

5-5-2015

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# Inactivation of Mandelate Racemase by 3-Hydroxypyruvate Reveals a Potential Mechanistic Link between Enzyme Superfamilies

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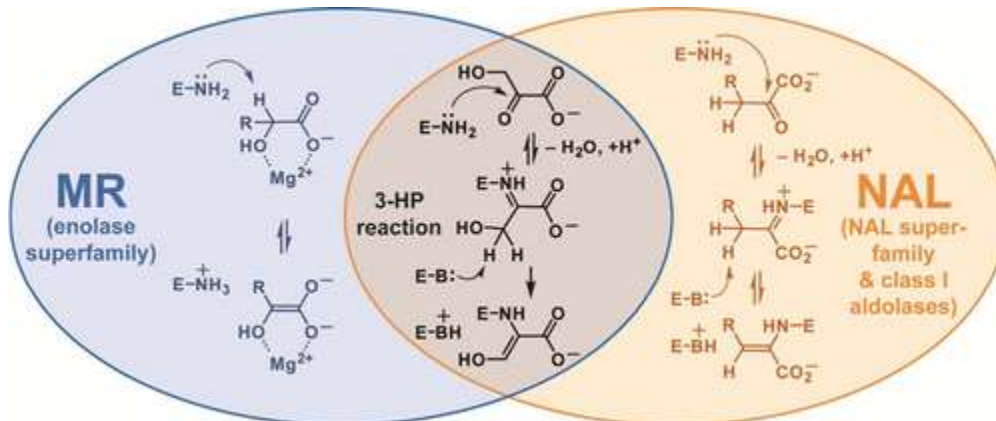
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## Abstract

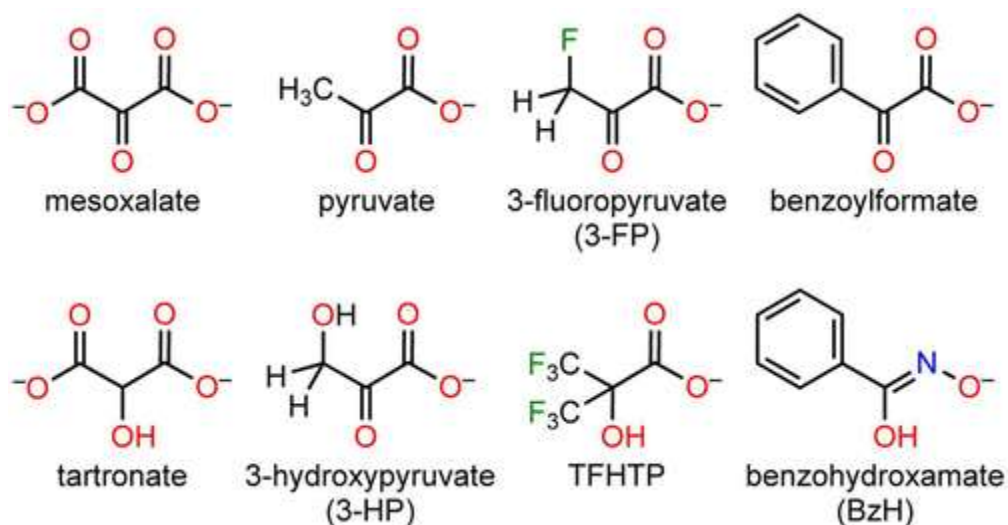


Mandelate racemase (MR), a member of the enolase superfamily, catalyzes the Mg<sup>2+</sup>-dependent interconversion of the enantiomers of mandelate. Several α-keto acids are modest competitive inhibitors of MR [e.g., mesoxalate ( $K_i = 1.8 \pm 0.3$  mM) and 3-fluoropyruvate ( $K_i = 1.3 \pm 0.1$  mM)], but, surprisingly, 3-hydroxypyruvate (3-HP) is an irreversible, time-dependent inhibitor ( $K_{inact}/K_I = 83 \pm 8$  M<sup>-1</sup> s<sup>-1</sup>). Protection from inactivation by the competitive inhibitor benzohydroxamate, trypsinolysis and electrospray ionization tandem mass spectrometry analyses, and X-ray crystallographic studies reveal that 3-HP undergoes Schiff-base formation with Lys 166 at the active site, followed by formation of an aldehyde/enol(ate) adduct. Such a reaction is unprecedented in the enolase superfamily and may be a relic of an activity possessed by a promiscuous progenitor enzyme. The ability of MR to form and deprotonate a Schiff-base intermediate furnishes a previously unrecognized mechanistic link to other α/β-barrel enzymes utilizing Schiff-base chemistry and is in accord with the sequence- and structure-based hypothesis that members of the metal-dependent enolase superfamily and the Schiff-base-forming *N*-acetylneuraminidase lyase superfamily and aldolases share a common ancestor.

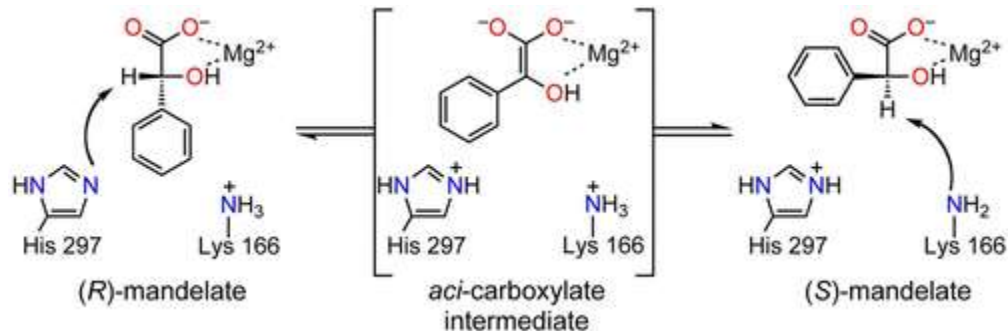
One of the first enzyme superfamilies recognized was the enolase superfamily,<sup>1</sup> whose members share a common partial reaction (i.e., the metal-assisted, Brønsted-base-catalyzed abstraction of the α-proton from a carboxylate substrate to form an enolate)<sup>2-5</sup> and a common protein fold [i.e., a modified (α/β)<sub>8</sub>-barrel (TIM barrel) bearing the catalytic residues at the C-terminal ends of the β-strands and an N-terminal domain that caps the active site conferring substrate specificity and excluding solvent].<sup>6-8</sup> Studies of members of the enolase superfamily have afforded important insights into enzyme evolution and developing strategies for assigning the correct functions to proteins identified in genome projects.<sup>3,9-12</sup> The well-characterized

archetype of this superfamily, mandelate racemase (MR, EC 5.1.2.2) from *Pseudomonas putida*, catalyzes the  $Mg^{2+}$ -dependent, 1,1-proton transfer reaction that interconverts the enantiomers of mandelate and has served as a useful paradigm for understanding how enzymes catalyze the rapid  $\alpha$ -deprotonation of a carbon acid substrate with a relatively high  $pK_a$  value.<sup>13</sup>

MR utilizes a two-base mechanism with Lys 166 and His 297 abstracting the  $\alpha$ -proton from (*S*)- and (*R*)-mandelate, respectively (Scheme 1). A variety of  $\beta/\gamma$ -unsaturated compounds,<sup>14-21</sup> as well as trifluorolactate,<sup>22</sup> serve as substrates for the enzyme. Recently, we showed that the substrate-product analogue 3,3,3-trifluoro-2-hydroxy-2-(trifluoromethyl)propionate [TFHTP (Chart 1)] is a potent competitive inhibitor of MR ( $K_i = 27 \mu M$ ) and that its unexpectedly high binding affinity arises from both intimate interactions between the trifluoromethyl groups and the hydrophobic pocket at the active site, and formation of salt bridges between the carboxylate of TFHTP and the active site Brønsted acid-base catalysts Lys 166 and His 297.<sup>23</sup> Recognizing that these active site bases can be exploited as binding determinants for inhibitors, we showed that tartronate is also a competitive inhibitor of MR ( $K_i = 1.8 \text{ mM}$ ), binding with its glycolate moiety chelating the  $Mg^{2+}$  ion and with the remaining carboxylate group interacting with and bridging the active site Brønsted acid-base catalysts.<sup>23</sup> Our interest in understanding the role of the active site Brønsted acid-base catalysts as binding determinants led us to interrogate the active site using three-carbon  $\alpha$ -keto acids as tartronate analogues. Most unexpectedly, we discovered that 3-hydroxypyruvate (3-HP) is an irreversible inhibitor of MR, reacting at the active site to form a Schiff-base intermediate that is subsequently deprotonated to form an aldehyde/enol(ate) adduct. Such a reaction has not been reported previously for any member of the enolase superfamily and suggests a mechanistic link to other enzyme superfamilies utilizing Schiff-base chemistry.



