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Structure, Function and Applications of Metal- Requiring Enzymes: Carbonic Anhydrase and Epi- Isozizaene Synthase

Julie A. Aaron

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Abstract

Cryptophane Biosensors for Targeting Human Carbonic Anhydrase

Cryptophanes represent an exciting class of xenon-encapsulating molecules that can be exploited as probes for nuclear magnetic resonance imaging. A series of carbonic anhydrase-targeting, xenon-binding cryptophane biosensors were designed and synthesized. Isothermal titration calorimetry and surface plasmon resonance measurements confirmed nanomolar affinity between human carbonic anhydrase II and the cryptophane biosensors. A 1.70 Å resolution crystal structure of a cryptophane-derivatized benzenesulfonamide human carbonic anhydrase II complex was determined, and shows how an encapsulated xenon atom can be directed to a specific biological target. Furthermore, this work illustrates the utility and promise of developing xenon biosensors to diagnose human diseases characterized by the upregulation of specific carbonic anhydrase biomarkers, specifically human carbonic anhydrase IX and XII.

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The X-ray crystal structure of recombinant epi-isozizaene synthase (EIZS), a sesquiterpene cyclase from *Streptomyces coelicolor* A3(2), has been determined at 1.60 Å resolution. Specifically, the structure of wild-type EIZS is that of its closed conformation in complex with three Mg²⁺ ions, inorganic pyrophosphate (PPi), and the benzyltriethylammonium cation (BTAC). Additionally, the structure of D99N EIZS has been determined in an open, ligand-free conformation at 1.90 Å resolution. Comparison of these two structures provides the first view of conformational changes required for substrate binding and catalysis in a bacterial terpenoid cyclase, and enables a comparison of substrate recognition amongst terpenoid synthases from different domains of life. Mutagenesis of aromatic residues in the enzyme active site alters the cyclization template and results in the production of alternative sesquiterpene products. The structure and activity of several active site mutants have been explored. The 1.64 Å resolution crystal structure of F198A EIZS in a complex with three Mg²⁺ ions, PPi, and BTAC reveals an alternative binding orientation of BTAC, whereas the crystal structures of L72V, A236G and V329A EIZS reveal an unchanged BTAC binding orientation. Alternative binding orientations of a carbocation intermediate could lead to the formation of alternative products.

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STRUCTURE, FUNCTION AND APPLICATIONS OF METAL-REQUIRING
ENZYMES: CARBONIC ANHYDRASE AND EPI-ISOZIZAENE SYNTHASE

JULIE ANNE AARON

A Dissertation in
Chemistry

Presented to the Faculties of the University of Pennsylvania in Partial Fulfillment of the
Requirements for the Degree of Doctor of Philosophy

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Dr. David W. Christianson, Professor, Chemistry
Supervisor of Dissertation

Dr. Gary A. Molander, Professor, Chemistry
Department Chairman

Dissertation Committee

Dr. Ronen Marmorstein, Professor, Chemistry and Wistar Institute

Dr. Ivan J. Dmochowski, Associate Professor, Chemistry

Dr. Barry S. Cooperman, Professor, Chemistry

“One must still have chaos in oneself to be able to give birth to a dancing star.”
- Nietzsche

I dedicate this thesis to my family,
for supporting me,
and to Dr. Li and Mr. Docketis,
for inspiring me.

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ABSTRACT

STRUCTURE, FUNCTION, AND APPLICATIONS OF METAL-REQUIRING ENZYMES: CARBONIC ANHYDRASE AND EPI-ISOZIZAENE SYNTHASE

Julie Anne Aaron

David W. Christianson

Cryptophane Biosensors for Targeting Human Carbonic Anhydrase

Cryptophanes represent an exciting class of xenon-encapsulating molecules that can be exploited as probes for nuclear magnetic resonance imaging. A series of carbonic anhydrase-targeting, xenon-binding cryptophane biosensors were designed and synthesized. Isothermal titration calorimetry and surface plasmon resonance measurements confirmed nanomolar affinity between human carbonic anhydrase II and the cryptophane biosensors. A 1.70 Å resolution crystal structure of a cryptophane-derivatized benzenesulfonamide human carbonic anhydrase II complex was determined, and shows how an encapsulated xenon atom can be directed to a specific biological target. Furthermore, this work illustrates the utility and promise of developing xenon biosensors to diagnose human diseases characterized by the upregulation of specific carbonic anhydrase biomarkers, specifically human carbonic anhydrase IX and XII.

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The X-ray crystal structure of recombinant *epi*-isozizaene synthase (EIZS), a sesquiterpene cyclase from *Streptomyces coelicolor* A3(2), has been determined at 1.60 Å resolution. Specifically, the structure of wild-type EIZS is that of its closed conformation in complex with three Mg²⁺ ions, inorganic pyrophosphate (PP_i), and the benzyltriethylammonium cation (BTAC). Additionally, the structure of D99N EIZS has been determined in an open, ligand-free conformation at 1.90 Å resolution. Comparison of these two structures provides the first view of conformational changes required for substrate binding and catalysis in a bacterial terpenoid cyclase, and enables a comparison of substrate recognition amongst terpenoid synthases from different domains of life. Mutagenesis of aromatic residues in the enzyme active site alters the cyclization template and results in the production of alternative sesquiterpene products. The structure and activity of several active site mutants have been explored. The 1.64 Å resolution crystal structure of F198A EIZS in a complex with three Mg²⁺ ions, PP_i, and BTAC reveals an alternative binding orientation of BTAC, whereas the crystal structures of L72V, A236G and V329A EIZS reveal an unchanged BTAC binding orientation. Alternative binding orientations of a carbocation intermediate could lead to the formation of alternative products.

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Part I: Cryptophane Biosensors for Targeting Human Carbonic Anhydrase

Chapter 1: Introduction

1.1 Carbonic Anhydrase: A Model System

Carbonic anhydrases (CAs) are ubiquitous zinc-metalloenzymes that catalyze the hydration of carbon dioxide to form bicarbonate (Equation 1). Four distinct, unrelated CA



gene families have been identified and characterized as follows; α -CAs found in vertebrates, bacteria, algae, and the cytoplasm of green plants, the β -CAs found in bacteria, algae, and chloroplasts of mono- and dicotyledons, γ -CAs found in archaea and some bacteria, and finally, the δ -CAs found in some marine diatoms (Supuran, 2008). There are 16 α -CA isozymes in mammals, with various sub-cellular localization including the cytosol, membrane-bound, mitochondrial, transmembrane, and secreted (Table 1.1) (Supuran, 2007). Not only are these enzymes important for pH homeostasis, they are also involved in a number of other biosynthetic pathways including gluconeogenesis, ureagenesis and lipogenesis. As such, CAs have been implicated in cellular proliferation, spermatozoan motility, and aqueous humor production, and are therefore targets to treat many diseases (Ekstedt, 2004; Pastorekova, 2006; Mincione, 2008).

The X-ray crystal structures of several human CA isozymes have been solved to-date, including CA I, II, III, IV, VI, VIII, IX, XII and XIII (Table 1.1). The molecular structures of human CAs are highly conserved. The structure is characterized by a central

anti-parallel β -sheet, and the active site contains a catalytically essential Zn^{2+} ion, coordinated by three histidine residues (Figure 1.1). The active site is located at the bottom of a roughly conical, 15 Å-deep cleft, which is predominantly hydrophobic on one side and predominately hydrophilic on the other (Liljas, 1972).

Kinetic experiments have helped to elucidate the catalytic mechanism of the enzyme (Figure 1.2). A zinc-bound water molecule makes a hydrogen bond to the hydroxyl moiety of T199. The pKa of the zinc-bound water is lowered to 7, and therefore can lose a proton and become a zinc-bound hydroxide. The hydroxide, a strong nucleophile, is well oriented to attack a CO_2 molecule bound in a nearby hydrophobic pocket (Figure 1.1), forming a zinc-bound bicarbonate. Next, the bicarbonate is displaced by a water molecule and diffuses into bulk solution. The active form of the enzyme is reformed by loss of proton from the new zinc-bound water via proton transfer to bulk solvent, assisted by H64.

The most effective CA inhibitors designed to date contain an aryl-sulfonamide moiety; the sulfonamide moiety coordinates to the Zn^{2+} ion as a sulfonamidate ion, and the aryl moiety binds in the hydrophobic cleft and acts as a scaffold from which the inhibitor can be tailored to target a specific isozyme (Krishnamurthy, 2008). Hundreds of CA inhibitors have been designed and tested as potential drugs for the treatment of a variety of diseases including glaucoma (CA II and CA XII), cancer (CA IX and CA XII), obesity (CA VA, CA VB), seizures (CA II, CA VII, CA XII, CA XIV), and bacterial infections (various CAs from pathogenic organisms) (Supuran, 2007).

