Crystal Structures Reveal that the Reaction Mechanism of Imidazoleglycerol-Phosphate Dehydratase Is Controlled by Switching Mn(II) Coordination

Graphical Abstract

Highlights

- IGPD forms open and closed complexes with IGP bound in two distinct conformations
- Mn(II) controls catalysis by switching between 6- and 5-coordination states
- A ligand-depleted 5-coordinate Mn(II) aids formation of the imidazolate intermediate
- Enzyme and substrate conformational changes are required for product formation

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In Brief

Bisson et al. determined structures that describe the reaction mechanism of IGPD, showing how the enzyme harnesses substrate binding energy to generate a ligand-depleted Mn(II) at the catalytic center. Conformational changes in the enzyme-substrate complex and switches in coordination chemistry control successive steps in catalysis.

Accession Numbers

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Crystal Structures Reveal that the Reaction Mechanism of Imidazolglycerol-Phosphate Dehydratase Is Controlled by Switching Mn(II) Coordination

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SUMMARY

Imidazolglycerol-phosphate dehydratase (IGPD) catalyzes the Mn(II)-dependent dehydration of imidazolglycerol phosphate (IGP) to 3-(1H-imidazol-4-yl)-2-oxopropyl dihydrogen phosphate during biosynthesis of histidine. As part of a program of herbicide design, we have determined a series of high-resolution crystal structures of an inactive mutant of IGPD2 from Arabidopsis thaliana in complex with IGP. The structures represent snapshots of the enzyme trapped at different stages of the catalytic cycle and show how substrate binding triggers a switch in the coordination state of an active site Mn(II) between six- and five-coordinate species. This switch is critical to prime the active site for catalysis, by facilitating the formation of a high-energy imidazolate intermediate. This work not only provides evidence for the molecular processes that dominate catalysis in IGPD, but also describes how the manipulation of metal coordination can be linked to discrete steps in catalysis, demonstrating one way that metalloenzymes exploit the unique properties of metal ions to diversify their chemistry.

INTRODUCTION

Metalloenzymes account for more than half of known enzymes (Foster et al., 2014) and are fundamental in facilitating many of the core metabolic processes that are essential for life. The chemical diversity and functionality of metalloenzymes is dependent upon their ability to utilize the unique chemistry of metal ions to facilitate specific reactions (Finkelstein, 2009; Frausto da Silva and Williams, 2001; Karlin, 1993). Numerous biophysical, biochemical, and structural studies have provided insights into how metal ions promote catalysis, highlighting their involvement in bond polarization and stabilization of transition states and intermediates (Andreini et al., 2008). Moreover, in the case of certain redox enzymes, for example manganese superoxide dismutase (Abreu and Cabelli, 2010) and oxalate decarboxylase (Tabares et al., 2009), changes in the oxidation state of the metal ions have been associated with variations in ligand coordination geometry during the reaction (Andreini et al., 2008). However, there is still much to be learned about the exact roles that different metal ions play in driving enzyme catalysis, specifically how the exquisite specificity with which they are selected is matched to the chemistry in which they are involved.

The metalloenzyme imidazolglycerol-phosphate dehydratase (IGPD) (EC 4.2.1.19) has been identified as a potential herbicide target due to its essential role in histidine biosynthesis (Ames, 1957; Hilton et al., 1965). It catalyzes the Mn(II)-dependent dehydration of (2R,3S)-2,3-dihydroxy-3-(1H-imidazol-5-yl) propyl dihydrogen phosphate (2R3S IGP) to 3-(1H-imidazol-4-yl)-2-oxopropyl dihydrogen phosphate (IAP). A previous structure of IGPD isoform 1 from Arabidopsis thaliana (IGPD1) (PDB: 2F1D) (Glynn et al., 2005) and a second from Mycobacterium tuberculosis (PDB: 4GUJ) (Ahangar et al., 2013) has shown that IGPD is a homo 24mer with two manganese ions, ~6.5 Å apart, bound at each of the catalytic centers. These structures, and other biochemical data, have revealed that, unlike other enzymes, where Mn2+ can be exchanged with Mg2+ or Zn2+ with little effect on activity (Andreini et al., 2008), IGPD has both a structural and mechanistic requirement for manganese (Petersen et al., 1997). Analysis of the IGPD1 structure, coupled with the observation that triazole-phosphonate compounds are potent inhibitors of the enzyme (Hawkes et al., 1993; Jin et al., 2015; Lindell et al., 1996; Mori et al., 1995), has suggested that the reaction (Figure 1A) proceeds via an initial high-energy deprotonation of the substrate imidazole ring (pK_a ~14.5) to yield an anionic imidazolate (Walba and Isensee, 1961), which is then converted to the product via diazafulvene and enol intermediates. Since the substrate lacks a carbonyl or imine adjacent to the leaving proton, which is thus non-acidic, IGPD uses an unusual mechanism for catalysis that is distinct from other dehydration reactions that have been reported to date (Glynn et al., 2005).

As part of a program of herbicide development designed to combat increasing resistance to glyphosate, we sought to expand our understanding of the enzyme mechanism by studying the structure of the IGPD in complex with its substrate, IGP,
and also with inhibitors. In this paper, we describe two distinct crystal structures of the enzyme-substrate complex, one that represents initial capture of the substrate by the enzyme and a second in which the substrate is bound as the activated high-energy imidazolate form that is primed for catalysis. Analysis of the interconversion between the two forms reveals that progression through the catalytic cycle is a highly dynamic process involving conformational changes to both the substrate and the protein. Central to this process is the repeated switch in coordination state of an active site Mn(II) between six- and five-coordinate species. These different coordination states are linked to specific steps in the reaction and serve to control progression through the catalytic cycle. This study therefore explains how an enzyme can exploit a feature of transition metals to augment the functionality of proteins, providing insights into one of the roles that divalent metal ions can play in catalysis.

**RESULTS**

The Form A Enzyme-Substrate Complex Contains a Ligand-Depleted Five-Coordinate Mn(II) Ion

To provide a foundation on which to analyze substrate binding, we determined the structure of A. thaliana IGPD isoform 2 (IGPD2) in complex with inorganic phosphate (1.75 Å resolution) (Table 1), which is a weak inhibitor of the enzyme ($K_i$ = $5$ mM) (Wiater et al., 1971a). The native protein crystallized as the same 24mer species that has been observed previously (Figure S2), and this structure shows that within the active site of the holoenzyme, both of the Mn(II) ions are octahedrally coordinated by four conserved protein ligands (Figures 1B and S1) and two water molecules; HOH3/HOH4 for Mn1 and HOH1/HOH2 (Figure 2A). The structure also shows that a phosphate ion binds in a site surrounded by a number of conserved positively charged residues (Q51, H55, R99, and K177) (Figure 2B), which are equivalent to those that surround the site of a bound sulfate ion in the IGPD1 complex (PDB: 2F1D) (Glynn et al., 2005). The holoenzyme has an open conformation where the C-terminal region (R193–R206, termed the C loop), which contains a number of conserved residues, is disordered (Figures 1B and S1).