**Chart 1.** Structures of Ligands



**Scheme 1.** Reaction Catalyzed by Mandelate Racemase

## Materials and Methods

### General

(*R*)-Mandelic acid, lithium 3-hydroxypyruvate (3-HP), sodium mesoxalate monohydrate, benzoylformic acid, sodium pyruvate, benzohydroxamic acid, sodium 3-fluoropyruvate monohydrate (3-FP), and all other reagents, unless mentioned otherwise, were purchased from Sigma-Aldrich Canada Ltd. Circular dichroism (CD)-based assays were conducted using a JASCO J-810 spectropolarimeter with a thermostated, water-jacketed cell holder. High-resolution electrospray ionization (ESI) mass spectrometry (MS) was conducted using a Brüker microTOF Focus orthogonal ESI-time-of-flight mass spectrometer operating in positive ion mode.

## *Concentration Correction for 3-HP*

Elemental analysis (Canadian Microanalytical Services Inc.) was conducted on the lithium salt of 3-HP purchased from Sigma-Aldrich to obtain an accurate estimate of the extent of hydration. On the basis of the presence of 26.13% carbon and 3.63% hydrogen in the sample, the calculated molecular weight of 3-HP was 137.89, which corresponds to  $C_3H_3O_4Li \cdot \sim 1.5H_2O$ .

## *Site-Directed Mutagenesis*

The pET52b(+)-wtMR plasmid<sup>24</sup> was used as the template to create the H297N mandelate racemase (MR) mutant by performing polymerase chain reaction (PCR)-based site-directed mutagenesis using the QuikChange Site-directed Mutagenesis Kit (Stratgene) and following the protocols described by the manufacturer. Polymerase chain reactions were conducted using *Pfu*Turbo DNA polymerase (Bio Basic Inc.). The forward and reverse synthetic deoxyoligonucleotide primers (Integrated DNA Technologies, Inc.) used to construct the mutant were 5'-CCAATGTCCAGCA**A**CCTGTTCCAAGAAATCAGC-3' and 5'-GCTGATTTCTTGGAACAGGT**T**GCTGGACATTGG-3', respectively. The codons specifying the relevant amino acid are underlined, and the altered bases are given in boldface. Potential mutant plasmids were used to transform competent *Escherichia coli* DH5a cells. DH5a cells were used for plasmid maintenance and for all sequencing reactions. The mutant open reading frame was sequenced using commercial automated DNA sequencing (Robarts Research Institute) to ensure that no other alterations of the nucleotide sequence had been introduced.

## *Enzyme Purification*

Recombinant wild-type (wtMR) and H297N MR from *Pseudomonas putida* were overexpressed in and purified from *E. coli* BL21(DE3) cells transformed with the pET-52b(+)-wtMR and pET-52b(+)-H297N-MR plasmids, respectively. The pET-52b(+)-wtMR plasmid containing the MR variant open reading frame encodes the MR gene product (MASWSHPQFEKGALEVLFGQPGYHM<sub>1</sub>...MR) with an N-

terminal StrepII tag (underlined; M<sub>1</sub> represents the first amino acid of wild-type MR).<sup>24</sup> The wild-type and mutant enzymes were purified by affinity chromatography using *Strep*-Tactin Superflow resin (IBA GmbH) and an ÄKTA FPLC instrument (GE Healthcare) as described previously.<sup>24</sup> Upon elution from the column, the enzyme was dialyzed against storage buffer [100 mM HEPES buffer (pH 7.5) containing 3.3 mM MgCl<sub>2</sub>, 200 mM NaCl, and 10% (v/v) glycerol] and stored at -20 °C. The purity of the enzyme preparation (≥99%) was checked using 12% sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS–PAGE). StrepII-tagged MR has the same activity as the wild-type enzyme;<sup>24</sup> consequently, the StrepII tag was not removed from the enzymes.

## Enzyme Assays

MR activity was assayed using a CD-based assay by following the change in ellipticity of mandelate at 262 nm using a thermostated quartz cuvette with a 1-cm light path (unless otherwise indicated) as described by Sharp et al.<sup>25</sup> All assays (total volume of 2 mL) were conducted at 25 °C in Na<sup>+</sup>-HEPES buffer (0.1 M, pH 7.5) containing MgCl<sub>2</sub> (3.3 mM), BSA [0.005% (w/v)], and (*R*)-mandelate as the substrate. The concentration of MR was determined from its absorbance at 280 nm using an extinction coefficient of 53400 M<sup>-1</sup> cm<sup>-1</sup>, which was calculated using the ProtParam tool available on the ExPASy server (<http://web.expasy.org/protparam>).<sup>26</sup> All kinetic parameters were determined in triplicate, and average values are reported. The reported errors are the standard deviations.

## Initial Screening of $\alpha$ -Keto Acids for Time-Dependent Inhibition of MR

Wild-type MR (32.7 nM) was incubated with either pyruvate (10.0  $\mu$ M), mesoxalate (10.0  $\mu$ M), 3-FP (10.0  $\mu$ M), benzoylformate (10.0  $\mu$ M), or 3-HP (8.0  $\mu$ M) in the presence of BSA [0.05% (w/v)] at 25 °C for 0, 6, 12, 18, and 24 min. At each time point, 200  $\mu$ L of the enzyme/inhibitor mixture was added to assay buffer containing (*R*)-mandelate to determine the MR activity. The residual MR activity was measured using the CD-based assay by following the change in ellipticity of mandelate at 262 nm for 3 min in a quartz cuvette with a

1-cm light path (0.5-cm light path length when the inhibition by benzoylformate was assessed). The final concentrations of MR, BSA, inhibitor, and (*R*)-mandelate in the 2.0 mL assay mixture were 3.3 nM, 0.005% (w/v), 1.0  $\mu$ M (except for that of 3-HP, which was 0.8  $\mu$ M), and 10 mM, respectively. Simultaneously, a control reaction mixture containing wild-type MR (32.7 nM) was incubated in assay buffer at 25 °C for 0 and 30 min to determine the rate of loss of MR activity under the incubation conditions in the absence of inhibitor. Similar time-dependent inhibition experiments were also conducted for pyruvate, mesoxalate, benzoylformate, and 3-FP at a greater inhibitor concentration (2 mM) for 0 and 10 min. The final concentrations of MR, inhibitor, and (*R*)-mandelate were equal to 3.3 nM, 0.2 mM, and 10 mM, respectively, in a total reaction volume of 2.0 mL for the activity assays.

### *Inhibition of MR by Mesoxalate and 3-FP*

Inhibition experiments were conducted in Na<sup>+</sup>-HEPES buffer (0.1 M, pH 7.5) containing MgCl<sub>2</sub> (3.3 mM), BSA [0.005% (w/v)], (*R*)-mandelate (0.25–10.0 mM), MR (3.0 nM), and either mesoxalate (0, 1.0, 2.0, and 4.0 mM) or 3-FP (0, 0.75, 1.5, and 3.0 mM) in a total reaction volume of 2.0 mL. The apparent kinetic constants  $V_{\max}$  and  $K_m$  were determined by fitting eq 1 to the initial velocity data using nonlinear regression analysis and KaleidaGraph version 4.02 from Synergy Software. Competitive inhibition constants ( $K_i$ ) were determined from plots of the apparent  $K_m/V_{\max}$  values versus inhibitor concentration in accord with eq 2.

$$v_i = \frac{V_{\max}[S]}{K_m + [S]} \quad (1)$$

$$v_i = \frac{V_{\max}[S]}{K_m \left(1 + \frac{[I]}{K_i}\right) + [S]} \quad (2)$$

### *Inactivation of MR by 3-HP*

The kinetic parameters for the time-dependent inhibition of MR by 3-HP were determined by incubating wild-type MR (32.7 nM) with



various concentrations of 3-HP (0, 3.99, 7.98, 11.97, and 15.95  $\mu\text{M}$ ) in the presence of BSA [0.05% (w/v)] at 25  $^{\circ}\text{C}$  for 0, 6, 12, 18, and 24 min. At each time point, an aliquot (200  $\mu\text{L}$ ) of the reaction mixture was added to the assay solution (total volume of 2.0 mL) containing (*R*)-mandelate (final concentration of 10 mM) to determine the remaining MR activity as described above. The observed pseudo-first-order rate constant for the inactivation of MR ( $k_{\text{obs}}$ ) at each inhibitor concentration was determined by fitting eq 4 (based on eq 3) to the data for the loss of MR activity as a function of time. The values of  $k_{\text{obs}}$  were then plotted against the concentration of 3-HP to determine the efficiency of inactivation ( $k_{\text{inact}}/K_{\text{I}}$ ) using eq 6 (based on eq 5, because  $[3\text{-HP}] \ll K_{\text{I}}$ ). Because saturation was not observed, only the apparent second-order rate constant ( $k_2$  or  $k_{\text{inact}}/K_{\text{I}}$ ) could be determined.

$$\frac{v_i}{v_0} = e^{-k_{\text{obs}}t} \quad (3)$$

$$\ln(\% \text{activity}) = 4.605 - k_{\text{obs}}t \quad (4)$$

$$k_{\text{obs}} = \frac{k_{\text{inact}}[3\text{-HP}]}{K_{\text{I}} + [3\text{-HP}]} \quad (5)$$

$$k_{\text{obs}} \approx \frac{k_{\text{inact}}[3\text{-HP}]}{K_{\text{I}}} = k_2[3\text{-HP}] \quad (6)$$

MR (32.7 nM) that had been completely inactivated by 3-HP (15.95  $\mu\text{M}$ ) was dialyzed for 4.5 and 10.5 h at 4  $^{\circ}\text{C}$  against HEPES assay buffer (pH 7.5) to test the reversibility of the inhibition. After dialysis, no significant MR activity was regained relative to that of a control sample containing wild-type MR in the absence of 3-HP.