Table 1.1. Catalytic steady-state constants¹ and protein data bank reference codes

for human carbonic anhydrase isozymes.

Isozyme	k_{cat} (s⁻¹)⁺	k_{cat}/K_M (M⁻¹ s⁻¹)	PDB code	Reference
CA I	2.0 x 10 ⁵	5.0 x 10 ⁷	2CAB	(Kannan, 1984)
CA II	1.4 x 10 ⁶	1.5 x 10 ⁸	1CA2	(Eriksson, 1988a)
CA III	1.0 x 10 ⁴	3.0 x 10 ⁵	1Z97	(Duda, 2005)
CA IV	1.1 x 10 ⁶	5.1 x 10 ⁷	3FW3	To be published. PDB released 12/01/2009
CA VA	2.9 x 10 ⁵	2.9 x 10 ⁷	-	-
CA VB	9.5 x 10 ⁵	9.8 x 10 ⁷	-	-
CA VI	3.4 x 10 ⁵	4.9 x 10 ⁷	3FE4	To be published. PDB released 12/16/2008
CA VII	9.5 x 10 ⁵	8.3 x 10 ⁷	-	-
CA VIII (CARP) ²	-	-	2W2J	(Di Fiore, 2009)
CA IX	3.8 x 10 ⁵	5.5 x 10 ⁷	3IAI	(Alterio, 2009)
CA X	-	-	-	-
CA XI	-	-	-	-
CA XII	4.2 x 10 ⁵	3.5 x 10 ⁷	1JCZ	(Whittington, 2001)
CA XIII	1.5 x 10 ⁵	1.1 x 10 ⁷	3D0N	(Di Fiore, 2009)
CA XIV	3.1 x 10 ⁵	3.9 x 10 ⁷	1RJ5 ²	(Whittington, 2004) ³
murine CA XV	4.7 x 10 ⁵	3.3 x 10 ⁷	-	-

¹Kinetic parameters were taken from a recent review by C. T. Supuran (Supuran, 2008).²CARP is a carbonic anhydrase-related protein.³Crystal structure of murine carbonic anhydrase XIV

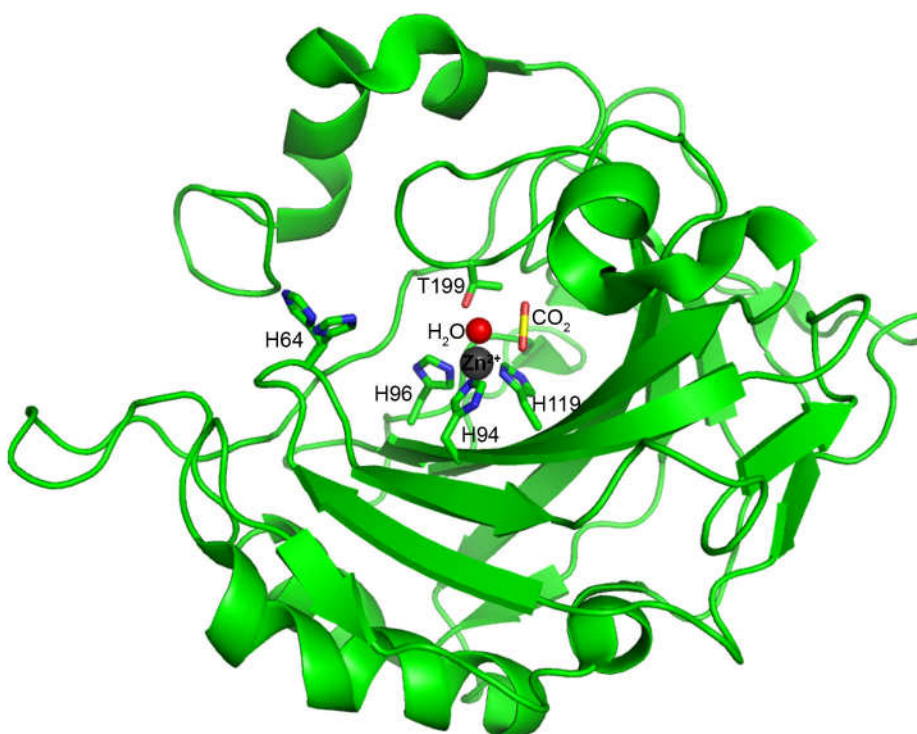


Figure 1.1. A cartoon representation of the crystal structure of human CA II, solved at 1.1 Å resolution (PDB 3D92) (Domsic, 2008) in complex with the substrate, CO₂. The catalytic Zn²⁺ ion is represented as a sphere (grey) coordinated by residues H94, H96 and H119, and is coordinated to water (red sphere), which makes a hydrogen bond to the side chain of T199. Carbon dioxide is bound in a hydrophobic pocket in the active site formed by V121, V143, and L198. A proton shuttle which transports a proton from the zinc-bound water molecule to bulk solvent is proposed to include H64, which is observed in two discrete conformations in the crystal structure.

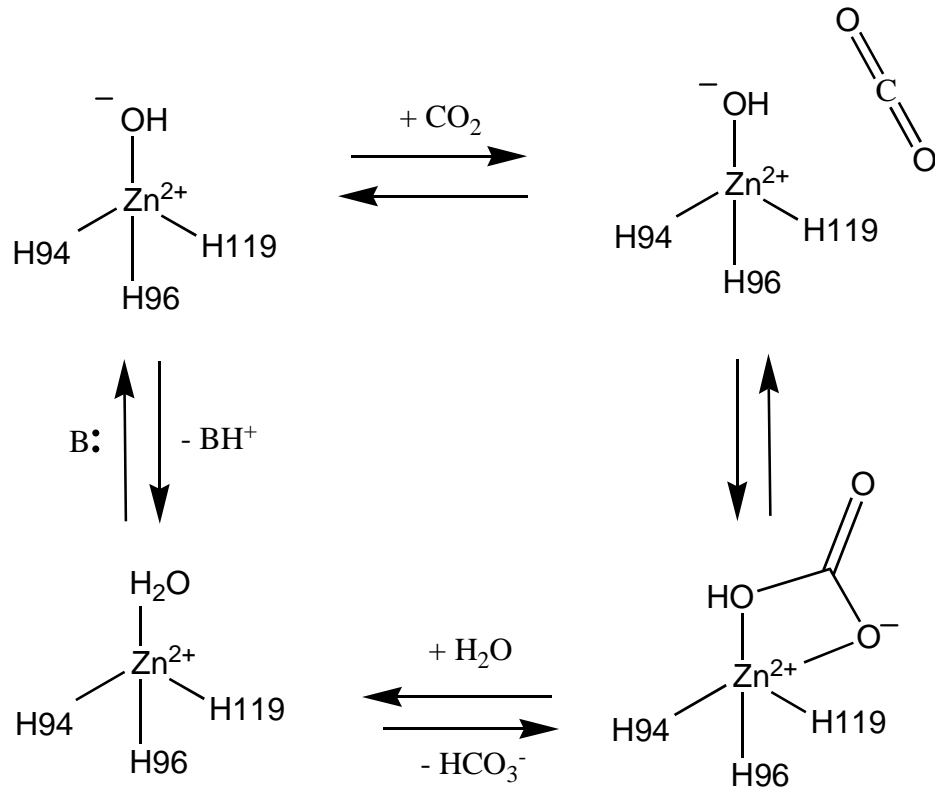


Figure 1.2. Catalytic mechanism of carbonic anhydrase. A schematic representation of the catalytic mechanism of the hydration of CO_2 by α -CAs. Carbon dioxide is bound in a hydrophobic pocket formed by V121, V143 and L198. Coordination to the zinc cation lowers the pKa of the bound water molecule from 15.7 to ~ 7 , by stabilizing its conjugate base, hydroxide. Furthermore, the positively charged zinc stabilizes the negatively-charged transition state leading to bicarbonate formation.