To study substrate binding in IGPD, we used site-directed mutagenesis to produce an inactive mutant of A. thaliana IGPD2 by replacing E21, a putative acid-base catalyst, with a glutamine residue. The E21Q IGPD2 mutant was confirmed to be inactive using a modified version of the stopped assay previously described by Ames (1957) and Hawkes et al. (1995), which showed no signal above background. E21Q IGPD2 was co-crystallized with substrate to yield a 1.12 Å resolution structure of the enzyme-substrate complex, termed form A (Table 1). The electron density clearly showed that the crystal contained a mixture of the enzyme bound to either the substrate, 2R3S IGP, or the 2S3S diastereoisomer, which is a by-product of the synthesis and is not a substrate of the enzyme (Hawkes et al., 1995; Saika et al., 1993) (Figure S3). Both diastereoisomers bind at the same site, with the enzyme adopting the same, open conformation equivalent to that of the holoenzyme, including the disordered C loop. The mode of binding of the substrate in the form A complex is comparable to that seen in the 2.1 Å resolution Mycobacterium tuberculosis IGPD-substrate complex (Ahangar et al., 2013), but the substantially increased resolution of the structure described here has enabled a much deeper analysis of how IGPD interacts with its substrate.
In the form A complex, the 2R3S substrate forms a bidentate interaction with Mn1 via the imidazole-N3 atom and the C3-OH group, which replace HOH3 and HOH4 as ligands to the metal ion, respectively (Figures 3A and 3B). Mn1 therefore retains the octahedral coordination seen in the holoenzyme complex. In contrast to Mn1, the second manganese ion, Mn2, is five-coordinate, sharing the same four protein ligands (E77, D108, H145, H73, and H170) and one water ligand (HOH1) with those seen in the holoenzyme structure. However, the position of the imidazole-N3 nitrogen atom of the substrate is such that it would be too close to that of HOH2, the sixth Mn2 ligand in the holoenzyme form. A potential steric clash in the form A complex is avoided by the prior displacement of HOH2 from the enzyme (Figure 3B). The substrate imidazole-N3 nitrogen atom forms a hydrogen bond to HOH1, which in turn is hydrogen bonded to two conserved carboxylic acid groups (E77 and D108). This arrangement implies that the imidazole ring is neutral, with its N3 nitrogen protonated and acting as a hydrogen bond donor, consistent with its high pKa (~14.5) for deprotonation. Compared with the holoenzyme structure, the position of Mn2 is shifted ~0.5 Å out of the plane of the four metal ligands HOH1, O-E77, N-H73, and N-H170 (Figures 3B and 3C). This movement is also accompanied by a shortening of the mean metal-ligand distances by approximately 0.1 Å compared with other octahedrally coordinated Mn(II) ions in complexes of IGPD (Table 2).

The Form B Enzyme-Substrate Complex Traps the Imidazole Form of IGP

A second set of crystallization conditions yielded crystals in the same space group and with equivalent cell dimensions to those observed in form A. However, in the structure determined from these crystals (1.41 Å resolution) (Table 1), termed form B, both the enzyme and the substrate have strikingly different conformations to those seen in form A. In this structure, the C loop is ordered and buries the substrate from solvent, forming a closed conformation of the enzyme, which has not been reported previously (Figure 3D). The position of the C loop in form B occludes the site where an ethylene glycol molecule, originating from the crystallization precipitant, is located in form A. Unlike the situation in form A, where the active site is occupied by a mixture of two diastereoisomers of IGP, in form B the electron density could be explained by the binding of a single molecule of the 2R3S IGP substrate, with only a small number of disconnected, very low-level difference features seen around the substrate binding site (Figure S4). The conformational changes observed in the C loop cause the side chain of S199 to occlude the phosphate binding site observed in form A, resulting in the phosphate group structure of form A therefore shows that IGPD links the binding of its substrate to the generation of a ligand-depleted, five-coordinate Mn(II) ion at the heart of the active site.

Table 1. Data Collection and Refinement Statistics

| Dataset                        | IGP2+Pi (PDB: 4QNK) | IGP2+1,2,4-Triazole (PDB: 4MU0) | E21Q IGP2 Form A + IGP (PDB: 4MU3) | E21Q IGP2 Form B + IGP (PDB: 4MU4) |
|--------------------------------|---------------------|---------------------------------|------------------------------------|------------------------------------|
| **Data Collection**            |                     |                                 |                                    |                                    |
| Beamline                       | Diamond I02         | Diamond I02                     | Diamond I02                        | Diamond I02                        |
| Wavelength (Å)                 | 0.9507              | 0.9686                          | 0.9794                             | 0.9686                             |
| Space group and unit cell      | a = b = c =         |                                 |                                    |                                    |
| parameters (Å)                 | 123                 | 225.13                          | 113.1                              | 112.9                              |
| Molecules per asymmetric unit  | 8                   | 1                               | 1                                  | 1                                  |
| Resolution (Å)                 | 71.19–1.75 (1.84–1.75) | 50.59–1.3 (1.33–1.3) | 46.17–1.12 (1.14–1.12) | 65.18–1.41 (1.45–1.41) |
| Unique observations            | 187811 (27,248)     | 61,097 (4421)                   | 94,992 (6558)                      | 47,547 (3450)                      |
| Rmerge b                       | 0.118 (0.579)       | 0.053 (0.762)                   | 0.041 (0.568)                      | 0.085 (0.849)                      |
| Rfactor/merge c                | 13.3 (22.0)/18.9 (30.1) | 13.0 (23.2)/14.8 (25.2) | 12.2 (23.1)/13.8 (23.0) | 12.4 (18.60)/15.4 (23.1) |
| Average B factors (Å²)         | 20                  | 15                              | 15                                 | 16                                 |
| Bond length rmsd (Å)           | 0.013               | 0.011                           | 0.012                              | 0.012                              |
| Bond angle rmsd (°)            | 1.5                 | 1.5                             | 1.8                                | 1.5                                |
| Multiplicity                   | 4.2 (4.0)           | 12.3 (12.5)                     | 9.5 (4.7)                          | 10.5 (10.8)                        |
| **Refinement**                 |                     |                                 |                                    |                                    |
| No. of non-H atoms             | 13,061              | 1654                            | 1753                               | 1818                               |
| R factor/Rfree (%)             | 13.3 (22.0)/18.9 (30.1) | 13.0 (23.2)/14.8 (25.2) | 12.2 (23.1)/13.8 (23.0) | 12.4 (18.60)/15.4 (23.1) |
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Values in parentheses are data in the highest-resolution shell. Rmsd, root-mean-square deviation.