### *Inactivation of MR by 3-HP in the Presence of Benzohydroxamate (BzH)*

MR (32.71 nM) was incubated with 3-HP (15.95  $\mu\text{M}$ ) in the presence BSA [0.05% (w/v)] and various concentrations of BzH (0, 5,

10, and 20  $\mu\text{M}$ ) at 25  $^{\circ}\text{C}$ . At various time points (0, 6, 12, 18, and 24 min), an aliquot (200  $\mu\text{L}$ ) of the reaction mixture was added to the assay solution (total volume of 2.0 mL) containing (*R*)-mandelate (final concentration of 10 mM) to determine the amount of MR activity remaining as described above. The observed pseudo-first-order rate constants for the inactivation of MR ( $k_{\text{obs}}$ ) by 3-HP (15.95  $\mu\text{M}$ ) in the presence of various concentrations of BzH were determined by fitting eq 4 to the data for loss of MR activity as a function of time.

### *Mass Spectrometry Analysis of MR Inactivated by 3-HP*

Wild-type MR (0.7 mg/mL) and H297N MR (0.98 mg/mL) were incubated with 3-HP (4.0 mM) at 25  $^{\circ}\text{C}$  until the activity of wild-type MR was undetectable using the CD-based assay [assay concentrations of MR and (*R*)-mandelate of 327 nM and 10 mM, respectively]. A control sample containing wild-type MR was incubated for the same time period in the absence of 3-HP. After complete inactivation of wild-type MR by 3-HP, an aliquot (20  $\mu\text{L}$ ) was removed from each reaction mixture and subjected to SDS-PAGE (12%).

### *In-Gel Digestions*

Gel bands from SDS-PAGE corresponding to MR and H297N MR were excised from the gel and rinsed with deionized water ( $2 \times 300 \mu\text{L}$ ). The excised gel strips were then cut into cubes ( $\sim 1 \text{ mm}^3$ ) and processed using an automated system for conducting in-gel protein digestions (Investigator Progest, Genomic Solutions). Briefly, the protein in the gel cubes was reduced using dithiothreitol (10 mM), carboxamidomethylated using iodoacetamide (100 mM), and digested for 12 h at 37  $^{\circ}\text{C}$  using sequencing-grade trypsin (Promega). The peptides were extracted from the gel pieces by three 20 min incubations with a solution (30  $\mu\text{L}$ ) containing acetonitrile (50%) and formic acid (5%) in LC-MS-grade water with gentle agitation. The extracts were pooled and dried using a vacuum concentrator (Speed Vac Concentrator, SPD 111 V-230, Thermo Electron Corp.) and finally resuspended in LC-MS-grade water (15  $\mu\text{L}$ ) containing acetonitrile (3%) and formic acid (0.5%).

## *LC-ESI-MS/MS and Data Analysis*

LC-ESI-MS/MS was conducted using a nanoflow HPLC system (UltiMate 3000, Dionex) connected to a hybrid ion trap-orbitrap high-resolution tandem mass spectrometer (Velos Pro, Thermo Scientific) operated in data-dependent acquisition (DDA) mode. Each sample (1  $\mu$ L) was injected onto a capillary column (C18 Onyx Monolithic, 0.10 mm  $\times$  150 mm, Phenomenex) at a flow rate of 300 nL/min. Samples were sprayed at 1.6 kV using fused silica noncoated emitters (20  $\mu$ m inside diameter with a 10  $\mu$ m ID tip, PicoTip Emitter from New Objective). Chromatographic separation was achieved using a linear gradient from 3 to 35% solvent B in solvent A over 30 min and then increasing the level of solvent B to 95% over 5 min (solvent A consisted of 0.1% formic acid in water and solvent B 0.1% formic acid in acetonitrile). The raw data files were acquired (Xcalibur, Thermo Fisher) and exported to Proteome Discoverer software version 1.4 (Thermo Fisher) for peptide and protein identification using the Sequest database search algorithm.<sup>27</sup> Carbamidomethyl cysteine and oxidized methionines were selected as static and dynamic modifications, respectively. Peptide to spectrum match (PSM) validation was conducted using high-confidence XCorr threshold values of 1.9, 2.3, and 2.6 for doubly, triply, and more highly charged precursors.

## *Analysis of H297N MR-3-HP Reduced with NaCNBH<sub>3</sub>*

H297N MR (0.98 mg/mL) was incubated with 3-HP (4.0 mM) in assay buffer at 25 °C for 30 min. Sodium cyanoborohydride (10  $\mu$ L of a 1 M stock solution) was then added to the reaction mixture (total volume of 200  $\mu$ L) to give a final concentration of 50 mM, and the pH was adjusted to 7.0 using HCl. The reaction mixture was then incubated for an additional 30 min. Finally, an aliquot of the reaction mixture (20  $\mu$ L) was subjected to SDS-PAGE (12%) and analyzed using mass spectrometry as described above.

## *X-ray Crystallography*

All crystals of wild-type MR were obtained at room temperature by the batch crystallization method. The protein solution consisted of

MR (6.0 mg/mL) purified as described above, in the presence of Na<sup>+</sup>-HEPES buffer (50 mM, pH 7.5) containing MgCl<sub>2</sub> (3.3 mM) and 3-HP (20 mM). The batch reservoir solution was composed of sodium acetate (140 mM, pH 4.5), KCl (280 mM), and pentaerythritol propoxylate 426 (PEP 426; 46%). The protein solution and reservoir solution were gently mixed in a 1:1 ratio to a final volume of 20  $\mu$ L, and 10  $\mu$ L of this mixture was pipetted into a lightly greased crystallization well and sealed with a glass coverslip lined with mineral oil. Crystals grew spontaneously for 17 h, resulting in hexagonally shaped crystals. These crystals were looped directly from the wells to be flash-cooled in liquid nitrogen for storage.

### *Data Collection, Structure Determination, and Refinement*

X-ray diffraction data for MR cocrystallized with 3-HP were collected at the Advanced Photon Source (APS) beamline LS-CAT-21-ID-G on a Rayonix MarMosaic 300 CCD detector, with an X-ray wavelength of 0.979 Å. Diffraction images were processed using HKL2000.<sup>28</sup> The structure was determined by molecular replacement using the wild-type MR enzyme [Protein Data Bank (PDB) 2MNR] as the search model with Phaser.<sup>29</sup> The molecular replacement models were extended by several rounds of manual model building with COOT<sup>30</sup> and refinement with REFMAC<sup>31</sup> using an automatically generated geometric/X-ray weighting term. Water molecules were added to the model in COOT with subsequent manual verification. On the basis of protein purification conditions and the well-established metal binding preference of MR, a Mg<sup>2+</sup> ion was modeled into the active site, despite the fact that a modeled Mg<sup>2+</sup> ion does not satisfy all of the observed electron density at this position.

## **Results and Discussion**

### *Structure–Activity Study*

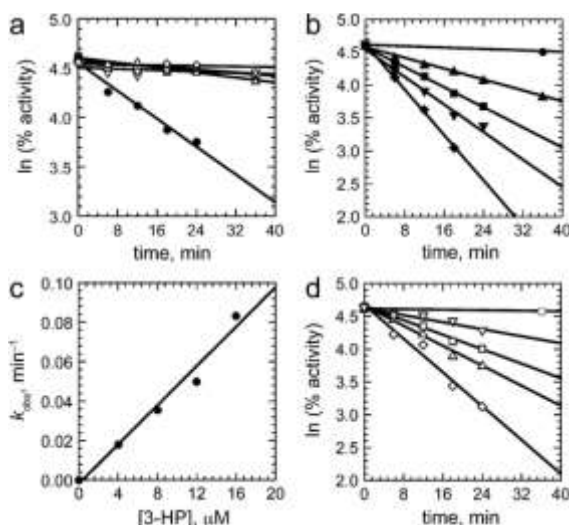
We examined the effect of the  $\alpha$ -keto acids 3-HP, mesoxalate, 3-fluoropyruvate (3-FP), and pyruvate on MR activity (Chart 1). Mesoxalate, which is fully hydrated in water (i.e., C2 *gem*-diol),<sup>32</sup> was a competitive inhibitor of MR with respect to (*R*)-mandelate [ $K_i = 1.8$