Of the 12 catalytically active mammalian CAs, CA II is the most thoroughly studied isozyme and is regarded as a robust model system for systematically studying protein-ligand binding (Elbaum, 1996; Krishnamurthy, 2008). Human CA II is a particularly good model system for many practical reasons, namely it is a 30 kDa monomeric enzyme that is simple to overexpress and purify from *E. coli* due to its exceptional stability. The catalytic mechanism of CA, as well as the mechanism of inhibition by Zn²⁺-binding ligands are well understood. Human CA II has the highest catalytic efficiency of the α -CAs, with $k_{\text{cat}}/K_M = 1.5 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, approaching the limit of diffusion control.

The X-ray crystal structures of several CA isozymes have been determined at high-resolution. These structures have enabled a thorough study of ligand/inhibitor binding. CA II, a soluble cytosolic isozyme, is particularly amenable to crystallization, resulting in hundreds of protein data bank (PDB) submissions, many at high resolution (better than 1.2 Å). The use of CA as a model for biophysical and physical-organic studies of protein-ligand interactions has been recently extensively reviewed (Krishnamurthy, 2008). The use of CA II as a model system for studying protein-ligand interactions is extended in this dissertation; CA II is utilized for the structure-based design of a xenon (¹²⁹Xe) biosensor for use as a magnetic resonance imaging (MRI) contrast agent.

1.2 ^{129}Xe MRI

Proton (^1H) magnetic resonance imaging (MRI) is one of the most widely used and versatile techniques for scanning deep tissue, with important applications in disease diagnosis. MRI offers many advantages for *in vivo* imaging; it is non-invasive, uses non-ionizing radiation, and produces tomographic images. Although the intrinsic ^1H MRI signals from water and fat typically have low sensitivity, contrast agents have been developed that contain gadolinium or iron-oxide particles, which improve the diagnostic power of the resulting images (Degani, 1997; Foster-Gareau, 2003). More recently however, research has shifted focus to investigate the use of other nuclear magnetic resonance (NMR) active nuclei, including ^3He , ^{13}C , ^{83}Kr and ^{129}Xe , which can be hyperpolarized to significantly increase the NMR signal. Hyperpolarization is achieved through a process known as spin-exchange optical pumping; angular momentum is transferred from circularly polarized light to an alkali metal, the newly polarized metal interacts with the nuclear spin of the isotope of interest through dipolar coupling to increase the population of unpaired spins (Kauczor, 1998). ^{129}Xe is particularly exciting for applications in imaging since it has a spin- $1/2$ nucleus, a >200 -ppm chemical shift window in water, and a natural isotopic abundance of 26% (commercially available up to 86%). Moreover, ^{129}Xe hyperpolarization can enhance MRI signals $\sim 10,000$ -fold (Cherubini, 2003).

Current *in vivo* ^{129}Xe MRI applications include functional lung imaging. Hyperpolarized ^{129}Xe MRI offers increased signal-to-noise ratios for lung tissue with

respect to conventional ^1H MRI. Typically, a mixture of hyperpolarized ^{129}Xe gas and N_2 is inhaled by a patient, where it acts as a contrast agent for visualizing the airways. Imaging the diffusion of xenon gas in the lungs has clinical applications in the diagnosis of asthma, chronic obstructive pulmonary disease, cystic fibrosis, and pediatric chronic lung disease (Fain, 2007). However, there are limitations to the application of hyperpolarized ^{129}Xe MRI imaging due to the reliance of the technique on the diffusion of xenon to the tissue of interest. This limitation can be overcome by the development of functional xenon biosensors, a strategy first proposed by Spence and colleagues in 2001 (Spence, 2001).

1.3 ^{129}Xe -Cryptophane Biosensors

Extending the application of hyperpolarized ^{129}Xe imaging beyond the lungs requires a biosensor that is able to bind xenon atoms, while simultaneously targeting the biological moiety of interest. An interesting class of organic supramolecular compounds known as cryptophanes can be used as the xenon cage. Cryptophane cages consist of two cup-shaped [1.1.1]orthocyclophane units connected by three bridging units (Figure 1.3). Cryptophanes of diverse shapes, sizes and chemical properties can be synthesized by varying the R1 and R2 substituents, the identity of the bridging units, Y, as well as the diastereomer (*syn* or *anti*). The type of bridging unit has a great effect on the size of the cage. The volume of a cryptophane-A cage [$\text{Y}=\text{O}(\text{CH}_2)_2\text{O}$, $\text{R}_1=\text{R}_2=\text{OCH}_3$] is 95 \AA^3 , and it can reversibly encapsulate xenon ($K_A \sim 3900 \text{ M}^{-1}$). The highest affinity measured to date for a cryptophane-Xe interaction is $K_A \sim 3.3 \times 10^4 \text{ M}^{-1}$, for tri-acetate cryptophane-A.

Furthermore, varying the R1 and R2 substituents facilitates the use of cryptophane cages as xenon carriers in a biosensor that can be targeted to specific proteins using an appropriate affinity tag (Lowery, 2006; Schroder, 2006). The affinity-tags and their respective targets that have been investigated include biotin-streptavidin, peptide-antigen and DNA-DNA hybrid (Spence, 2001; Roy, 2007; Schlundt, 2009). In this work, a series of racemic biosensors have been designed to target the active site of the CA isozymes, and structural and binding studies of the biosensor with human CA I and CA II follow.

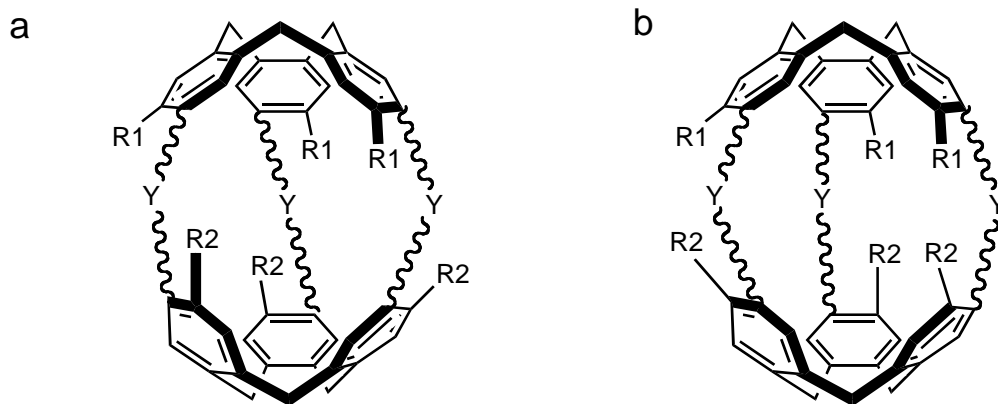


Figure 1.3. General structures of cryptophanes. (a) *anti* diastereomer. (b) *syn* diastereomer. Varying the bridging atoms (Y) and substituents (R1 and R2) give rise to cages of diverse size, shape and chemical properties.

Chapter 2: Binding Studies of ^{129}Xe -Cryptophane Biosensors and Carbonic Anhydrase

2.1 Design and Synthesis of ^{129}Xe -Cryptophane Biosensors

The design of a functional ^{129}Xe -cryptophane biosensor to target CA is quite simple. First, an appropriate cryptophane cage must be chosen as the functional xenon binding moiety. Next, an appropriate CA affinity tag must be synthesized and attached at the R1 or R2 position, and finally, the remainder of R1 and R2 positions must be derivatized with appropriate side groups to impart sufficient biosensor solubility. As a starting point for the design, the cryptophane-A cage was chosen due to its superior xenon binding capabilities. To target the catalytically active Zn^{2+} ion in the active site of CA a benzene-sulfonamide functional group was chosen, and added at the R1 position. In order to increase the aqueous solubility of the biosensor, additional carboxylic acid functional groups were coupled to the two additional R1 positions, and $-\text{OCH}_3$ groups were added at the R2 positions. A series of three biosensors were synthesized with varying linker lengths between the sulfonamidate functional group and the cryptophane cage to investigate the optimal biosensor construct. A summary of the synthesis, performed by J. M. Chambers is summarized in Figure 2.1. The CA targeting portion of the biosensor was synthesized starting from **1**. The benzene-sulfonamide moiety was conjugated to an azide linker, varying the number of methylene groups between the benzene ring and the azide moiety from 0-2 to form **2-4**. Next, starting with tripropargyl cryptophane (R1= CH_2CCH and R2= OCH_3), **5**, the azide functionality on **2-4** was stoichiometrically coupled to the cryptophane via a copper catalyzed [3+2] cycloaddition.

Following silica column chromatography to remove unstoichiometric side products, the remaining R1 positions were coupled to 3-azidopropionic acid, **6**, to form a series of water-soluble CA targeting cryptophane biosensors **7-9**.