In the form A complex, the 2R3S substrate forms a bidentate interaction with Mn1 via the imidazole-N1 atom and the C3-OH group, which replace HOH3 and HOH4 as ligands to the metal ion, respectively (Figures 3A and 3B). Mn1 therefore retains the octahedral coordination seen in the holoenzyme complex. In contrast to Mn1, the second manganese ion, Mn2, is five-coordinate, sharing the same four protein ligands (E77, H145, H73, and H170) and one water ligand (HOH1) with those seen in the holoenzyme structure. However, the position of the imidazole-N3 nitrogen atom of the substrate is such that it would be too close to that of HOH2, the sixth Mn2 ligand in the holoenzyme. This potential steric clash in the form A complex is avoided by the prior displacement of HOH2 from the enzyme (Figure 3B). The substrate imidazole-N3 nitrogen atom forms a hydrogen bond to HOH1, which in turn is hydrogen bonded to two conserved carboxylic acid groups (E77 and D108). This arrangement implies that the imidazole ring is neutral, with its N3 nitrogen protonated and acting as a hydrogen bond donor, consistent with its high pKa (~14.5) for deprotonation. Compared with the holoenzyme structure, the position of Mn2 is shifted ~0.5 Å out of the plane of the four metal ligands HOH1, O-E77, N-H73, and N-H170 (Figures 3B and 3C). This movement is also accompanied by a shortening of the mean metal-ligand distances by approximately 0.1 Å compared with other octahedrally coordinated Mn(II) ions in complexes of IGPD (Table 2).
of the IGP being repositioned in a new site, approximately 2.2 Å away from that seen in form A (Figure 4). This forces the substrate to adopt a radically different mode of binding, such that the IGP-phosphate group in form B makes new interactions with R121, S199, and K201, while maintaining those with R99 and K177, albeit with minor conformational changes to these side chains. Moreover, the shift in phosphate position is accompanied by major changes in the torsion angles of the substrate. While these changes do not affect the position of the imidazolate-N1 atom within the coordination sphere of Mn1, they do result in the exchange of the C3-OH for the C2-OH as the second ligand to Mn1 (Figures 3D and 3E). This generates a different geometric arrangement around the metal ion and is accompanied by a shift in the position of the IGP-imidazolate, which in form B sits equidistant between the two metal ions. Thus, the imidazolate-N1 and -N3 nitrogen atoms interact directly with Mn1 and Mn2, respectively, with both of the metal ions octahedrally coordinated and lying in the plane of the imidazolate ring (Figure 3F). This implies that both nitrogen atoms in the imidazolate ring are deprotonated and that the ring is bound as an imidazolate anion, providing the first direct evidence for the existence of the imidazolate as an intermediate in the reaction. Furthermore, the conformation of the substrate is such that the C2, C3, and the imidazolate ring of the IGP lie in approximately the same plane, closely resembling the sp2 geometry of the carbon backbone in the proposed diazafulvene intermediate. The structure of form B therefore shows that when IGPD is in complex with the imidazolate intermediate, its active site is in a closed conformation and is solvent inaccessible, with both metal ions octahedrally coordinated.

As the C loop forms a crucial part of the active site in the form B complex, and since the conserved residues in this region form part of the phosphate binding site (Figure 4), we investigated whether the C loop was required for activity. A C-terminal deletion mutant (residues 170–176) was made in the IGPD homolog from Pyrococcus furiosus (Pf). Activity tests on the wild-type protein, using a modified version of a stopped assay protocol at 65°C (Ames, 1957; Hawkes et al., 1995) gave a specific activity of 2450 m mol IAP mg−1 min−1. In contrast, no activity above background could be recorded for the ΔC-loop IGPD, demonstrating that the C loop is essential for catalysis.

1,2,4-Triazole Mimics the Binding of the Imidazolate

IGPD is inhibited by 1,2,4-triazole and 3-amino-1,2,4-triazole (amitrole) (Hilton et al., 1965; Wiater et al., 1971b), both of which have a chemical structure related to that of an imidazolate anion. To determine whether the mode of binding of 1,2,4-triazole resembles that of the substrate imidazolate seen in the form B structure, we co-crystallized wild-type IGPD2 with 1,2,4-triazole and obtained a 1.3 Å resolution structure of the complex (Figure 2). For clarity. The atom colors, metal interactions, and electron density are shown as in (A). See also Figure S2. As the C loop forms a crucial part of the active site in the form B complex, and since the conserved residues in this region form part of the phosphate binding site (Figure 4), we investigated whether the C loop was required for activity. A C-terminal deletion mutant (residues 170–176) was made in the IGPD homolog from Pyrococcus furiosus (Pf). Activity tests on the wild-type protein, using a modified version of a stopped assay protocol at 65°C (Ames, 1957; Hawkes et al., 1995) gave a specific activity of ~50 μmol IAP mg−1 min−1. In contrast, no activity above background could be recorded for the ΔC-loop IGPD, demonstrating that the C loop is essential for catalysis.
The conformation of the enzyme in this structure was similar to that of the holoenzyme, including the disordered C loop, and a number of water molecules were seen to occupy the available hydrogen bonding positions within the phosphate binding site. Electron density between the two metal ions clearly showed that 1,2,4-triazole binds between the metal ions, with the N1 and N4 nitrogen atoms coordinating Mn1 and Mn2, respectively, replacing HOH3 and HOH2 from the holoenzyme structure (Figure 2C). Both of the manganese ions are therefore octahedrally coordinated and lie in the plane of the triazole ring. 1,2,4-Triazole binds in an equivalent way to that of the imidazolate in the form B complex, suggesting that inhibition of IGPD by 1,2,4-triazole is based on it mimicking the mode of binding of the high-energy imidazolate intermediate.

**DISCUSSION**

**Interconversion between the A and B Forms Reveals Novel Features of the IGPD Reaction Mechanism**

We take the view that the two enzyme-substrate complexes presented here represent two different snapshots of the early steps in the reaction pathway of IGPD. The form A complex demonstrates how the open form of the enzyme can capture the neutral imidazole form of IGP. This is consistent with the expectation that, at biological pH, the neutral form of IGP would predominate over the imidazolate in solution by ~10^2-fold, given the pK_a of ~14.5. In contrast, in form B the enzyme is in a closed conformation due to the ordered C loop. This is associated with the production of an imidazolate anion, the charge on which is critical for the subsequent dehydration step of the reaction, where pi-electron donation leads to loss of the C3 hydroxyl, facilitated by the weak electron-withdrawing nature of Mn(II). The form B complex thus represents a trapped intermediate state of the reaction. The interconversion of the form A and form B structures therefore describes progress through the reaction, in a process that is clearly associated with a number of dramatic torsion angle changes to the IGP, the switch in coordination chemistry at Mn2, and the ordering of the C loop. We presume that these structural changes reflect the molecular processes that dominate the early steps in catalysis, and to investigate the processes that facilitate production of the imidazolate we have analyzed the conformational changes that occur during interconversion between the

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open and closed states of the enzyme-substrate complex (Figure 5 and Movie S1). In the following sections, we analyze the implications of this for the mechanism of the enzyme.