$\pm 0.3$  mM (Figure 1S of the Supporting Information)], binding with the same affinity as tartronate ( $K_i = 1.8$  mM),<sup>23</sup> suggesting that the glycolate-like moiety present in both ligands chelates the  $Mg^{2+}$  and the second carboxylate forms a salt bridge with the active site Brønsted acid–base catalysts, thereby compensating for the loss of hydrophobic interactions.<sup>23</sup> Pyruvate, which exists as the keto form under the assay conditions,<sup>33,34</sup> exhibited only 40% inhibition of MR activity at a concentration of 25 mM [i.e.,  $K_i \approx 20$  mM, assuming competitive inhibition with 1 mM (*R*)-mandelate;  $K_m = 1.2$  mM]. This low binding affinity is similar to that of lactate ( $K_i \approx 30$  mM)<sup>20</sup> and likely arises because the methyl group is not sufficiently hydrophobic, nor can it form salt bridges with the Brønsted acid–base catalysts. MR bound 3-FP with an affinity [ $K_i = 1.3 \pm 0.1$  mM (Figure 2S of the Supporting Information)]  $\sim 15$ -fold greater than the affinity it exhibited for pyruvate, due to either dipolar interactions (rather than true H-bonds)<sup>35,36</sup> with the active site acid–base catalysts or enhanced hydrophobic interactions.<sup>37</sup>

In aqueous solutions,  $\alpha$ -keto acids can exist as hydrates (*gem*-diols) to varying degrees.<sup>33</sup> The hydration state will determine the hybridization of C2 and could affect the binding of  $\alpha$ -keto acids at the active site of MR. Pyruvate exists as the keto form (>92%) at pH 6.8,<sup>33,34</sup> and benzoylformate (an  $\alpha$ -keto acid examined previously)<sup>38</sup> is also expected to be in the keto form (>90%) at the assay pH.<sup>39</sup> Our observations that MR binds pyruvate ( $K_i \sim 20$  mM) and benzoylformate ( $K_i = 0.65$  mM)<sup>38</sup> with affinities similar to those for lactate ( $K_i = 30$  mM)<sup>20</sup> and (*R*)-mandelate ( $K_m = 0.81$  mM),<sup>38</sup> respectively, suggest that the enzyme does not distinguish between  $sp^2$  and  $sp^3$  hybridization at C2 in the ground state. Mesoxalate and 3-FP are 100% and >90% hydrated, respectively, in aqueous solutions.<sup>32,40-42</sup> However, in the active site of MR, it is possible that coordination of the metal ion could lead to dehydration of the *gem*-diols, yielding the corresponding keto species.<sup>43</sup> It is unlikely, however, that this latter possibility would result in a significant alteration of the binding affinity considering our observations with pyruvate and lactate, and benzoylformate and mandelate, mentioned above.

## Inactivation by 3-HP

Surprisingly, 3-HP (8.0  $\mu\text{M}$ ) showed time-dependent inhibition, while the other  $\alpha$ -keto acids did not show such behavior at a concentration of 10  $\mu\text{M}$  (Figure 1a) or 2 mM (Figure 3S of the Supporting Information). Dialysis of the inactive 3-HP-treated MR against assay buffer for  $\sim 11$  h did not restore enzymatic activity, indicating that the inactivation was irreversible. The inactivation of MR by 3-HP followed pseudo-first-order kinetics (Figure 1b), and the observed pseudo-first-order rate constants ( $k_{\text{obs}}$ ) exhibited a linear dependence on the concentration of 3-HP (Figure 1c). Saturation was not observed at 3-HP concentrations of  $\leq 16$   $\mu\text{M}$ , and at higher concentrations of 3-HP, we were not able to obtain accurate  $k_{\text{obs}}$  values because the loss of the MR activity was too rapid (relative to the assay duration of 3 min). Consequently, we were not able to differentiate between a kinetic mechanism for inactivation following a single step and a two-step mechanism with the reversible formation of a binary complex between MR and 3-HP. The slope from the plot of  $k_{\text{obs}}$  versus the concentration of 3-HP gave an apparent second-order rate constant for inactivation ( $k_{\text{inact}}/K_{\text{I}}$  or  $k_2$ ) with a value of  $83 \pm 8 \text{ M}^{-1} \text{ s}^{-1}$ . The observed rate of inactivation by 3-HP was reduced in the presence of the reversible competitive inhibitor benzohydroxamate (BzH) (Figure 1d), indicating that 3-HP likely binds at the active site.



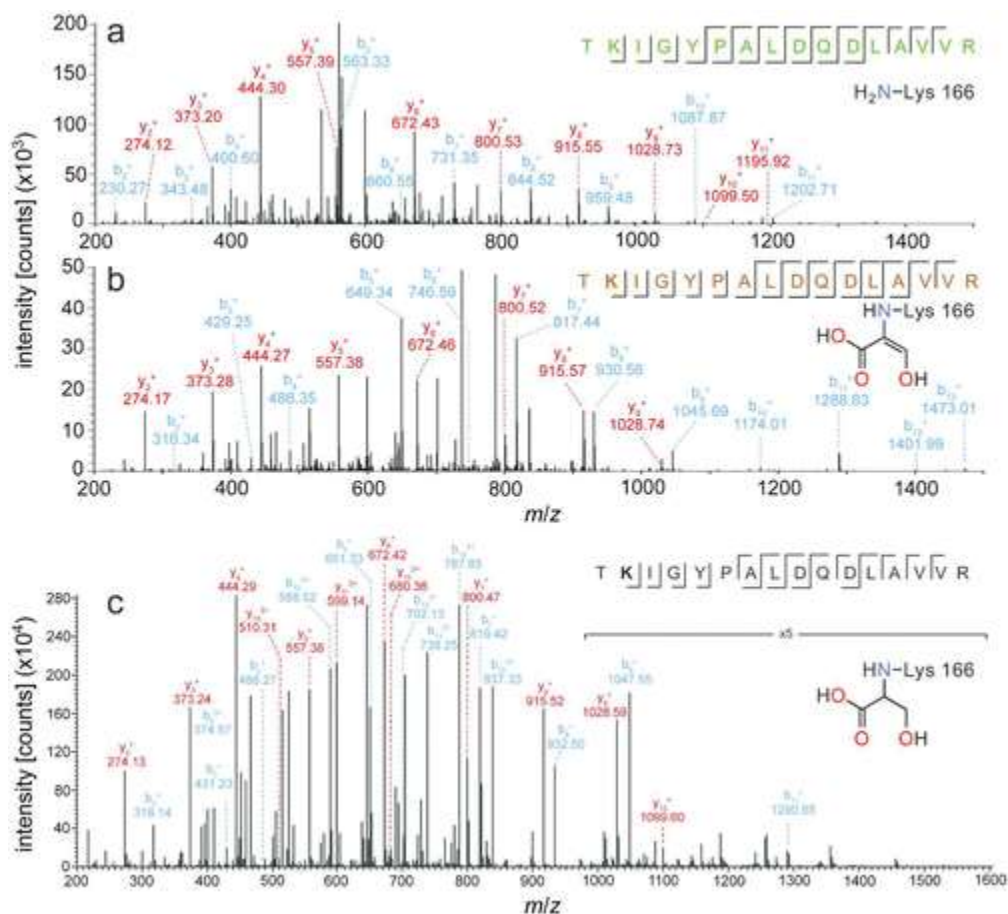
**Figure 1.** Time-dependent inhibition of MR activity. (a) Time-dependent inactivation of MR (32.7 nM) incubated with either 3-HP (8.0  $\mu\text{M}$ , ●), mesoxalate (10  $\mu\text{M}$ , ▽), pyruvate (10  $\mu\text{M}$ , □), benzoylformate (10  $\mu\text{M}$ , Δ), 3-FP (10  $\mu\text{M}$ , ◇), or no inhibitor (○)

at 25 °C. The reaction mixture was diluted 10-fold by addition of assay buffer containing (*R*)-mandelate (final concentration of 10 mM), and the initial rates were measured at the indicated times. (b) Time-dependent inactivation of MR (32.7 nM) incubated with various concentrations of 3-HP: 0 (●), 4.0 (▲), 8.0 (■), 12.0 (▼), and 16.0 (◆) μM at 25 °C. The initial rates were measured at the indicated time points. (c) Observed pseudo-first-order rate constants for inactivation ( $k_{\text{obs}}$ ) plotted as a function of 3-HP concentration. The slope yields a value for the apparent second-order rate constant for the efficiency of inactivation ( $k_{\text{inact}}/K_i$ ) equal to  $83 \pm 8 \text{ M}^{-1} \text{ s}^{-1}$ . (d) Protection against 3-HP-dependent inactivation of MR by 0 (◇), 5 (Δ), 10 (□), and 20 (▽) μM benzohydroxamate at 25 °C. MR was incubated with 3-HP (16.0 μM), and the initial rates were measured at the indicated time points as described above. The value for the control reaction, in the absence of 3-HP, is also shown (○). Experimental details may be found in Materials and Methods.