2.2 Isothermal Titration Calorimetry Measurements

2.2.1 Introduction

Isothermal titration calorimetry (ITC) is a technique used to measure the thermodynamic parameters of interactions between molecules by measuring the change in heat upon mixing of the analytes of interest. Specifically, ITC directly measures the binding affinity (K_a), enthalpy change (ΔH), and binding stoichiometry (n) of an interaction between two molecules, in our case CA and a cryptophane biosensor. Then, using the Gibbs free energy equation (2), the entropy (ΔS) and Gibbs free energy (ΔG) of the interaction can be determined (Falconer, 2010).

$$\Delta G = -RT \ln K_a = \Delta H - T\Delta S \quad (2)$$

An ITC experiment is performed in a calorimeter that contains two identical small cells surrounded by an adiabatic jacket. One cell serves as a control/reference cell while the second cell serves as a sample cell. To begin an experiment, one of the analytes of interest, in our case the protein CA, is placed in the sample cell. Next, the biosensor is slowly titrated into the protein solution in aliquots of a few microliters per addition. Binding of the cryptophane biosensor to CA is exothermic. The instrument detects temperature differences between the sample and reference cell and measures the time-dependent input of power required to maintain both cells at the same temperature. The raw data is integrated with respect to time to determine the total amount of heat released per analyte injection, and together with the molar ratio of biosensor to protein the thermodynamic parameters of the interaction can be determined. The advantages of ITC include the ability to measure free energy, enthalpy, entropy, association constant, and stoichiometry of an interaction simultaneously without incorporating unnatural labels,

which could introduce bias to the measurement. The association constant for the interaction can be converted into a dissociation constant from the relationship $K_d = 1/K_a$ (Freyer, 2008).

2.2.2 Experimental Methods

All calorimetry experiments were conducted at 298 K on a VP-ITC titration microcalorimeter from MicroCal, Inc. (Northampton MA), using standard protocols and data analysis (Wiseman, 1989; Fisher, 1995). Human CA I was purchased from Sigma and used without further purification, human CA II was overexpressed in *Escherichia coli* and purified as previously described (Alexander, 1993). Biosensors **7**, **8** and **9**, as well as the benzenesulfonamide-linker, **4**, were synthesized by J. M. Chambers (Chambers, 2009). CA I and CA II were diluted to ~ 20 μ M and exhaustively dialyzed against 50 mM Tris-SO₄ (pH 8.0). Biosensors (~10 mM stock solutions in DMSO) were dissolved at a concentration of 135-300 μ M in an aliquot of the same buffer, and an equivalent concentration of DMSO was added to the enzyme solution. Prior to the titration experiment, samples were degassed under vacuum for 5 min. The sample cell (effective volume = 1.4 mL) was overfilled with 1.8 mL of CA at a concentration of 14-26 μ M, and the reference cell was filled with water. The contents of the sample cell were titrated with 30 aliquots (10 μ L each) of inhibitor (two initial 2 μ L injections were made, but not used in data analysis). After each injection, the heat change was measured and converted to the corresponding enthalpy value. The reaction mixture was continuously stirred at 300 rpm during titration. Control experiments were carried out by titrating the inhibitor into the buffer solution under identical experimental conditions. The

calorimetric data are presented with the background titrations subtracted from the experimental data. The amount of heat produced per injection was calculated by integration of the area under each peak. Data were fit to the equation $q = V \Delta H[E]_t K[L]/(1 + K[L])$, where q is the heat evolved during the course of the reaction, V is the cell volume, ΔH is the binding enthalpy per mole of ligand, $[E]_t$ is the total enzyme concentration, K is the binding constant, and $[L]$ is inhibitor concentration. Nonlinear regression fitting to the binding isotherm (ORIGIN 5.0 software, MicroCal) using a one-site model gave the equilibrium dissociation constant of the ligand, K_d , and estimates of the standard error. Representative isothermal calorimetric data and binding isotherms are shown in Figures 2.2 and 2.3 and a summary of dissociation constants for the CA-biosensor complexes are summarized Table 2.1. The error is $\sigma_i = \sqrt{(C_{ii}\chi^2)}$, where C_{ii} is the diagonal element of the variance-covariance matrix.

2.2.3 Results and Discussion

ITC binding measurements indicate that all three biosensors have nanomolar affinity for human CA I and CA II (Table 2.1). The K_d of the benzenesulfonamide affinity tag, **4**, was measured as a control to determine the contribution of the cryptophane on the binding constant. ITC binding studies indicate that the presence of cryptophane and length of the linker between the sulfonamide and the cage has little effect on the K_d . Overall, all three biosensors exhibited modestly higher affinity for CA I (20 – 80 nM) versus CA II (60-100 nM). Biosensor **9** exhibited the highest affinity for CA II (60 nM), while biosensor **7** exhibited the highest affinity for CA I (20 nM) (Chambers, 2009).

Table 2.1. Summary of dissociation constants for biosensors-carbonic anhydrase complexation determined by ITC at 298 K.

CA Isozyme	Ligand	K_d (nM)
I	4	30 ± 10
	7	20 ± 10
	8	80 ± 10
	9	30 ± 20
II	4	100 ± 10
	7	100 ± 20
	8	110 ± 30
	9	60 ± 20

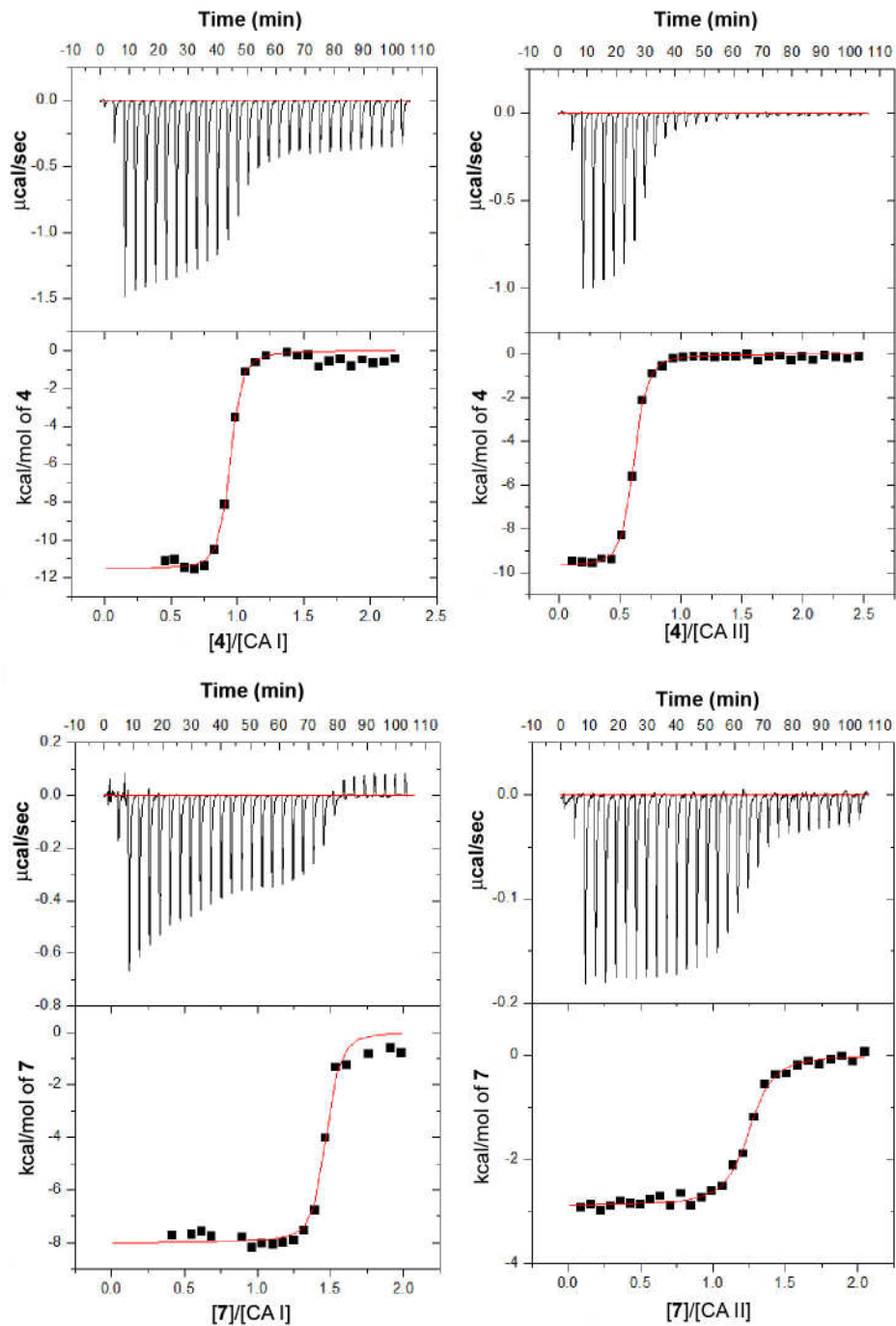


Figure 2.2. Isothermal calorimetric data for the interactions of **4** and **7** with CA I (left panel) and CA II (right panel). CA I (19.9 μM) titrated with **4** (200.0 μM); CA II (25.9 μM) titrated with **4** (292.3 μM); CA I (20.0 μM) titrated with **7** (183.0 μM); CA II (19.32 μM) titrated with **7** (182 μM).