Electrostatic and Kinetic Roles for the Manganese Ions in Catalysis

The structures described here clearly point to two critical electrostatic roles for the manganese ions in facilitating the deprotonation of the imidazole during the first step in catalysis. In the form A structure, where the imidazole-N1 nitrogen atom ligates Mn1 and the N3 nitrogen is hydrogen bonded to HOH1 (Figure 3B), the electron-withdrawing effect of Mn1 is likely to reduce the $pK_a$ of the protonated imidazole-N3 nitrogen atom. This suggests that the way that the substrate interacts with Mn1 during the initial binding phase of the reaction lowers the energy barrier to deprotonation of N3 and formation of the imidazolate. Analysis of the form B structure provides evidence for a second electrostatic role for the manganese ions. In this structure, the mode of binding of the negatively charged imidazolate between the pair of manganese ions results in the metal ions becoming shielded from the repulsion that they exert on each other (Figure 3E). The pair of manganese ions in IGPD therefore acts as a coulombic trap for the negatively charged imidazolate intermediate.

The Five-Coordinate Mn(II) Ion Leads to Weak Substrate Binding in IGPD

In the reaction, the equilibrium between the imidazole/five-coordinate Mn2 ion and the imidazolate/six-coordinate Mn2 ion is presumably significantly shifted compared with the deprotonation of imidazole to imidazolate in water. Although it is not possible to reliably estimate the free energy changes associated with generating a five-coordinate manganese ion, pentacoordinate-Mn complexes have only been observed in metalloenzymes associated with changes in redox state of Mn(II) to Mn(III),

| Table 2. Average Metal-Ligand Distances (Å) |
|---------------------------------------------|
| Structure                        | IGPD2 + Pi | IGPD2+ 1,2,4-Triazole | E21Q IGPD2 Form A + IGP | E21Q IGPD2 Form B + IGP |
|---------------------------------------------|
| Mn1 coordination number                  | 6          | 6                     | 6                        | 6                        |
| HOH3 or equivalenta                     | 2.30 (2.22–2.47) | 2.26               | 2.35                    | 2.30                    |
| HOH4 or equivalentb                     | 2.42 (2.36–2.47) | 2.29               | 2.41                    | 2.33                    |
| OE1 Glu173                              | 2.19 (2.15–2.23) | 2.17               | 2.15                    | 2.16                    |
| NE2 His47                               | 2.27 (2.22–2.29) | 2.29               | 2.25                    | 2.26                    |
| NE2 His169                              | 2.24 (2.20–2.30) | 2.31               | 2.26                    | 2.30                    |
| NE2 His74                               | 2.25 (2.22–2.28) | 2.30               | 2.26                    | 2.30                    |
| Mean                                     | 2.27          | 2.27                 | 2.28                    | 2.28                    |
| Mn2 coordination number                 | 6          | 6                     | 5                       | 6                        |
| HOH1                                    | 2.20 (2.13–2.26) | 2.27               | 2.06                    | 2.19                    |
| HOH2 or equivalentc                     | 2.25 (2.20–2.26) | 2.25               | 2.29                    | –                       |
| NE2 His170                              | 2.24 (2.22–2.27) | 2.27               | 2.28                    | 2.16                    |
| NE2 His145                              | 2.28 (2.22–2.32) | 2.31               | 2.45                    | 1.99                    |
| NE2 His73                               | 2.24 (2.20–2.26) | 2.25               | 2.19                    | 2.19                    |
| NE2 His74                               | 2.22 (2.26–2.35) | 2.30               | 2.33                    | 2.33                    |
| Mean                                     | 2.25          | 2.28                 | 2.26                    | 2.17                    |

*aEquivalents are either the IGP-imidazole, IGP-imidazolate, or 1,2,4-triazole N1 atom.
*bEquivalents are either the C2-OH or C3-OH substituents of IGP.
*cThe ligand distance around Mn2 for the two diastereoisomers of IGP in the form A structure are shown separately.
*dEquivalent is the 1,2,4-triazole N4 atom.
*eMean distances across all eight monomers in the asymmetric unit; range of measured distances is shown in parentheses.
is accompanied by changes to the IGP torsion angles and an exchange of the C3-OH of IGP for the C2-OH as a ligand to Mn1. We have modeled these torsion angle changes to determine the optimal path of rotation (Movie S1). The only movement that avoids potential steric hindrance between the C3-OH and other residues in the active site involves the C2-OH and C3-OH passing each other as they move toward, and away from, Mn1, respectively. During this rotation the C3-OH is transiently perpendicular to the imidazolate ring, maximizing the overlap of its anti-bonding orbital with the pi-electron system of the ring. This is the exact requirement for the geometry of the optimal transition state that leads to the loss of the C3 hydroxyl from the imidazolate intermediate, to form the diazafulvene intermediate (Nasipuri, 1991). When the C3-OH is perpendicular to the imidazolate ring it is within 3.3 Å of one of the carboxylate oxygens of E21 (Figure 5B), which is the proposed acid-base catalyst involved in diazafulvene formation. Furthermore, oscillation of the substrate torsion angles around a form B-like structure would allow this transition state to be accessed at a higher frequency, favoring production of the diazafulvene (Figure 5C).

The Form B Structure Indicates How IGPD Catalyzes the Conversion of the Diazafulvene to IAP
In the next stage of the reaction, the diazafulvene is converted to an enolic intermediate by abstraction of the C2 proton, prior to an enzyme-catalyzed tautomerization that leads to the product, IAP (Glynn et al., 2005). The similarity of the mode of binding of IGP in the form B structure to a diazafulvene allows us to examine conversion to the enol intermediate and the final product, IAP. In form B, the C2-OH of the IGP ligates Mn1 and, thus, the C2-H is rendered more acidic due to its proximity to the metal ion. This would favor the loss of the proton from the diazafulvene during the reaction, in a process that is likely to be catalyzed by E173, the carboxyl of which is immediately adjacent to the C2 (3.5 Å) (Movie S1). This residue is also conveniently placed to participate in the enzyme-catalyzed tautomerization of the enol to IAP.