## Site of Covalent Modification

To identify the residue(s) modified upon inactivation by 3-HP, 3-HP-modified MR (3-HP-MR) and an untreated sample of MR (control-MR) were subjected to SDS-PAGE followed by in-gel digestion with trypsin. The resulting proteolytic cleavage peptides were analyzed using LC-ESI-MS/MS. For both control-MR and 3-HP-MR, ≥95% sequence coverage was obtained. MS data showed identical peptides in both the control-MR and 3-HP-MR samples with the exception of the cleavage fragment TKIGYPALDQDLAVVR (TK-peptide, residues 165–180 with one missed cleavage site). The TK-peptide in the 3-HP-MR sample gave two *m/z* peaks. One peak corresponded to an unmodified peptide ( $\text{MH}^+$ , *m/z* 1758.97) that was also observed in control-MR, suggesting that either the enzyme was partially modified or some modification was reversed during sample preparation. The other TK-peptide had a higher *m/z* value ( $\text{MH}^+$ , *m/z* 1844.98), consistent with MR being modified by addition of a group with a mass of 86 Da. Internal fragmentation of the triply charged modified TK-peptide (monoisotopic, *m/z* 615.66) and the triply charged unmodified TK-peptide (monoisotopic, *m/z* 586.99) precursors gave a mass increment of 86 Da between the  $\text{b}_2^+$  ions arising from the modified (316.34 Da) and unmodified (230.27 Da) TK-peptides, suggesting modification of either Thr 165 or Lys 166 (Figure 2a,b). Modification of Thr 165 seemed unlikely because the side chain of this residue is directed away from the active site, and it is difficult to account for the increment of 86 Da. The remaining b and y ions corresponded to the expected amino acid sequence of the TK-peptides. Upon treatment of H297N MR with 3-HP and subsequent LC-ESI-MS/MS analysis, no modified tryptic peptides were observed, suggesting that His 297 is required for

irreversible inhibition. Thus, our kinetic and MS/MS results suggest that 3-HP modifies Lys 166, with the resulting adduct having a molecular mass of 86 Da.



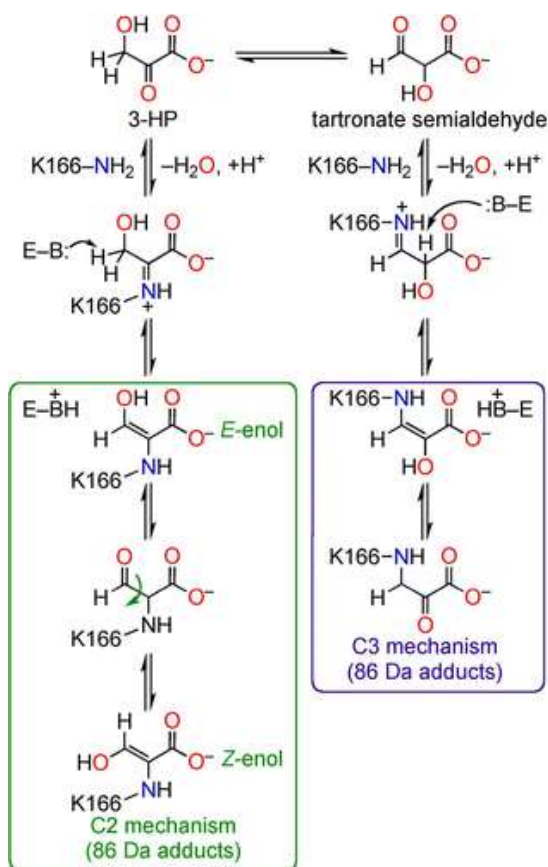
**Figure 2.** ESI-MS/MS analysis of TK-peptides. Mass spectra for the triply charged ion of the TK-peptide from (a) untreated wild-type MR ( $m/z$  586.995;  $MH^+$  = 1758.971 Da) and (b) 3-HP-treated wild-type MR ( $m/z$  615.664;  $MH^+$  = 1844.976 Da) reveal modification of the tryptic peptide (with one missed cleavage site), increasing the mass by 86 Da. The increase in mass of the  $b_2^+$  ion by 86 Da suggests modification of Lys 166 (bold in panel b). (c) ESI-MS/MS analysis of the triply charged ion ( $m/z$  616.329;  $MH^+$  = 1846.97 Da) of the TK-peptide (residues 165–180) arising from the trypsin-digested 3-HP-modified H297N MR treated with NaCNBH<sub>3</sub> reveals modification of the active site Lys 166, increasing the mass by 88 Da. This mass is consistent with reduction of the Schiff-base adduct.

## Mechanism of Inactivation

A possible mechanism accounting for our observations involves the reversible formation of a Schiff base (imine) between the  $\epsilon$ -NH<sub>2</sub> group of Lys 166 and the  $\alpha$ -carbonyl of 3-HP [C2 mechanism (Scheme



2)], similar to the mechanism proposed for the inactivation of *N*-acetylneuraminase lyase (NAL)<sup>44</sup> and dihydrodipicolinate synthase (DHDPS)<sup>45</sup> by 3-HP. Subsequent deprotonation of the Schiff base (presumably by His 297) would yield an enol or enolate [i.e., enol(ate)] (86 Da), which could tautomerize to yield a covalent aldehyde adduct. Incubation of H297N MR with 3-HP, followed by treatment with NaCNBH<sub>3</sub>, and subsequent separation by SDS-PAGE and analysis using LC-ESI-MS/MS revealed that Lys 166 of the TK-peptide was modified by a group with a mass of 88 Da (Figure 2c), consistent with reduction of the Schiff base and in accord with the proposed inactivation mechanism. In addition, we observed two 88 Da adducts arising from the nonspecific modification of Lys 207 and Lys 238 located away from the dimer interface on the surface of the enzyme (data not shown). Such adducts would be expected considering that imines may form spontaneously at other sites bearing a primary amino group.<sup>46,47</sup>

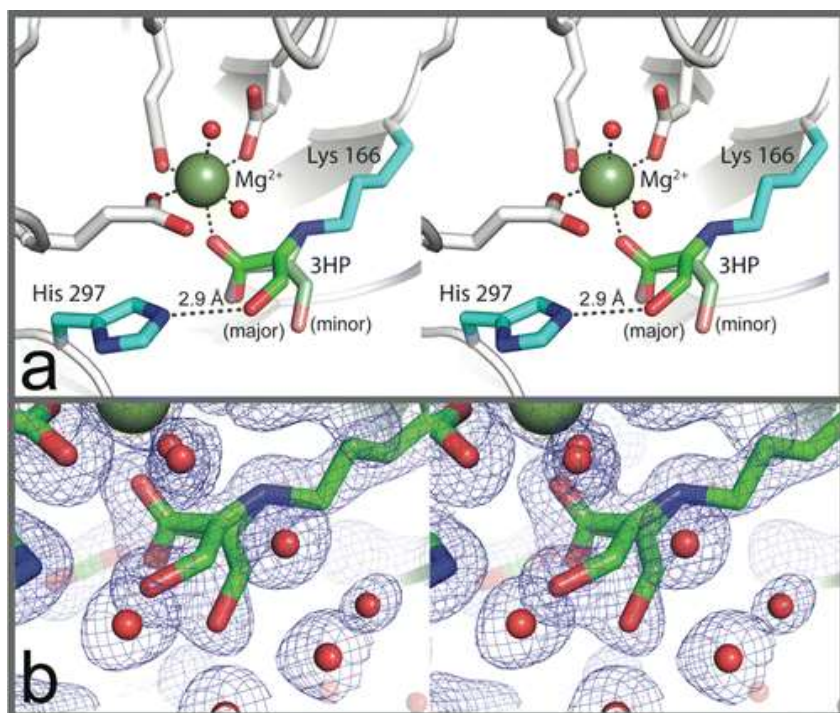


**Scheme 2.** Proposed Mechanisms for Inactivation of MR by 3-HP Giving Rise to an 86 Da Adduct

Alternatively, 3-HP may undergo tautomerization either in solution or within the active site to yield tartronate semialdehyde.<sup>48</sup> Schiff-base formation between Lys 166 and the aldehyde [C3 mechanism (Scheme 2)], followed by deprotonation, also yields an adduct with a mass of 86 Da and is consistent with our LC-ESI-MS/MS observations.

## *Structural Studies*

To distinguish between these two mechanistic possibilities, we determined the X-ray crystal structure of MR inactivated by 3-HP at 1.65 Å resolution (Table 1). The electron density clearly indicates bond formation occurs between the  $\epsilon$ -NH<sub>2</sub> group of Lys 166 and C2 of 3-HP in accord with the C2 mechanism (Figure 3 and Figure 4S of the Supporting Information). The structure was refined by defining a single bond between the  $\epsilon$ -NH<sub>2</sub> group of Lys 166 and C2 of 3-HP, and the 3-HP adduct was further refined in two alternative conformations: major (70%) and minor (30%) conformations of either the (*E*)-enol(ate) or the aldehyde. Although it is not possible to determine whether the product adduct exists as the aldehyde or enol(ate) tautomer from the crystal structure, the enol(ate) may be preferred because enolase, which has a similar active site architecture, is strongly inhibited by the enolate of d-tartronate semialdehyde phosphate.<sup>49,50</sup> Certainly, the active site of MR is optimized to stabilize an enolate in accord with its ability to catalyze racemization, and N $\epsilon$ 2 of His 297 is 2.9 Å from the terminal oxygen of the major adduct so that it is capable of stabilizing the enol(ate) through H-bonding (Figure 3). The inability of 3-FP to inactivate MR indicates that the formation of either the enol(ate) or aldehyde by 3-HP is essential for inactivation.