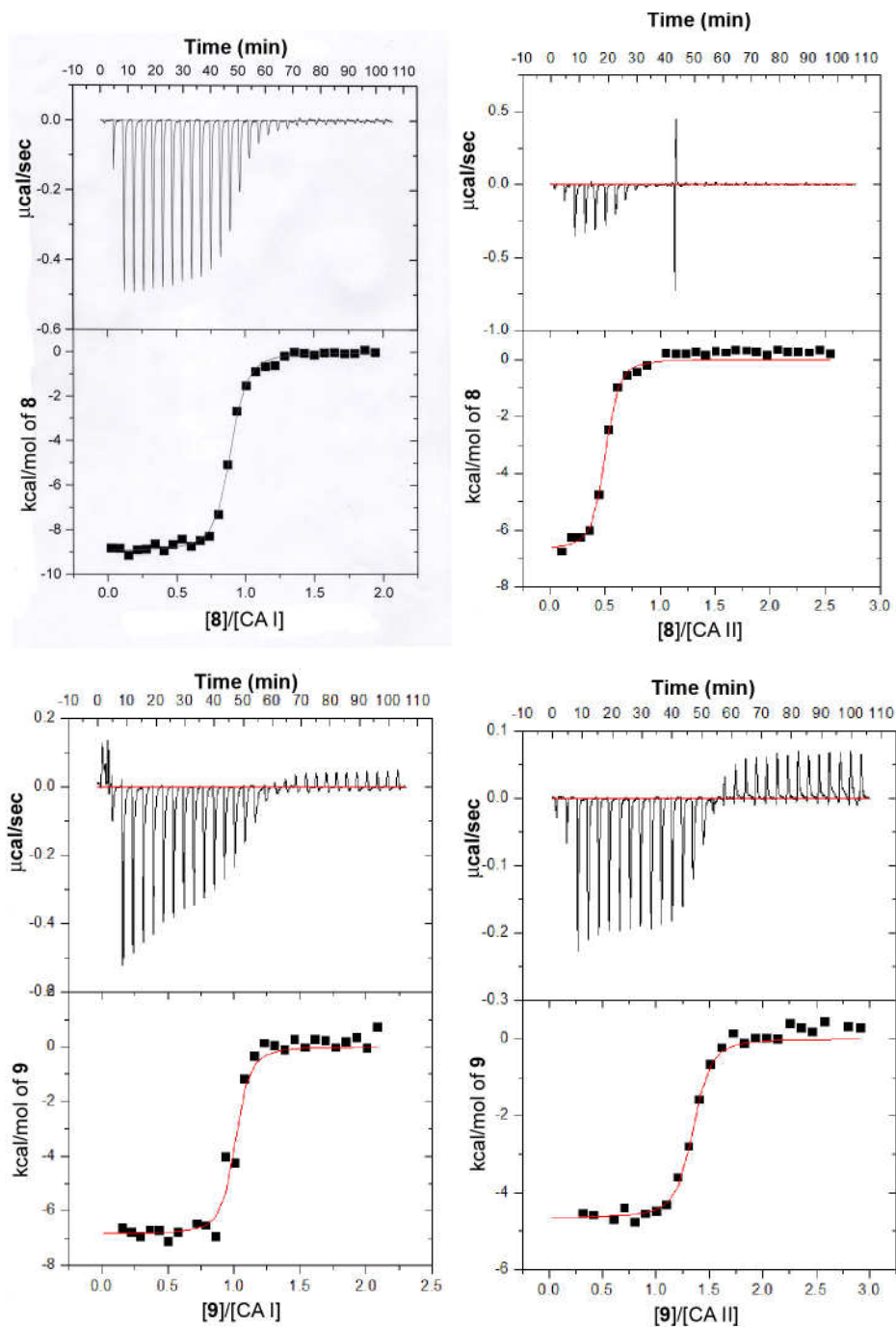


Figure 2.3. Isothermal calorimetric data for the interactions of **8** and **9** with CA I (left panel) and CA II (right panel). CA I (22.4 μM) titrated with **8** (200.0 μM); CA II (16.6 μM) titrated with **8** (194.9 μM); CA I (14.2 μM) titrated with **9** (135.7 μM); CA II (14.9 μM) titrated with **9** (200 μM).

2.3 Surface Plasmon Resonance Measurements

2.3.1 Introduction

Surface plasmon resonance (SPR) is a physical phenomenon that can be used to detect very small changes at a surface. Specifically, SPR experiments measure a change in the local index of refraction at a surface, resulting in a change in resonance conditions of surface plasmon waves (Pattnaik, 2005). Recently, a number of instruments based on SPR have become commercially available and are specifically designed to quantify macromolecular interactions (Jason-Moller, 2006). The experimental setup involves immobilization of one of the binding partners of interest (typically referred to as the “ligand”) to a sensor-chip. A typical sensor-chip is a glass slide coated with a thin layer of gold followed by a specific surface matrix, to which the ligand is attached (examples include carboxymethylated dextran, streptavidin, nickel chelation, and hydrophobic monolayer). Surface plasmons are excited by incident light beam on the opposite side of the gold surface. The incident photons induce an evanescent light field into the gold film and at a certain incident angle are able to excite surface plasmons. When a plasmon is excited, the change in the reflected light is observed at that incident angle is measured by a charged couple device (CCD) chip (Pattnaik, 2005). The first step of an SPR experiment is to immobilize the “ligand” on the sensor chip surface. Next, the “analyte” flows over the chip, where it interacts with the “ligand” (the association phase), this interaction is correlated to the change in mass at the sensor surface, resulting in an observable change in the SPR angle, detected by a change in the intensity of the reflected light (measured in resonance units (RU)). Finally, the chip is regenerated by flowing buffer over the surface of the chip (dissociation phase). The time-dependant change in

SPR signal recorded during an experiment can be fit to calculate the binding constant, K_d , for the interaction of the specific ligand and analyte (Pattnaik, 2005).

SPR techniques offer many advantages for studying binding of small molecules and macromolecular targets; the technique can be label-free, interactions can be studied in real time, it requires very small volumes of protein, and one immobilized sample on a chip can be reused many times to study a variety of different analytes. However there are a number of challenges as well since the change in refractive index is relatively small, accurate results require optimization of experimental parameters and high-quality data for consistent, accurate results (Cannon, 2004).

2.3.2 Experimental Methods

Interaction analyses were performed using a Biacore 3000 SPR instrument (Biacore AB, Uppsala, Sweden) at The Protein Core Facility, Children's Hospital of Philadelphia. Recombinant human CA II was coupled to a carboxymethylated dextran (CM5) chip using amine coupling reagents *N*-ethyl-*N'*-(3-dimethylaminopropyl)carbodiimide, *N*-hydroxysuccinimide (NHS), and ethanolamine HCl, using previously published procedures (Cannon, 2004). The CA II stock solution was prepared in 100 mM sodium acetate (pH 4.9). Samples of the analytes, biosensors **7-9**, were prepared in the running buffer (20 mM Na₂HPO₄-NaH₂PO₄, pH 7.4, 1.5 M NaCl, 3% DMSO) at the following analyte concentrations, 62.5 nM, 125 nM, 250 nM, 500 nM, 1000 nM, 2000 nM. Each concentration of analyte was tested in duplicate. Kinetic data

were fit to a simple 1:1 interaction model (Langmuir binding) using the program BIAevaluation.