Switching Coordination Chemistry: Understanding the Role of Specific Metal Ions in Controlling Enzyme Catalysis
The studies described here provide a molecular explanation of the unusual dehydration mechanism catalyzed by IGPD. Central to this is the enzyme-catalyzed formation of a high-energy imidazolate intermediate, without which the reaction rate at pH 7 would be limited by the rate of productive substrate binding. We have demonstrated how changes in manganese coordination chemistry dominate all aspects of catalysis in IGPD. In the first part of this process, the enzyme harnesses the substrate binding energy to create a distorted, ligand-depleted metal center, which serves to remove kinetic barriers to the production of the imidazolate intermediate. Subsequently, a second switch in coordination chemistry restores the octahedral coordination of the metal ion, leading to critical torsion angle changes to the substrate that are essential for concomitant production of the diazafulvene. More widely in biology, the loss of a ligand from an octahedrally coordinated metal ion is not without precedence, for example the changes in coordination state of iron in both soluble...
methylene monoxygenase (Rosenzweig et al., 1993) and hemo-globin (Perutz, 1970). Although five-coordinate Mn(II) centers and changes in coordination states have also been observed in the structures of both manganese superoxide dismutase and oxalate decarboxylase (Tabares et al., 2005, 2009) their roles, unlike IGPD, are associated with changes in metal redox state during the reaction. To our knowledge, the work on IGPD presented here is the first description of how an enzyme can exploit the unique ability of transition metals to switch coordination states at specific steps in catalysis. In IGPD this facilitates a high-ly choreographed and dynamic process that controls the reaction mechanism at every stage. This role of metal ions may well be a more important driver to catalysis than is currently recognized.

**EXPERIMENTAL PROCEDURES**

**Cloning and Mutagenesis**

Two ΔN constructs of the gene encoding for IGPD2 from *A. thaliana* were designed to remove an N-terminal signal peptide and cloned into pET24a vectors (Novagen). Construct A (residues 54–272), was provided as a synthetic gene (GeneArt) and construct B (residues 69–272), which was further truncated at the N terminus, was cloned from the original construct. The E21Q mutant was made by introducing a point mutation to construct B by QuickChange site-directed mutagenesis (Stratagene), subsequently verified by sequencing (GeneServices). The IGPD2 complex with phosphate was determined from crystals grown with construct A, while the 1,2,4-triazole complex was determined using construct B. The gene encoding for IGPD from *Pf* was PCR amplified with a mutation to the start codon, which replaced a GTG with an ATG, before ligation into pETBlue (Novagen). A ΔC construct of the *Pf* enzyme was also cloned from the full-length gene, introducing a premature stop codon, which removed the C-terminal loop region of the protein (residues 170–176). The ΔC construct was subsequently cloned into a pET24a vector for protein expression.

**Expression and Purification**

Plasmids encoding for IGPD2 were transformed into Escherichia coli BL21(DE3) cells (Novagen) and protein expression was induced for 5 hr with 1 mM isopropyl-β-D-thiogalactopyranoside at 37°C in Luria broth, supplemented with 5 mM MnCl₂. Plasmids encoding for the *Pf* IGPD were transformed into Tuner DE3 cells (Novagen) and expressed using the same method, but for 3 hr and supplemented with 4 mM MnCl₂. Cells were harvested and lysed by sonication. For IGPD2, insoluble material was removed by centrifugation and the cell-free extract was purified in a process involving anion-exchange and size-exclusion chromatography, combined with ammonium sulfate precipitation. For *Pf* IGPD, cell-free extract was heated to 70°C for 20 min, before purification using anion-exchange and size-exclusion chromatography. All proteins were analyzed after each stage by SDS-PAGE.

**Crystallization and Data Collection**

Prior to crystallization, substrate (Toronto Research) or inhibitor (Sigma) were added to the protein (10 mg ml⁻¹ in 50 mM Bis-Tris propane buffer [pH 8.0], 50 mM NaCl, 10 mM β-mercaptoethanol, and 0.4 mM MnCl₂) at a final concentration of 5 mM. Initial crystallization conditions were determined by automated screening (Nextal) using a Matrix Hydra II crystallization robot. Crystals were de-

**Figure 5. The Reaction Mechanism of IGPD**

(A) Stereo view to show the initial binding mode of the substrate to IGPD. IGP (green carbons) binds with a neutral imidazole ring, displacing HOH2 and leading to five-coordinate geometry around Mn2 (maroon sphere). The protonated imidazole-N3 atom forms a hydrogen bond (orange dashes) to HOH1 (red sphere), which is also hydrogen bonded to D108 and E77. The protein backbone and side-chain carbon atoms are shown in white; all non-carbon atoms are colored as in Figure 4. E21 is modeled on the basis of its position in the wild-type complexes, and other relevant side chains are shown as sticks, with labels omitted for clarity. Schematics representing the first two steps in the proposed reaction mechanism accompany the stereo views with the five-coordinate manganese ion, Mn2, colored maroon and enclosed in a pentagon. Schematic produced with ChemDraw. See also Movie S1.

(B) Ordering of the C loop (not shown) induces a conformational change, triggering deprotonation of the IGP-imidazole to imidazolate (yellow carbons) and restoring the octahedral coordination of Mn2. This is accompanied by the exchange of the C3-OH for the C2-OH as the ligand to Mn1 and repositioning of the IGP-phosphate. During substrate rearrangement between the form A and form B states, the C3-OH is transiently perpendicular (partially transparent) to the imidazolate, which is required for C3-OH elimination. A double-headed arrow represents oscillation around the intermediate. E21 is within close proximity (3.3 Å) to the C3-OH (pink dashes). Figure is drawn as in (A).

(C) The modeled position of the diazafulvene intermediate (lime green) in which the C3-OH has been eliminated. Figure is drawn as in (A).
were optimized by hanging-drop vapor diffusion using a 5:1 ratio of protein to precipitant, which yielded cubic crystals after 5–10 days when equilibrated against a 1-ml reservoir of precipitant at a temperature of 17°C. Crystals were cryoprotected in their crystallization solution containing 25%–30% ethylene glycol, prior to flash-cooling in liquid nitrogen. All data sets were collected on the MX beamlines at the Diamond Light Source. The two mutant complexes with IGP and the 1,2,4-triazole complex belonged to the space group P432 with cell dimensions of a = 113 Å. The structure with phosphate was determined in the space group I22 with cell dimensions of a = 225 Å. The data collection statistics are summarized in Table 1.

Structure Determination and Refinement
Data were processed using Xia2 (Winter, 2010) or by integration using iMosflm (Leslie and Powell, 2007). All additional data processing was carried out with CCP4i (Winn et al., 2011). The IGPΔ2 complex with phosphate was determined by molecular replacement using PHASER (McCoy et al., 2007) with a monomer of A. thaliana IGPΔ1 (PDB: 2F1D) (Glynn et al., 2005) as a search model (sequence identity ~96%). The 1,2,4-triazole complex was determined by the same method, using the refined protein coordinates from the phosphate complex as a search model. All other structures were directly refined using the protein atoms from the phosphate complex, but omitting ligands, metal ions, and solvent. Model building and refinement was carried out in Coot (Emsley and Cowtan, 2004) and Refmac5 (Murshudov et al., 1997), with ligand libraries generated by JLigand (Lebedev et al., 2012). Refinement statistics are summarized in Table 1. For all structures, no electron density was visible for the residues before S75, which was renumbered S9 for consistency with the structure of IGPD1 from A. thaliana (PDB: 2F1D) (Glynn et al., 2005). Models were validated using MolProbity (Chen et al., 2010) and figures were generated using Pymol (Shrodinger, LLC).

