the manganese ion with a Mn-O distance of 2.3 Å. In that position it replaces a water molecule that is bound to the ion in the crystal structure of AxDTA. Another water molecule forms a bridge between the OH-group of the substrate and the imidazole ring of His193. The phenyl ring is located in the entrance funnel to the active site and is pointing towards the exterior of the enzyme.

**Catalytic mechanism and stereospecificity**

Based on the crystal structure and the modeling results we propose the following mechanism for the degradation of β-hydroxy amino acids catalyzed by AxDTA (Fig 5). As usual for PLP-dependent enzymes the reaction cycle starts with a transaldimination to form the external aldimine. Once the external aldimine is formed, deprotonation of the β-OH-group of the substrate is facilitated by its coordination to the manganese ion and, most likely, involves His193 as the base. Proton abstraction is mediated by a water molecule. In the next step the Cα-Cβ bond is cleaved yielding an aldehyde and the resonance stabilized, deprotonated aldimine of glycine and PLP. Reprotonation of this intermediate could either involve Lys59 or His193. In the latter case, however, reprotonation would very likely have to be water mediated. The reaction cycle is completed by another transaldimination reaction to reform the resting state of the enzyme with the internal aldimine linkage to Lys59.

A PSI-Blast search using the amino acid sequence of AxDTA against the non-redundant NCBI-database yielded 25 additional sequences annotated as D-threonine aldolases. The application of a 90% sequence identity cutoff reduced the list to 19 sequences (including AxDTA) with identities ranging from 20 to 85% (with an average of 40%). A multiple sequence alignment using ClustalOmega [34] shows that all residues implicated in our proposed mechanism (including His193, the manganese coordinating residues and amino acids interacting with the PLP cofactor) are conserved among these enzymes even for homologues with low overall sequence identity.

The putative mechanism involves the manganese ion as a crucial Lewis acid, which is in line with previous findings showing that divalent metal ions are important for enzyme activity [7]. In contrast to well-studied PLP-dependent enzyme reactions assisted by monovalent ions, where the metal ion can be either directly involved in catalysis or via allosteric effects [35], detailed analyses of PLP-dependent reactions involving divalent ions have been described to a lesser extent [7,31,36,37]. A search in the PDB using the program Relibase+ [38] yielded only two structures, which resemble AxDTA in terms of metal binding architecture and the relative arrangement of the metal binding site and the PLP cofactor. One of them is an enzyme from *Idiomarina loihiensis* annotated as "predicted amino acid aldolase or racemase". Its structure (PDB entry 3LLX) was determined by the Joint Center for Structural Genomics (JCSG), but no paper has so far been published. It should be noted that this structure was used as the molecular replacement search template in the structure determination of AxDTA. The second enzyme is a D-serine dehydratase from chicken kidneys (PDB-entry 3ANU) [37]. In both cases, a zinc ion is bound at the same position as the manganese in AxDTA (Fig 6). In the case of the serine dehydratase it has been reported that the zinc ion can be replaced by manganese without loss of activity [37]. Based on crystal structures and molecular modeling a catalytic mechanism has been proposed for this enzyme, which is very similar to our proposed mechanism for AxDTA (Fig 5) and also involves activation of the substrate OH-group by coordinating to the metal ion [37]. Metal coordination in the dehydratase and the predicted aldolase/racemase, however, is achieved by a histidine and a cysteine instead of a histidine and an aspartate in AxDTA (Fig 6). In addition, the potential base His193 in AxDTA is not present in the other two enzymes.
Fig 5. Proposed catalytic mechanism of AxDTA.

doi:10.1371/journal.pone.0124056.g005
Because of the structural similarities of D-threonine aldolases and alanine racemases Seebeck et al. were able to redesign the racemase from Geobacillus stearothermophilus into a D-threonine aldolase by replacing a single active site tyrosine (Tyr265) by alanine [9]. The proposed retro-aldolase mechanism of this redesigned variant does not involve metal ion contributions but features the direct deprotonation of the substrate OH-group by a histidine (His166). This histidine forms π-stacking interactions with the PLP-cofactor and is structurally equivalent to Tyr187 in AxDTA (Fig 3A). While it is in principle conceivable that Tyr187 is the active site base in AxDTA, this residue appears not to be activated by surrounding residues as has been shown necessary in mechanisms of similar PLP-dependent enzymes [39]. In those cases, the tyrosine OH-group is hydrogen bonded to a histidine residue. In the structure of AxDTA no such or similar interactions of Tyr187 were observed. Therefore, we consider the direct involvement of this residue in the mechanism less likely.

L-specific threonine aldolases (LTAs) are more common in nature than D-specific enzymes. The overall structure of AxDTA presented here is completely different to the fold exhibited by L-threonine aldolases (e.g. the enzyme from Thermotoga maritima, PDB-entry 1LW4 [5]), which involves a 3-layer α/β-sandwich instead of an (α/β)8-barrel and is closely related to the fold of aspartate amino transferases. The superposition of the PLP-cofactor provides a rationale for the inverted enantiopreference of the two types of enzymes (Fig 7). For the LTA from Thermotoga maritima a mechanism has been proposed, in which the β-OH group of the substrate is deprotonated by an active site histidine residue [5] which is located at the si-face of the cofactor. In contrast to that, the proposed base His193 is positioned at the opposite, re-face of PLP. Thus, the cofactor can be seen as a pseudo mirror plane in the superposition (Fig 7). This situation is reminiscent of a recent study of enantio-complementary ene-reductases, where the FMN-cofactor serves as the approximate mirror plane [40]. According to a recent classification of enantio-complementarity in enzymes [41] the LTA/DTA-pair is a member of group 1, which includes enzyme pairs with different folds and mirror-image active sites.

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Fig 6. Superposition of the active site regions in AxDTA and the serine dehydratase from chicken kidney [37]. Amino acids are shown as gray (AxDTA) and pink (PDB-entry: 3ANU) sticks. The cofactors are shown in yellow and the metal ions are shown as gray and pink spheres. Metal coordination is indicated by gray (AxDTA) and pink (3ANU) dashed lines. Water molecules are shown as small red (AxDTA) and dark red (3ANU) spheres.

doi:10.1371/journal.pone.0124056.g006
Conclusions

The crystal structure of AxDTA clearly shows the relationship of D-threonine aldolases and alanine racemases. The role of the essential metal (manganese) ion is most likely to activate the β-hydroxy group of the substrate for deprotonation by His193. The enantio-complementarity of DTAs and LTAs can be explained by an approximate mirror symmetry of crucial active site residues relative to the PLP-cofactor.

Acknowledgments

We acknowledge the support of the beamline staff at SLS (Villigen, Switzerland) during diffraction data collection.

Author Contributions

Conceived and designed the experiments: MKU GO GS DM MS KG. Performed the experiments: MKU GO GS LRB FVA. Analyzed the data: MKU GO GS DM MS KG. Wrote the paper: MKU GO GS DM MS KG.

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