2.3.3 Results and Discussion

Dissociation constants for the interactions of human CA II and biosensors **7-9** were determined using SPR (sensorgrams are shown in Figure 2.4 and a summary of the results in Table 2.2). Comparison of the dissociation constants obtained from ITC and SPR confirm nanomolar-binding affinity of the cryptophane biosensors to CA II. Furthermore, SPR measurements indicate that biosensor **7** binds tightest to human CA II, followed by biosensor **9** then **8**. Interestingly, there is a greater than 10-fold discrepancy between the K_d determined by SPR and ITC for the biosensor **7**-CA II interaction. The origins of this discrepancy are unclear; as the fitting parameters for the SPR and ITC data are satisfactory. Overall, the SPR experiments illustrate the usefulness of this technique for studying cryptophane-biosensor-protein interactions, specifically in a high-throughput manner.

Table 2.2. Dissociation constant determined by surface plasmon resonance for biosensors **7-9** with human CA II.

Analyte	K_d (nM)	χ^2
7	7.33	0.853
8	207	0.442
9	40.9	1.1

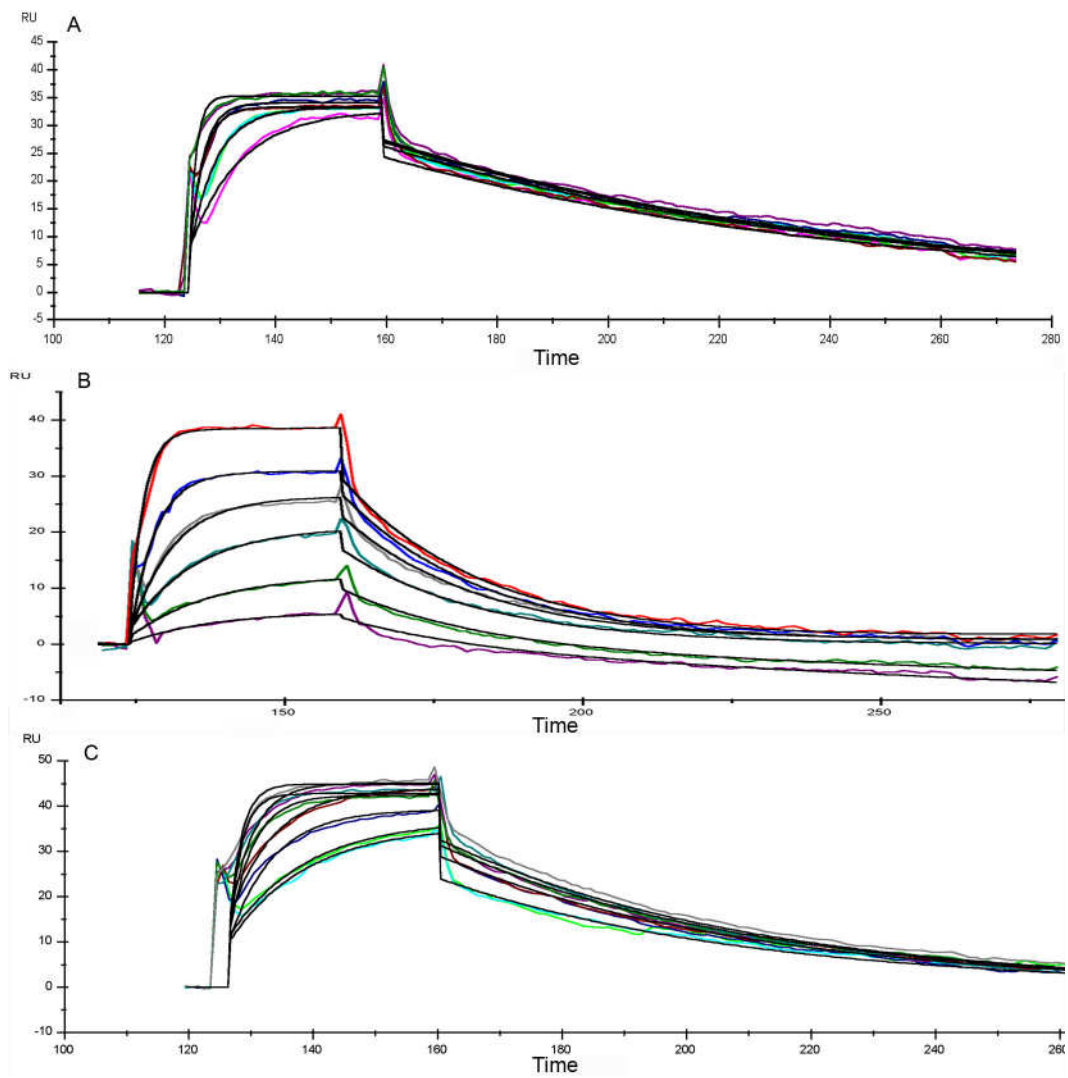


Figure 2.4. Surface plasmon resonance sensorgrams for the interaction between human CA II and (A) biosensor **7**, (B) biosensor **8**, and (C) biosensor **9**.

Chapter 3: Structural Studies of ^{129}Xe -Cryptophane-CA II

3.1 Materials and Methods

Human CA II was overexpressed in *Escherichia coli* and purified as previously described (Alexander, 1993). Crystals of the CA II-biosensor complex were formed by adding a two-fold excess of biosensor **9** (10 mM stock in DMSO) to 0.5 mg/mL CA II (50 mM Tris-SO₄, pH 7.5) and incubating at 4 °C for one hour. The mixture was concentrated using a YM-10 filter to a final CA II concentration of 10 mg/mL. Crystals were grown using the hanging drop method: a 5 µL drop of CA II solution was added to a 5 µL drop of precipitant solution (50 mM Tris-SO₄, 16% PEG 3350, 3.5 mM β-mercaptoethanol) and suspended over a reservoir containing 1 mL 50 mM Tris-SO₄, 27-32% PEG 3350 and 3.5 mM β-mercaptoethanol at 4 °C. Crystals formed within 1-2 weeks and were improved with seeding. Cubic crystals grew to typical dimensions 0.2 mm x 0.2 mm x 0.2 mm (Figure 3.1). Crystals were cryoprotected by augmentation of the mother liquor with 15% glycerol and then looped and pressurized with Xe_(g) for 30 minutes at 20 atm using a Xenon Chamber (Hampton Research). Crystals were flash cooled 10 seconds after Xe pressurization. Crystals yielded diffraction data to 1.70 Å at the Cornell High Energy Synchrotron Source (CHESS) beamline F-2 ($\lambda = 0.9795$ Å, 100 K), using an ADSC Quantum 210 CCD detector (Szebenyi, 1997). Diffraction data were indexed, integrated and scaled using HKL2000 (Otwinowski, 1997). Crystals belonged to space group C2 (unit cell parameters $a = 67.4$ Å, $b = 50.0$ Å, $c = 81.0$ Å, $\beta = 107.1^\circ$) and were isomorphous with those of T199P CAII complexed with thiocyanate (PDB 1LG6) (Huang, 2002). Initial phases were obtained by molecular replacement using the program

Phaser (Storoni, 2004) with PDB 1LG6 (less water molecules and ligand) as a search probe for rotation and translation functions. The programs CNS (Brunger, 1998) and O (Jones, 1991) were used in refinement and rebuilding, respectively. Figures were generated using PyMOL. Molecular surface area was calculated with protein interfaces, surfaces and assemblies service PISA at the European Bioinformatics Institute (http://www.ebi.ac.uk/msd-srv/prot_int/pistart.html) (Krissinel, 2007). Data collection and refinement statistics are summarized in Table 3.1.