Determining the Kₘ of IGP Binding to IGPD2
The Kₘ was determined using a modified coupled assay (Hawkes et al., 1985). Assay buffer was combined with bovine glutamate dehydrogenase (Sigma) and a large excess of imidazoleacetate-phosphate transaminase (S. Singh, Syngenta). Reduced nicotinamide adenine dinucleotide (Sigma) and IGPD2 were added to a final concentration of 6 mM and 45 nM, respectively, before incubation at 30°C in a temperature-controlled spectrophotometer. When the signal at 340 nm had stabilized, IGP was added at various concentrations to start the reaction. Initial rates from the linear part of the curve were recorded, and Kₘ and Kₙa were determined by non-linear least-squares fitting in GraphPad Prism.

ACCESSION NUMBERS
The atomic coordinates and structure factors have been deposited in the PDB under accession codes PDB: 4MU0, 4MU3, 4MU4, and 4QNK.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, five figures, and one movie and can be found with this article online at http://dx.doi.org/10.1016/j.str.2015.05.012.

AUTHOR CONTRIBUTIONS
C.B. and K.L.B. conducted the work on A. thaliana IGPD2 and, together with D.W.R. and P.J.B., interpreted the data and wrote the manuscript. R.C.V. conducted modeling studies and, together with T.R.H., contributed to discussions on the enzyme mechanism. S.E.S. established the purification procedures for the enzymes; T.C.E. conducted studies on the enzyme mechanism. S.E.S. established the purification procedures for the enzymes; T.C.E. conducted studies on

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Supplemental Information

Crystal Structures Reveal that the Reaction

**Mechanism of Imidazoglycerol-Phosphate Dehydratase**

*Is Controlled by Switching Mn(II) Coordination*

Claudine Bisson, K. Linda Britton, Svetlana E. Sedelnikova, H. Fiona Rodgers, Thomas C. Eadsforth, Russell C. Viner, Tim R. Hawkes, Patrick J. Baker, and David W. Rice
Figure S1. Related to Figure 1. Multiple sequence alignment of IGPD homologues from a range of species.

Examples are from Arabidopsis thaliana (ARATHA2; isoform 2 and ARATHA1; isoform 1), Pisium sativum (PISISAT), Triticum aestivum (TRITAES), Saccharomyces cerevisiae (SACCERV), Burkholderia pseudomallei (BURKPSE), Staphylococcus aureus (STAPAUR), Mycobacterium tuberculosis (MYCOTUB) and Pyrococcus furiosus.
Sequences are numbered according to the full-length polypeptide, including the N-terminal signal peptides for the plant homologues. Black and gray shading indicates identical or similar residues, respectively. Black dots indicate residues that are involved in metal ion binding. Residues implicated in recognition of the substrate-phosphate are marked with a black triangle. Sequences were retrieved from UniProt, the alignment was produced using Tcoffee and the figure was drawn using boxshade.

**Figure S2. Related to Figure 2. The biological assembly of IGPD2.** The 24mer is viewed down the 4-fold axis, chains are represented as cartoons and each is drawn in a different colour. The pair of Mn$^{2+}$ ions in each active site are coloured purple and represented as spheres.
Figure S3. Related to Figure 3A-C. The mode of binding of the 2R3S and 2S3S diastereoisomers of IGP in the form A complex.

(A) Both the substrate, 2R3S IGP, (green) and the contaminant, 2S3S IGP, (gray) form a bidentate interaction with Mn1 via their imidazol(at)e N1 nitrogen and C3-OH atoms. The phosphate groups of both diastereoisomers bind in an equivalent position to each other and to that of the phosphate moiety observed in the holo enzyme complex. 2S3S IGP binds between the metal ions with an imidazolate anion whereas the substrate, 2R3S IGP, binds with a neutral imidazole. This leads to two distinct positions for Mn2, one octahedral (gray), associated with the 2S3S IGP and one square pyramidal (green), associated with the substrate. Important hydrogen bonds and metal ion interactions are shown as orange and black dashes, respectively.

(B) Diagram to show the interactions made between the metal ions and the 2S3S IGP. The imidazole ring is deprotonated, forming an imidazolate anion that interacts with both metal ions, each of which is octahedral.

(C) An imidazolate anion (gray) (0.4 occupancy) and a phosphate anion (occupancy
1.0) were modelled into the electron density map and refined. The resulting difference map (green mesh) revealed a second ring position (inset), which was interpreted as an imidazole based on the pattern of hydrogen bonding. The final position of the substrate (green) is shown partially transparent. The difference map is contoured at 4.0 σ and the 2mFo-DFc map (gray mesh) is contoured at 1.0 σ.

(D) Refinement of the imidazole (green) (0.6 occupancy) and a phosphate anion (1.0 occupancy) revealed difference density for the imidazolate anion that had been modelled previously (inset). Residual density in the difference maps (green mesh) indicated two positions corresponding to two hydroxyl groups due to the different chirality at C2. A steric clash between the 2S position of the hydroxyl and the C4 atom of the neutral imidazole (1.9Å) (black arrow) allowed for the 2R hydroxyl to be matched to the imidazole and the 2S hydroxyl to the imidazolate. The observation of positive difference density around Mn2 in both difference maps indicated that there were two positions for the metal ion. The final position of 2S3S IGP (gray) is shown partially transparent. The difference map is contoured at 4.0 σ and the 2mFo-DFc map (gray mesh) is contoured at 0.9 σ.

(E) The 1.15 Å 2mFo-DFc map (gray mesh) after refinement with 2R3S IGP and 2S3S IGP with the ligands modelled at occupancies of 0.6 and 0.4, respectively, is shown at 1.5 σ.
Figure S4. Related to Figure 3D-E. The mode of binding of 2R3S IGP in the form B complex.

The 1.4 Å 2mFo-DFc electron density map (gray mesh) is shown surrounding the modelled substrate at 1.5 σ. After refinement some low-level density features could be observed in the difference map, suggesting the binding of other minor species, but it was not possible to interpret the density. The difference map is shown in green at 4.5 σ. The IGP is coloured as in Figure 3D-E. The two manganese ions are labelled and shown as purple spheres.
Figure S5. Related to experimental procedure: Determining the Km of IGP binding to IGPD2.

The Km was determined over various IGP concentrations from 125 µM to 2 mM using a coupled assay (Hawkes et al., 1995). After a period of lag, initial rates were recorded and plotted against the substrate concentration. The Vmax was calculated to be 12 nmoles (+/- 2 nmoles) of IGP consumed min⁻¹ and the Km was estimated to be 170 µM (+/- 70 µM). Km and Kcat were determined by non-linear least squares fitting in Graph Pad Prism.