**Figure 3.** X-ray structure of 3-HP-MR. (a) Stereoview of the active site showing the major (70%) and minor (30%) conformations of the 3-HP adduct with Lys 166. The aldehyde/enol(ate) oxygen of the major conformation is within H-bonding distance of the Brønsted acid–base catalyst His 297. (b) Stereoview of representative  $2F_o - F_c$  electron density maps, contoured at  $1.0\sigma$ , for the 3-HP-MR structure showing a close-up view of the Lys 166–3-HP adduct.

**Table 1.** X-ray Data Collection and Refinement Statistics

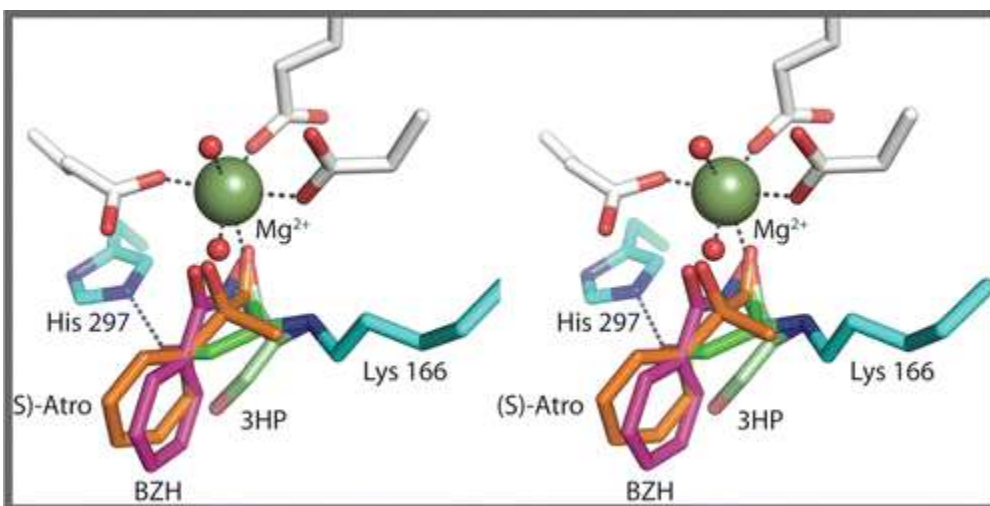
	<b>3-HP-MR</b>
PDB entry	4X2P
space group	<i>I</i> 422
cell dimensions	
<i>a</i> , <i>b</i> , <i>c</i> (Å)	123.88, 123.88, 105.30
$\alpha$ , $\beta$ , $\gamma$ (deg)	90, 90, 90
resolution range (Å)	50.0–1.65 (1.68–1.65) <sup>a</sup>
redundancy	13.0 (8.4) <sup>a</sup>
completeness (%)	99.6 (94.0) <sup>a</sup>
no. of unique reflections	49040
$R_{\text{merge}}$	0.082 (0.437) <sup>a</sup>
average $I/\sigma$	29.55 (4.125) <sup>a</sup>
	Refinement
resolution range (Å)	40–1.65 (1.69–1.65) <sup>a</sup>
$R_{\text{cryst}}$	0.145 (0.181) <sup>a</sup>
$R_{\text{free}}$	0.177 (0.232) <sup>a</sup>
no. of protein atoms	2757

### 3-HP-MR

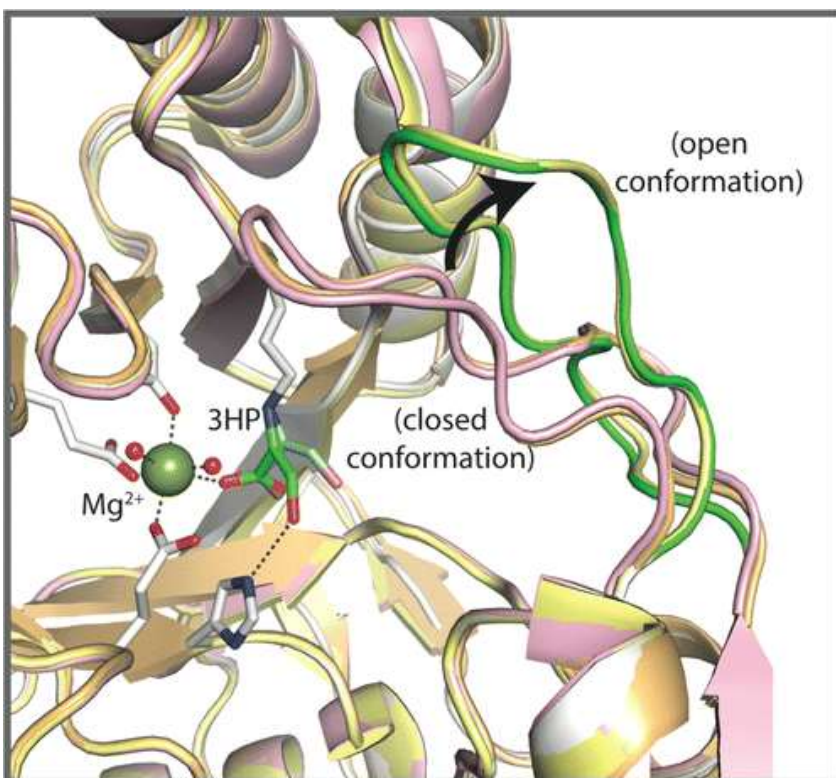
no. of water molecules	297
Wilson <i>B</i> value (Å <sup>2</sup> )	17.0
average <i>B</i> factor (Å <sup>2</sup> )	
protein	16.1
ligand	23.9
Mg <sup>2+</sup>	2.3
solvent	26.1
Ramachandran plot (%)	
most favored	91.5
additionally allowed	7.5
generously allowed	0.7
disallowed	0.3
root-mean-square deviations	
bond lengths (Å)	0.028
bond angles (deg)	2.54

<sup>a</sup>Values in parentheses are for the highest-resolution bin.

C2 of the 3-HP enamine occupies a position approximately equivalent to that of the  $\alpha$ -methyl group of (*S*)-atrolactate (Figure 4). Both ground state and transition state analogues bind to MR with one carboxylate oxygen and the  $\alpha$ -OH group coordinated to Mg<sup>2+</sup> in the active site. In the 3-HP-MR structure, however, the  $\alpha$ -OH group is replaced with a free water molecule, while one carboxylate oxygen maintains coordination with Mg<sup>2+</sup> in a manner analogous to that of the substrate and transition state analogues. This coordination with the active site Mg<sup>2+</sup> likely facilitates deprotonation of the Schiff base by His 297, similar to its role in stabilizing the *aci*-carboxylate intermediate during the racemization of mandelate. Interestingly, the active site lid (20s loop) that typically closes over the active site when either ground state or transition state analogues are bound adopts an “open” conformation in the 3-HP-MR structure (Figure 5). This conformation may arise because the hydrophobic group (i.e., aromatic ring) present on substrates and intermediate analogues is not present to drive lid closure.<sup>51</sup> The 20s loop in the structure of MR (PDB entry 4M6U) with bound tartronate, which also lacks the phenyl ring, exhibited an approximately 1:1 ratio of both the open and closed conformations;<sup>23</sup> however, it is important to note that the 20s loop of MR with only sulfate bound (PDB entry 2MNR) was closed, suggesting that other factors may also contribute to the observed conformation of the loop in crystals.<sup>8</sup>



**Figure 4.** Stereoview of the active site of MR containing the adduct with 3-HP (green), overlaid with the bound substrate analogue (*S*)-atrolactate [(*S*)-Atro; PDB entry 1MDR, orange],<sup>(71)</sup> and the intermediate analogue benzohydroxamate (BzH; PDB entry 3UXK, pink).<sup>(72)</sup> The Brønsted acid–base catalysts (His 297 and Lys 166) are colored cyan.