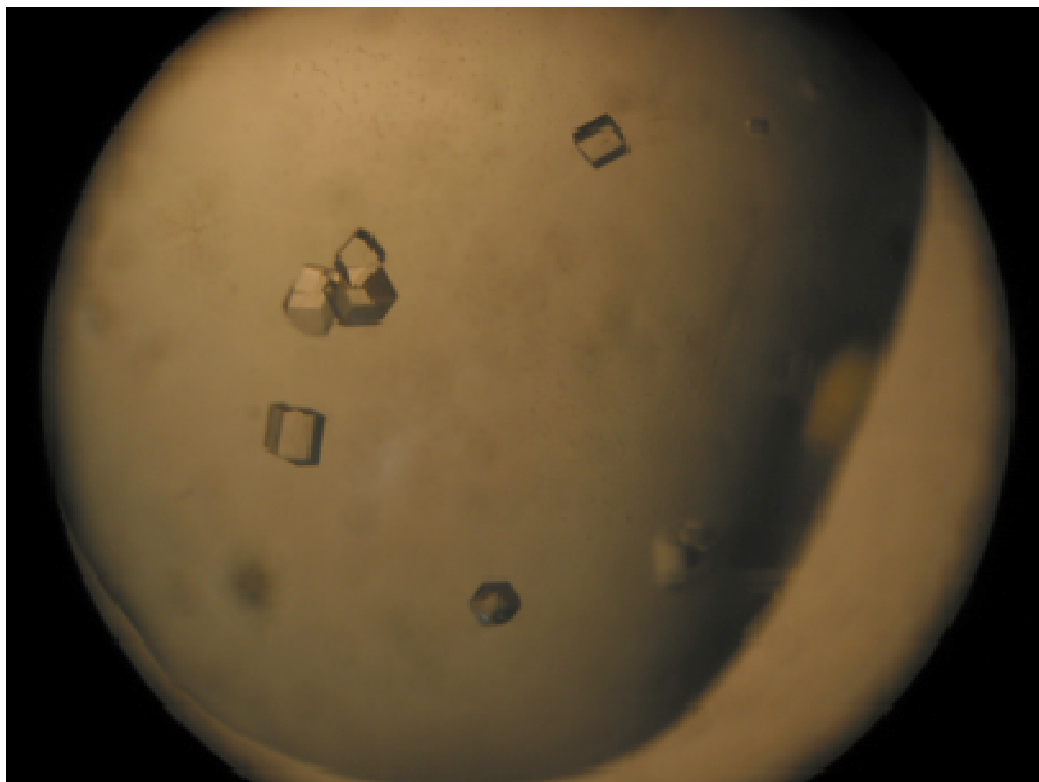


Figure 3.1. Crystals of Biosensor-9-CA II.

Table 3.1. Data Collection and Refinement Statistics for CA II-9-Xe complex.

CA II-9-Xe complex	
Data Collection	
PDB Code	3CYU
Resolution, Å	38.7 – 1.70
Total reflections measured ^a	52826 (4698)
Unique reflections measured ^a	27728 (2556)
R _{merge} ^{a,b}	0.078 (0.496)
I/σ(I) ^a	27.1 (2.3)
Completeness (%) ^a	97.0 (90.3)
Redundancy ^a	3.9 (3.7)
Refinement	
Reflections used in refinement/test set	24730/1139
R _{work}	0.226
R _{free}	0.249
Protein atoms ^c	2049
Water molecules ^c	185
Xe atoms ^c	2
Cryptophane-A-benzenesulfonamide atoms ^c	103
R.m.s deviations	
Bond lengths, Å	0.016
Bond angles, °	1.8
Dihedral angles, °	22.4
Improper dihedral angles, °	0.7
Average B-factors, Å²	
Main chain	31
Side chain	35
Xe atoms	43
Zn atom	28
Cryptophane-A-benzenesulfonamide atoms	42
Solvent	40
Ramachandran Plot^d	
Allowed (%)	86.6
Additionally allowed (%)	12.5
Generously allowed (%)	0.9
Disallowed (%)	0.0

^aNumber in parentheses refer to the outer 0.1 Å shell of data.

^b $R_{\text{merge}} = \sum |I - \langle I \rangle| / \sum I$, where I is the observed intensity and $\langle I \rangle$ is the average intensity calculated for replicate data.

^cPer asymmetric unit

^dRamachandran plot statistics calculated for non-proline and non-glycine residues using PROCHECK (Laskowski, 1993).

3.2 Results and Discussion

The X-ray crystal structure of the CA II-**9**-Xe complex was solved to 1.70 Å resolution, and refined to final R_{work} and R_{free} values of 0.23 and 0.25, respectively (Aaron, 2008). Two xenon sites were identified by inspection of the Bijvoet difference Fourier map calculated from anomalous data. The first site is near the opening of the active site cleft, 18 Å from Zn^{2+} and 8 Å from the protein chain, and corresponds to the Xe atom encapsulated by the cryptophane (Figure 3.2). The encapsulation of Xe within the cryptophane cage of **9** is confirmed by inspection of the Bijvoet difference Fourier map calculated from anomalous scattering data. X-ray diffraction data was collected at a wavelength $\lambda = 0.9795$ Å, which is far from the Xe L_1 edge of 2.27 Å (Watanabe, 1965). Nevertheless, the anomalous scattering component f'' is 3.4 e^- for Xe, so the anomalous signal is still prominent at the wavelength of data collection. The second Xe site is a hydrophobic pocket defined by A116, L148, V218, L157, V223 and F226, which is consistent with the known binding interactions of Xe in other systems (Figure 3.3) (Prange, 1998). The crystallographic occupancies of these Xe sites refine to 0.50 and 0.37, respectively. Anomalous scattering peaks are absent from crystals not subject to Xe pressurization.

The occupancy of the active site zinc ion was refined at 0.5, which was consistent with the occupancy of 0.5 determined for biosensor **9**. Because the crystallographic occupancy was thus 0.5 for Xe encapsulated within the cryptophane moiety, and the electron density map indicated the binding of both cryptophane enantiomers, each enantiomer was refined with an occupancy of 0.25 (average B-factor = 42 Å²). A total of

185 water molecules were included in later cycles of refinement. Data reduction and refinement statistics are recorded in Table 2. The N-terminus (N1-H3) was disordered and is omitted from the final model.

Biosensor **9** coordinates to the active site Zn^{2+} ion as the sulfonamidate anion, displacing the zinc-bound hydroxide ion of the native enzyme as previously observed in other complexes of CA II with benzenesulfonamide derivatives (Eriksson, 1988b; Elbaum, 1996; Supuran, 2007; Krishnamurthy, 2008). The crystallographic occupancies of **9** and Zn^{2+} are refined at 0.5. It is unusual to observe diminished Zn^{2+} occupancy in a CA II-inhibitor complex, and the molecular origins of this effect are not clear. Notably, **9** contains a chiral axis and the electron density map reveals the binding of equal populations of both enantiomers (Figures 3.4 and 3.5), (each refined with an occupancy of 0.25) (Eliel, 1994; Collet, 1996; Ruiz, 2006). Overall, the binding of **9** does not cause any significant structural changes in the active site, and the root-mean-square deviation is 0.34 Å for 256 $\text{C}\alpha$ atoms between the current structure and the unliganded enzyme (PDB 2CBA) (Hakansson, 1992). The total surface area of **9** is $\sim 1500 \text{ \AA}^2$, of which $\sim 500 \text{ \AA}^2$ becomes solvent inaccessible due to contacts of **9** within the active site cleft of CAII designated molecule I in Figure 3.6. Crystal contacts bury an additional 540 \AA^2 of the surface of **9** as follows: 270 \AA^2 with molecule III, and 240 \AA^2 and 30 \AA^2 with the front and back faces of molecule II, respectively. Molecule IV does not contact **9** bound to molecule I.

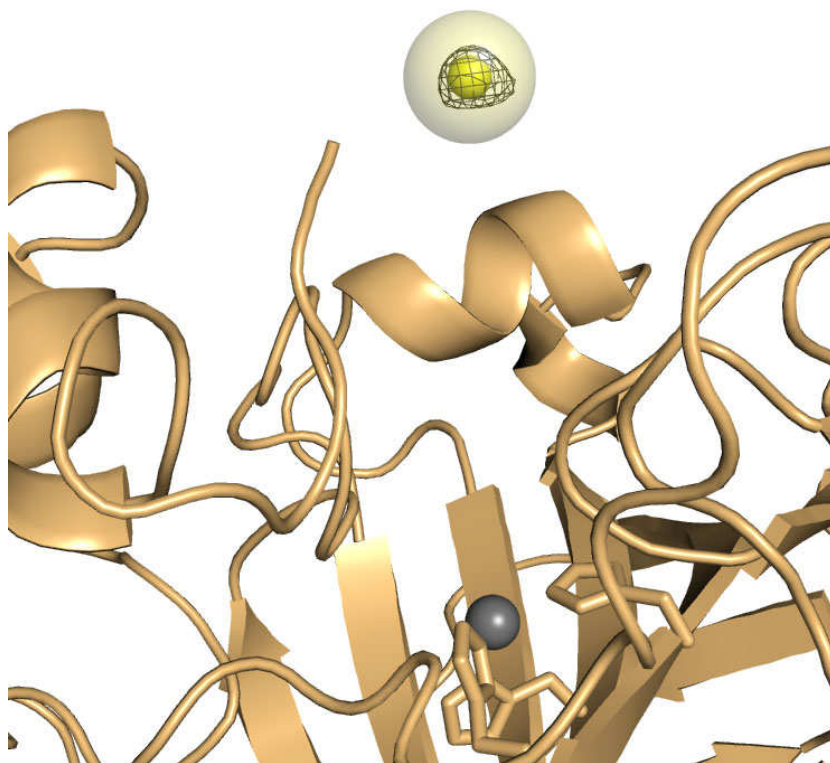


Figure 3.2. Anomalous Fourier map showing Xe location at opening of CA II active site. Xe (yellow sphere) was identified upon inspection of Bijvoet difference Fourier map (black) calculated from anomalous data to be 18 Å from Zn²⁺ (gray sphere) at opening of active site cleft. The Xe occupancy is refined at 0.50. The van der waals radius of Xe is shown as a translucent yellow sphere.