Movie S1. Related to Figure 5. The reaction mechanism of IGPD. A model of the reaction mechanism was produced using Pymol (Schrodinger, LLC) and interpolation between the form A and form B complex structure using LSQMAN (Kleywegt et al., 2001) from the Uppsala crystallography software factory (USF).
### Supplemental Experimental Procedures:

| Primer Name | Sequence 5’>3’ |
|-------------|----------------|
| IGPD2-ΔN_f  | CATATGGAATCAGCTCGAATAGGTGAA |
| IGPD2-ΔN_r  | GGATCCTTATGAACGTGACAAGACTCTCT |
| E21Q_sense  | GAAGAGAGAAACAAACAAACAAATGTATCAGTGAAG |
| E21Q_antisense | CTTCACTTGATACATTTGTTTGCTTTT |
| Pf-IGPD_f   | ATGGAGAGAAACTCAAAAGGAAC |
| Pf-IGPD_r   | TTCAATACCCCTTTTGCT |
| Pf-IGPD-ΔC_f | CATATGAGAGAAACTCAAAAGGAAC |
| Pf-IGPD-ΔC_r | GGATCCTTAAACTCTCCTCTCTCCTTT |

### Construct Design

Two N-terminally truncated constructs of the *HISN5B* gene from *Arabidopsis thaliana* were designed in order to remove the N-terminal signal peptide from the over-expression product. The longer length construct, construct A (residues 54-272), was provided as a synthetic gene (GeneArt) and subsequently ligated into a pET24a (Novagen) expression plasmid using Ndel and BamHI restriction sites. The shorter construct, construct B (residues 69-272), which had a further 16 residues deleted from the N-terminus to completely remove the signal peptide, was cloned from the original construct using the pair of synthetic oligonucleotides, IGPD2-ΔN_f and IGPD2-ΔN_r, before ligation into a pET24a plasmid using Ndel and BamHI restriction sites. The IGPD2 complex with phosphate was determined from crystals grown with construct A, whilst the mutant complexes with substrate and the 1,2,4-triazole complex were determined using construct B.

### Mutagenesis

Template plasmid encoding for construct B was propagated in *Escherichia coli* sub-cloning strain DH5α (Invitrogen) and purified by mini-prep (Qiagen). A pair of synthetic oligonucleotides, E21Q_sense and E21Q_antisense (Eurofins MWG Operon), were used to create an E21Q mutation using the Stratagene QuikChange site directed
mutagenesis procedure. The reaction was carried out in a 25 µl volume using an 18 cycle thermal cycling reaction that incorporated a 12 min elongation step. The template plasmid was then degraded by incubation with DpnI for 1 hr at 30 °C and the resulting plasmid was transformed into XL1-Blue E. coli cells (Stratagene). Plasmid was harvested and sequenced (UoS Geneservice) to validate the mutation.

**Protein over-expression and purification**

Both the wild-type and mutant IGPD2 proteins were over-expressed in E. coli strain BL21 (DE3) (Novagen). After incubation at 37 °C for 5 hr post-addition of 1 mM IPTG and 5mM MnCl₂, cells were harvested by centrifugation, re-suspended in buffer A (50 mM Tris-HCl pH 8 and 50 mM NaCl) and lysed by sonication (2 x 20 s cycles at 16 micron amplitude on a Soniprep 150 sonicator). The lysate was centrifuged at 70000 g for 10 min to pellet the insoluble material and the cell free extract was applied to a 5 ml Hi-Trap DEAE-Sepharose Fast Flow cartridge (GE Healthcare). IGPD2 does not bind to the DEAE matrix under the conditions used, so the column was washed with buffer A and a column-volume of eluted material was retained. Protein was precipitated in 1 M ammonium sulphate solution, pelleted by centrifugation at 40000 g for 5 min and dissolved in buffer B (50 mM Tris-HCl pH 8.0, 0.8 M ammonium sulphate). Hydrophobic chromatography was carried out with a Phenyl-Toyopearl 650S (TOSOH) column or using a 5 ml Hi-Trap Phe-HP cartridge (GE Healthcare) that had been pre-equilibrated with buffer B. The sample was applied and IGPD2 was eluted by a reverse gradient of ammonium sulphate concentration from buffer B to buffer A. Fractions were collected and analysed for protein content by Bradford Assay (Bio-Rad). Peak protein fractions containing IGPD2 were combined and the protein was precipitated in 2 M ammonium sulphate solution. Precipitate was collected by centrifugation at 40000 g for 5 mins and dissolved in 1 ml of buffer C (50 mM Tris-HCl...
pH 8.0 and 0.5 M NaCl). The sample was applied to a 16 x 60 Superose 6 gel filtration column pre-equilibrated in buffer C. Gel filtration was performed at a flow rate 1.5 ml min\(^{-1}\). 2 ml fractions were collected and those with the highest protein concentration were combined. The final samples were concentrated with a 30000 MW cut-off VivaSpin concentrator and buffer exchanged into buffer D (50 mM Bis-Tris propane pH 8.0, 50 mM NaCl, 10 mM β-mercaptoethanol and 0.4 mM MnCl\(_2\)) using a diafiltration cup, then analysed by SDS-PAGE.

**Crystallisation**

Prior to crystallisation, both wild-type IGPD2 and the E21Q mutant were concentrated to 10 mg ml\(^{-1}\) in buffer D. Na\(_2\)HPO\(_4\) or 1,2,4-triazole (Sigma) were added to the wild-type enzyme to a final concentration of 5 mM prior to crystallisation. The same procedure was followed for the E21Q mutant with synthetic IGP (Toronto Research). Initial conditions were determined by automated screening (Nextal) using a Matrix Hydra II crystallisation robot. The conditions were optimized by hanging-drop vapor diffusion using a 5:1 ratio of protein to precipitant, to yield cubic crystals after 5-10 days when equilibrated against a 1 ml reservoir of precipitant at a temperature of 290 K. Crystals of the 1,2,4-triazole complex were grown in optimized hanging-drop experiments from 0.1 M Tris-HCl pH 9.0 and 13 % PEG 200. Crystals of the phosphate complex were grown in sitting-drop experiments from 0.1 M Bicine pH 9.0 and 10% PEG 6000. The E21Q mutant complexes with IGP were crystallized in hanging drop experiments with 25 % ethylene glycol (form A) or 0.1 M Hepes pH 7.5, 0.05 % PEG 2000 and 0.4 M succinic acid pH 7.0 (form B).
Data collection

The wild-type co-crystals with 1,2,4-triazole or phosphate were cryoprotected in their crystallisation buffer containing 25 % ethylene glycol. The form A E21Q IGPD2 crystals were cryoprotected with 30 % ethylene glycol and the form B crystals were cryoprotected in their crystallisation buffer containing 30 % ethylene glycol, prior to cooling in liquid nitrogen. All datasets were collected on the MX beamlines at the Diamond Light Source (Table 1). The two mutant complexes with IGP and the 1,2,4-triazole complex belonged to the spacegroup P432 with cell dimensions of approximately a=113.5 Å and with a monomer in the asymmetric unit. The structure with phosphate was determined to be in the spacegroup I23 with cell dimensions of a=225 Å and eight subunits of IGPD2 in the asymmetric unit. Data collection statistics are summarized in Table 1.