**Figure 5.** Structural overlay of the 3-HP–MR structure (colored white, with the 20s loop colored green) with the structures of MR in the presence of (*S*)-atrolactate (PDB entry 1MDR, orange),<sup>71</sup> benzohydroxamate (PDB entry 3UXK, pink),<sup>72</sup> and benzilate

(PDB entry 4HNC, yellow).<sup>23</sup> Aside from 3-HP, all other ligands are not shown for the sake of clarity. The 3-HP-MR structure exhibits an "open" conformation of the 20s loop lid, similar to the conformation observed when the bulky substrate-product analogue benzilate is bound to C92S/C264S/K166C MR.<sup>23</sup>

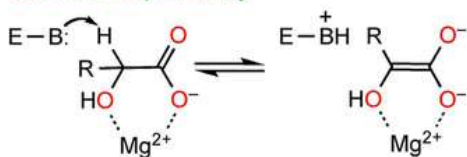
## *Mechanistic Link between Superfamilies*

The formation of a Schiff base is unprecedented in the enolase superfamily. However, irreversible inhibition by 3-HP has been demonstrated for members of the NAL superfamily. This superfamily includes NAL, DHDPS, *trans*-*o*-hydroxybenzylidene-pyruvate hydratase-aldolase, and d-4-deoxy-5-oxoglucarate dehydratase,<sup>2,7,52</sup> which have also been included within the class I aldolases<sup>53</sup> by the SCOP database<sup>54,55</sup> on the basis of their similar Schiff-base chemistry and common ( $\alpha/\beta$ )<sub>8</sub>-fold. 3-HP inactivates NAL from *Clostridium fringens*<sup>56</sup> and *E. coli*<sup>44</sup> and DHDPS from *E. coli*,<sup>45</sup> and the X-ray crystal structures of the covalently modified NAL and DHDPS enzymes revealed the presence of an aldehyde/enol(ate)-lysine adduct similar to the one we observe for MR inactivated by 3-HP.<sup>44,45</sup> Although the mechanism-based inhibition of these two NAL superfamily members by 3-HP also proceeds via initial Schiff-base formation followed by deprotonation,<sup>44,45</sup> the apparent second-order rate constant for the inactivation of MR exceeds that observed for inactivation of DHDPS ( $0.56 \text{ M}^{-1} \text{ s}^{-1}$ ) by ~138-fold. This observation is in accord with the active site of MR being optimized to promote enolate formation.

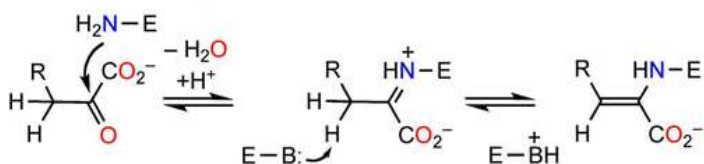
Many of the enzymes sharing the highly prevalent ( $\alpha/\beta$ )<sub>8</sub>-barrel (or TIM barrel) folding topology<sup>53,57-62</sup> are believed to have arisen via divergent evolution from a common ancestor or a limited number of progenitor enzymes.<sup>53,60,63-67</sup> Many aldolases share this fold with members of the enolase and NAL superfamilies, as well as sharing a similar mechanistic theme, i.e., the formation of a "masked" carbanionic intermediate (Scheme 3). Formation of such an intermediate is facilitated via formation of an imine/enamine "electron sink" with an active site lysine residue in the class I aldolases, and by utilizing a transition metal ion (usually  $\text{Zn}^{2+}$ ) as an essential Lewis acid cofactor in the class II aldolases. Although the reaction mechanisms differ between these two classes of aldolases, their ancestral aldolase has been postulated to have been a Schiff-base-forming enzyme that subsequently gave rise to a lineage of metal-dependent class II

aldolases.<sup>53</sup> Several comparisons of sequences, structures, and functions have suggested common ancestry between the class I and II aldolases and the enolase superfamily.<sup>53,57,63,67</sup> Considering the metal ion dependence of the class II aldolases, the mechanistic link to the enolase superfamily was clear (Scheme 3),<sup>68</sup> but such a link between the class I aldolases and the enolase superfamily was not evident. Allen and co-workers conducted a comparison of the primary and tertiary structures of class I aldolases and members of the NAL superfamily, and concluded that these enzymes could all be included in a large superfamily utilizing Schiff-base chemistry and related by divergent evolution from a common progenitor enzyme.<sup>69</sup> Interestingly, these authors employed MR as a “negative control” because it was deemed not to be mechanistically related to the aldolases. This study suggests that a mechanistic relationship does indeed exist. Thus, while over evolutionary time scales the catalytic mechanism of enzymes may not be fully conserved,<sup>67</sup> the ability of MR to form and deprotonate a Schiff-base intermediate furnishes a mechanistic link to other  $\alpha/\beta$ -barrel enzymes utilizing Schiff-base chemistry and may be a relic of an activity possessed by a promiscuous progenitor enzyme.<sup>70</sup> Delineating such chemistry-based relationships and their overlap with structure space and sequence space affords a more complete understanding of nature's strategies for the evolution of new catalysts.<sup>3</sup>

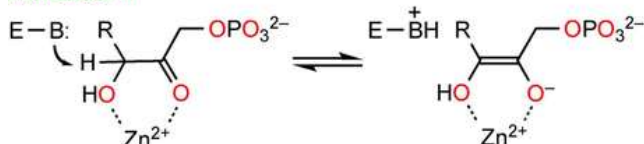
#### Enolase Superfamily



#### NAL Superfamily / Aldolase I



#### Aldolase II



**Scheme 3.** Summary of the Reactions Catalyzed by the Enolase and NAL Superfamilies and the Class I and II Aldolases

## Accession Codes

The atomic coordinates of the 3-HP–MR structure have been deposited in the Protein Data Bank as entry 4X2P.

## Funding Information

This work was supported by a Discovery Grant from the Natural Sciences and Engineering Research Council (NSERC) of Canada (S.L.B.) and National Science Foundation MRI-R2 Grant DBI-0959442 (M.St.M.). Use of the Advanced Photon Source was supported by U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences, under Contract DE-AC02-06CH11357. Use of LS-CAT Sector 21 was supported by the Michigan Economic Development Corp. and the Michigan Technology Tri-Corridor for the support of this research program (Grant 085P1000817). B.N.W. was supported by a GAANN award (Graduate Assistance in Areas of National Need) from the U.S. Department of Education.

The authors declare no competing financial interest.

## Acknowledgment

We thank Alejandro Cohen of the Dalhousie Faculty of Medicine Proteomics Core Facility for his assistance in conducting and analyzing LC–ESI-MS/MS data.

## Abbreviations

BzH	benzohydroxamate
CD	circular dichroism
DHDPS	dihydrodipicolinate synthase
ESI	electrospray ionization
3-FP	3-fluoropyruvate
3-HP	3-hydroxypyruvate
3-HP–MR	wild-type MR modified by 3-HP
LC	liquid chromatography
MR	mandelate racemase
MS	mass spectrometry



NAL        *N*-acetylneuraminase lyase  
TFHTP     3,3,3-trifluoro-2-hydroxy-2-(trifluoromethyl)propanoate.

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## Supporting Information

Inhibition by  $\alpha$ -keto acids (Figures 1S–4S). This material is available free of charge via the Internet at <http://pubs.acs.org>.

# Supporting Information

for

## **Inactivation of Mandelate Racemase by 3-Hydroxypyruvate Reveals a Potential Mechanistic Link Between Enzyme Superfamilies**

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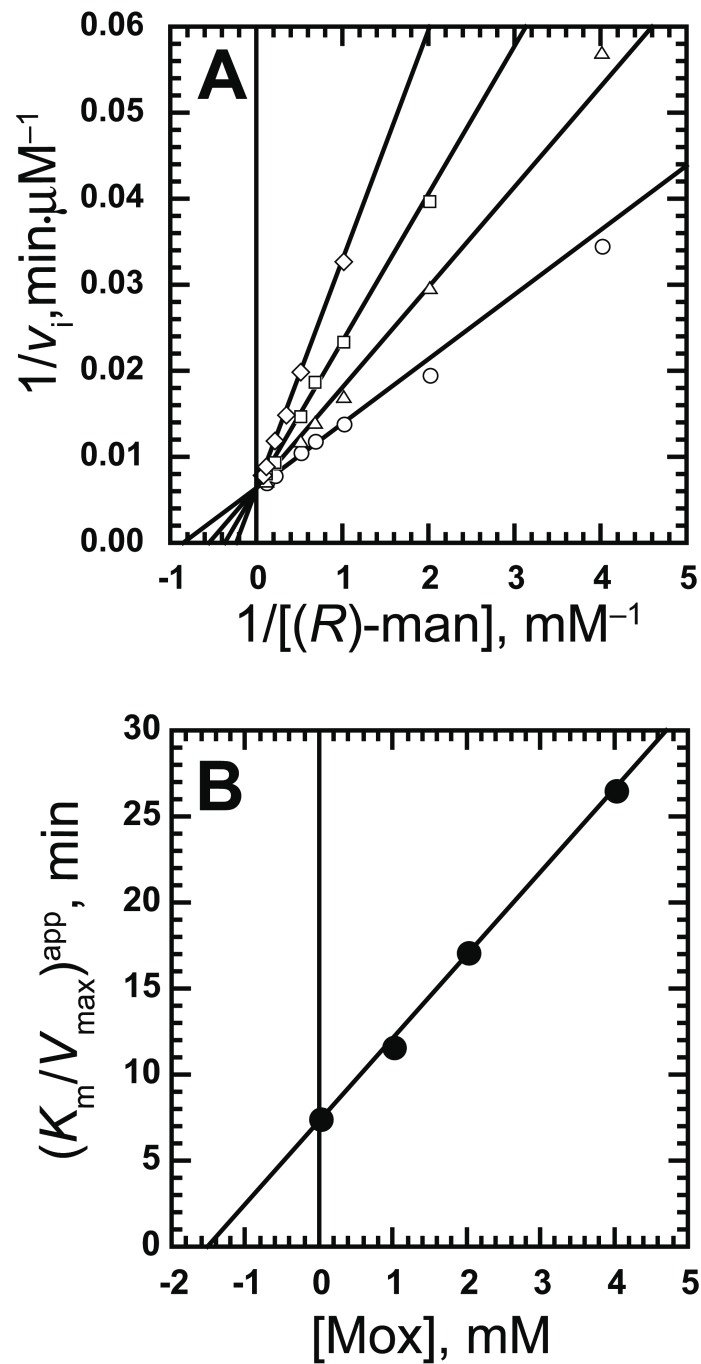
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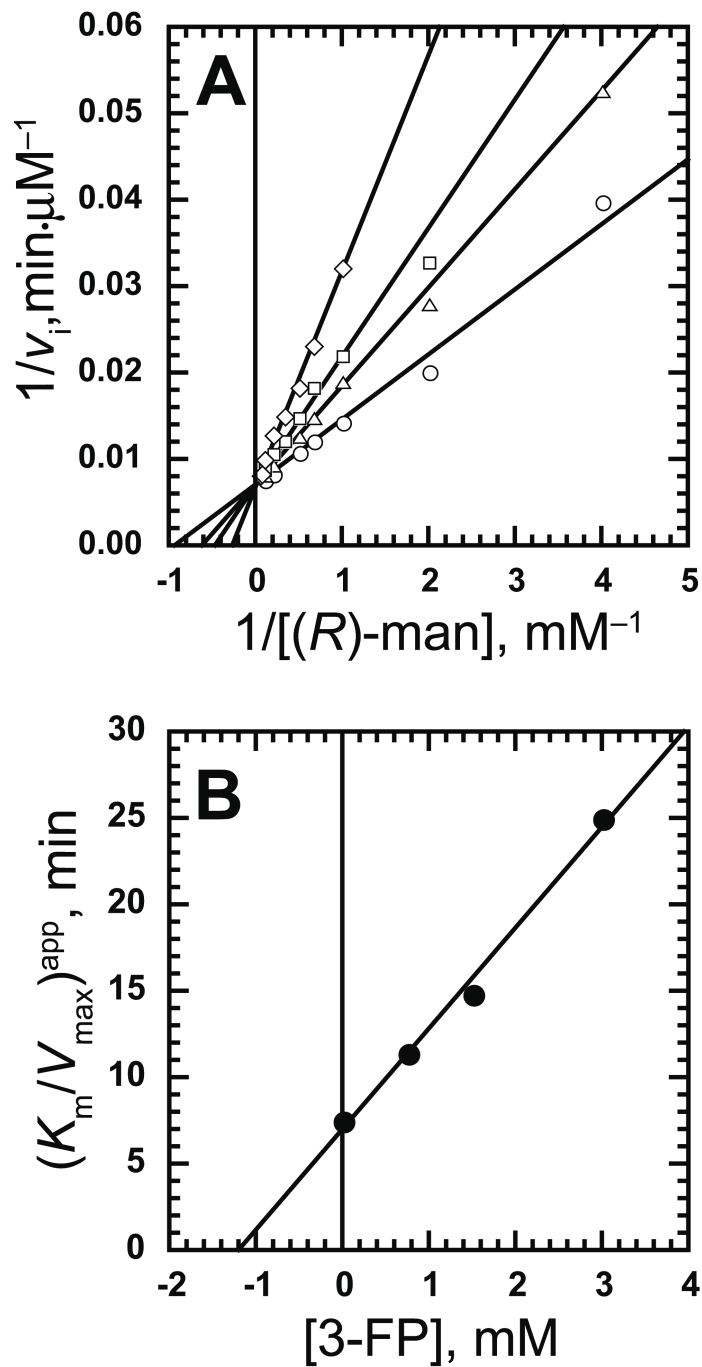
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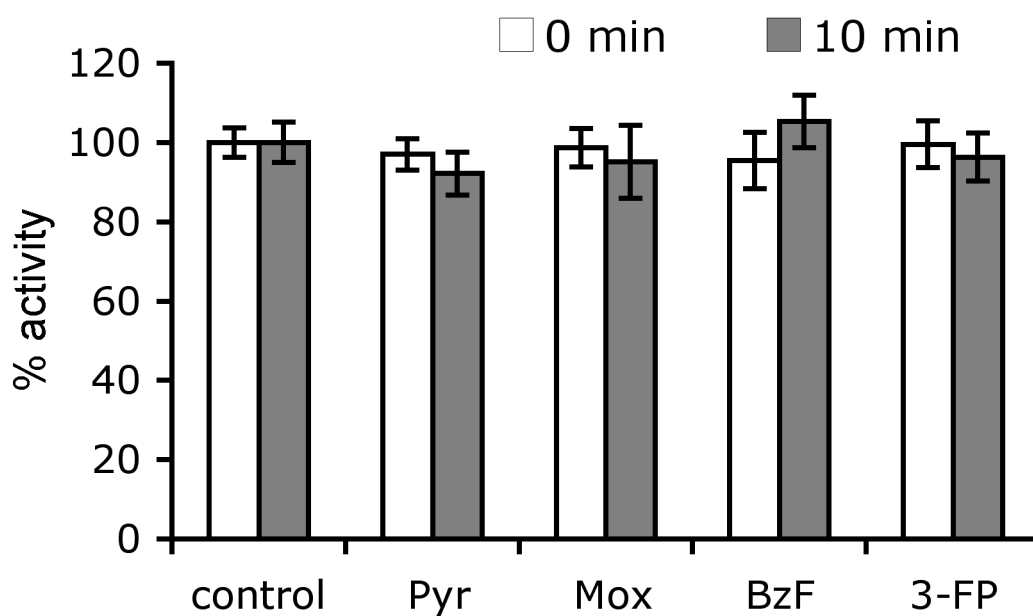




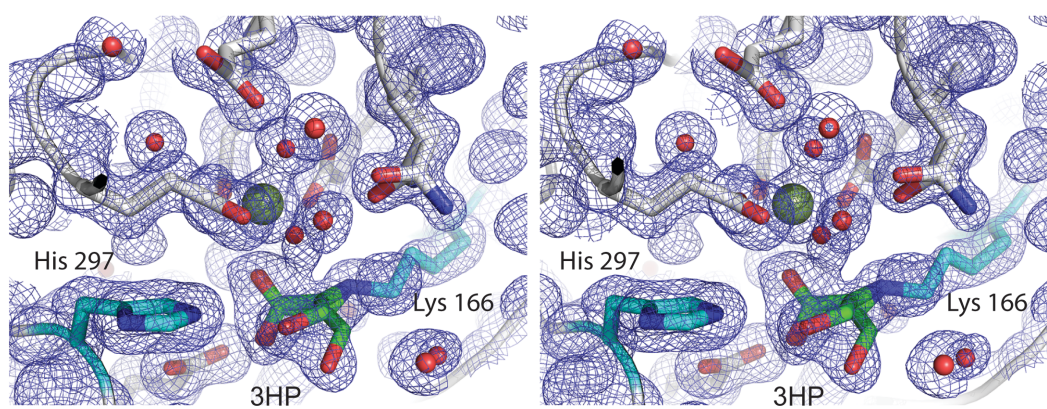
**Supplementary Figure 1S.** Competitive inhibition of MR by mesoxalate (Mox). (A) Representative Lineweaver-Burk plot. (B) Replot of the apparent  $K_m/V_{max}$  values against inhibitor concentration. Assay conditions are as described in the Materials and Methods.



**Supplementary Figure 2S.** Competitive inhibition of MR by 3-fluoropyruvate (3-FP). (A) Representative Lineweaver-Burk plot. (B) Replot of the apparent  $K_m/V_{\max}$  values against inhibitor concentration. Assay conditions are as described in the Materials and Methods.



**Supplementary Figure 3S.** Time-dependence for the inhibition of MR by  $\alpha$ -keto acids. Pyruvate (Pyr), mesoxalate (Mox), benzoylformate (BzF), and 3-fluoropyruvate (3-FP) do not show significant reduction in MR activity when incubated at concentrations of 2 mM with wild-type MR for 10 min at 25 °C.



**Supplementary Figure 4S.** Stereoview of a representative  $2F_o-F_c$  electron density map, contoured at  $1.0 \sigma$ , for the active site of 3-HP-MR. The Brønsted acid-base catalysts (His 297 and Lys 166) are colored cyan and the 3-HP adduct is colored green.