Structure determination and Refinement

Data were processed at the Diamond Light Source using the Xia2 pipeline (Winter, 2010) or by integration using imosflm (Leslie and Powell, 2007). All additional data processing was carried out using CCP4i (Winn et al., 2011). The phosphate complex was determined by molecular replacement using PHASER (McCoy et al., 2007) with a monomer of A. thaliana IGPD1 (PDB: 2F1D) (Glynn et al., 2005) as a search model (sequence identity ~96%). The complex with 1,2,4-triazole was determined by the same method, but using a monomer of IGPD2 from the phosphate complex as the search model. All subsequent structures were determined directly by refinement using protein coordinates from the 1,2,4-triazole complex, but omitting ligands, metal ions and solvent. Model building and refinement was carried out in COOT (Emsley and Cowtan, 2004) and Refmac5 (Murshudov et al., 1997), with ligand libraries generated by JLigand (Lebedev et al., 2012). Refinement statistics are summarized in Table 1.
Models were validated using Molprobity (Chen et al., 2010) and figure were made in Pymol (Schrodinger, 2010). Refinement statistics are summarized in Table 1.

In the case of the form A E21Q complex with substrate, the very high resolution (1.12 Å) of the data meant that there was sufficient evidence to interpret the maps as corresponding to the binding of both the 2R3S (substrate) and 2S3S (contaminant) diastereoisomers of IGP (Figure S3). Initial maps showed that the two ligands share a common position for the phosphate moiety and the C1 and C2 atoms, but since the diastereoisomers differ in chirality at C2, two positions could be observed in the electron density map for the C2-OH. The substrate and 2S3S IGP share the same chirality at C3, consistent with the observation of a single electron density feature representing the C3-OH, which is a ligand to Mn1. The structure was refined cautiously with each of the diastereoisomers of IGP first modeled on their own into the corresponding density. Difference maps subsequently revealed the unmodelled position for the binding of the alternate diastereoisomer. Only one of the ring positions was compatible with the chirality of the substrate, due to a steric clash with the C2-OH, resulting in the substrate and 2S3S IGP being assigned the neutral imidazole and an imidazolate, respectively. Unlike the electron density at Mn1, which was spherical, that at Mn2 was ellipsoidal, suggesting two different metal positions, each associated with the binding of the different diastereoisomers.

**Determining the Km of IGP binding to IGPD2**

The Km was determined using a modified coupled assay (Hawkes et al., 1995). Assay buffer (0.1 M potassium-Hepes pH 7.0, 0.22 M Ammonium chloride and 0.22 M L-glutamate sodium salt) was aliquoted into 10 ml volumes and frozen at -80 °C. Prior to use the assay buffer was defrosted and supplemented with 0.2 mM MnCl₂ and 0.01
mM pyridoxal-5-phosphate, both of which were made up in 0.1 M potassium-Hepes pH 7.0. Stock solutions of NADH and bovine glutamate dehydrogenase (GDH) (both Sigma) were prepared in 0.1 M potassium-Hepes pH 7.0 to 6 mM and 2 mg ml⁻¹, respectively. Imidazoleacetol-phosphate transaminase (IAPT), which had been previously purified and frozen as droplets (S. Singh, Syngenta), was defrosted on ice. IGP, supplied as a salt (Toronto Research), was made up to 0.2 M in potassium-Hepes pH 7.0. Mn-IGPD2 from frozen stocks was diluted 40-fold in assay buffer to a working concentration of 4.2 µM. 650 µl of supplemented assay buffer was combined with 100 µl of GDH, 100 µl of IAPT (large excess) and 50 µl of NADH in a cuvette and incubated at 30 °C in a temperature-controlled spectrophotometer. 10 µl of IGPD2 was added and the signal was recorded at 340 nm until it had stabilized at which point 10 µl of IGP at various concentrations (125 µM to 2 mM) was added to start the reaction. After a lag time of ~120 s, initial rates from the linear part of the curve were recorded. Kₘ and Kₖₐₜ were determined by non-linear least squares fitting in Graph Pad Prism.

**Cloning, expression and purification of the ΔC mutant of Pf IGPD**

The gene encoding IGPD from *Pyrococcus furiosus* was PCR amplified using the synthetic oligonucleotides Pf-IGPD_f and Pf-IGPD_r. These were designed to incorporate a 5' mutation to replace the start codon GTG, commonly observed in Archea, with an ATG, prior to cloning into pETBlue (Novagen). The subsequent plasmid, pIGPDKLB, was transformed into Tuner (DE3) *E. coli* (Novagen) and Pf IGPD was over-expressed at 37 °C for 3 hr post-addition of 1 mM IPTG and 4 mM MnCl₂. A second construct was amplified from pIGPDKLB using a pair of synthetic oligonucleotide primers, Pf-IGPD-ΔC_f and Pf-IGPD-ΔC_r, to introduce a premature stop codon into the 3’ end of the gene encoding *Pf* IGPD, thereby removing the part of
the sequence that encodes for the C-loop. The amplified gene was subsequently cloned into a pET24a vector, producing the plasmid pΔC-IGPDKLB, which was over-expressed using the same method as the original construct. The wild-type protein was purified using an initial 20 min heat step at 70 °C in buffer E (40 mM Tris pH 8.0 and 2 mM EDTA), followed by centrifugation to pellet the cell debris. The supernatant was applied onto a DEAE-sephrose fast flow column (Amersham Biosciences) and eluted on a gradient of 0-0.5 M NaCl in buffer E. The wild-type protein was then concentrated and applied to a Hi-Load Superdex 200 gel filtration column, eluting as the 24mer (~500 kDa) species. Purification of the ΔC mutant followed the same method as the wild-type protein except that the ΔC mutant was precipitated by addition of 2 M ammonium sulphate and pelleted by centrifugation at 40000g for 5 min, before being resuspended in buffer F (40 mM Tris pH 8.0, 2 mM EDTA and 0.5 M NaCl) prior to gel filtration. The ΔC mutant eluted as the trimeric (~60kDa) species, which was reapplied to the gel filtration column in the presence of 2 mM MnCl₂ in order to reassemble it.

Both proteins were concentrated using a 30000 MW cut-off VivaSpin concentrator and buffer exchanged into 0.1 M Hepes pH 7.0 using a diafiltration cup.

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