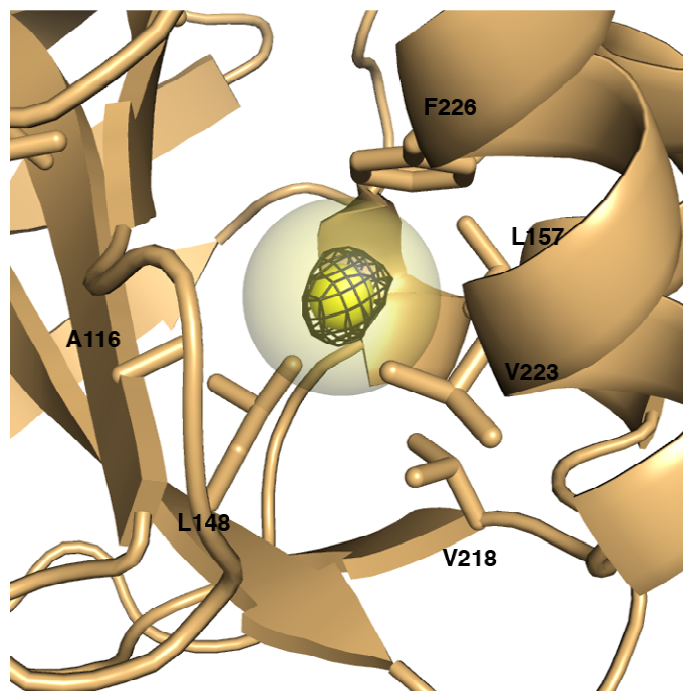


Figure 3.3. Anomalous Fourier map showing Xe location in CA II hydrophobic pocket. A second binding site is observed in a hydrophobic pocket defined by A116, L148, V218, L157, V223 and F226. Binding at this site is consistent with Xe binding in other proteins. Occupancy is 0.37. The van der Waals radius of Xe is shown as a translucent yellow sphere.

Some structural changes are observed near the outer rim of the active site cleft where the cryptophane binds. The most notable change is observed for Q136, which rotates $\sim 180^\circ$ to make van der Waals contacts with the cryptophane and the symmetry-related cryptophane bound to molecule III in the crystal lattice. Other residues at the active site rim of molecule I that make close contacts with the cryptophane are G132 and P202. Additional structural changes in the crystal lattice result from the binding of **9** to molecule I: in molecule II, H36 rotates $\sim 90^\circ$ to make a van der Waals contact with the cage, and Q137 of molecule III rotates $\sim 90^\circ$ to donate a hydrogen bond to an ether oxygen atom of **9**.

Although the pendant propionates appear to be more disordered than the cryptophane and are characterized by correspondingly weaker electron density, a hydrogen bond between a propionate moiety and Q53 of molecule II is observed. The relative dearth of strong cryptophane-protein interactions may explain why the affinity of **9** measured by ITC is only slightly better than that measured for the parent triazole-benzenesulfonamide lacking the cryptophane ($K_D = 100 \pm 10$ nM). Larger refined thermal (B) factors for **9** ($\langle B \rangle = 42 \text{ \AA}^2$) compared with the overall CA II model ($\langle B \rangle_{\text{main chain}} = 31 \text{ \AA}^2$; $\langle B \rangle_{\text{side chain}} = 35 \text{ \AA}^2$) reflect the mobility and conformational heterogeneity of the bound biosensor.

Limited hydrogen bond interactions between CA II and the cryptophane moiety of **9** may be advantageous for the use of cryptophanes as ^{129}Xe biosensors. Translational and rotational freedom, the consequence of a flexible linker between the cryptophane and the

benzenesulfonamide, could allow the cage to reorient rapidly *in situ*, independently of the protein, to result in decreased correlation times and narrower line widths that increase the sensitivity of ^{129}Xe NMR measurements in solution. In conclusion, this work reveals the first experimentally determined structure showing how an encapsulated ^{129}Xe atom can be specifically directed to a biomedically relevant protein target. The possible implications for cancer diagnosis are profound, given that CA isozymes IX and XII are overexpressed on the surface of certain cancer cells (Pastorekova, 2006). Moreover, a search of the Protein Data Bank reveals that with its molecular mass of 1554, the **9**-Xe complex is one of the largest synthetic organic ligands ever cocrystallized with a protein. Thus, this work demonstrates the feasibility of preparing crystalline complexes between proteins and nonbiological, nanometer-scale ligands.

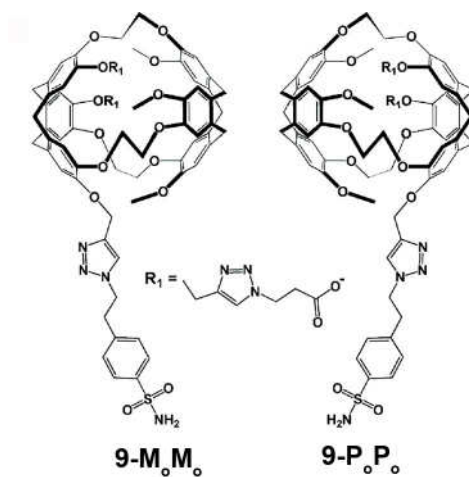


Figure 3.4. The M_0M_0 and P_0P_0 enantiomers of the cryptophane-A-derived CA biosensor. The benzenesulfonamide moiety serves as an affinity tag that targets the Zn^{2+} ion, and the R_1 substituents contain triazole propionate moieties that enhance aqueous solubility.

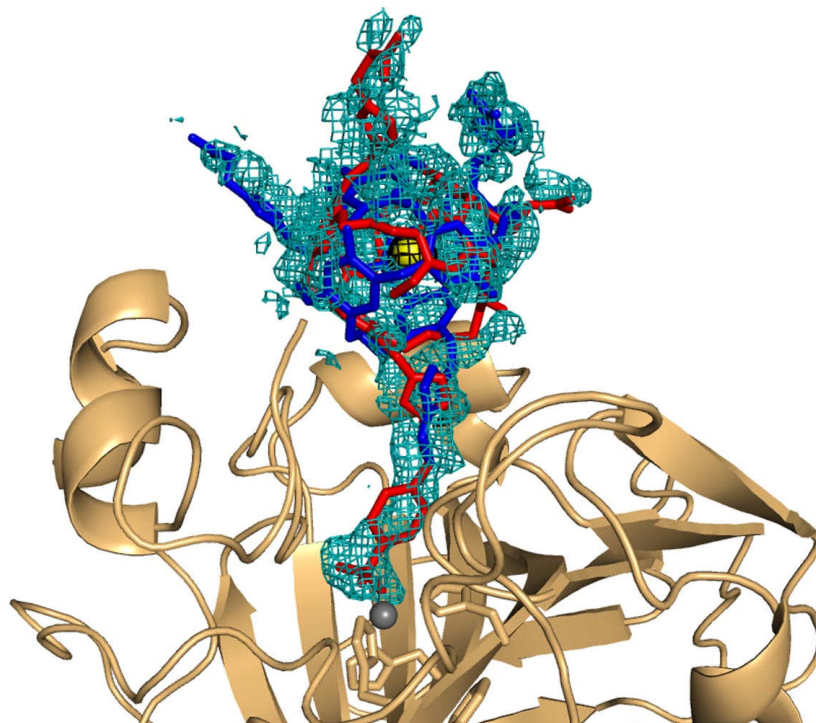


Figure 3.5. The crystal structure of biosensor **9** bound to CAII. Refinement revealed the binding of equal populations of both enantiomers of **9**. A simulated annealing omit map showing **9**-M_oM_o (blue) and **9**-P_oP_o (red) bound in the active site (1.9 σ contour, teal).

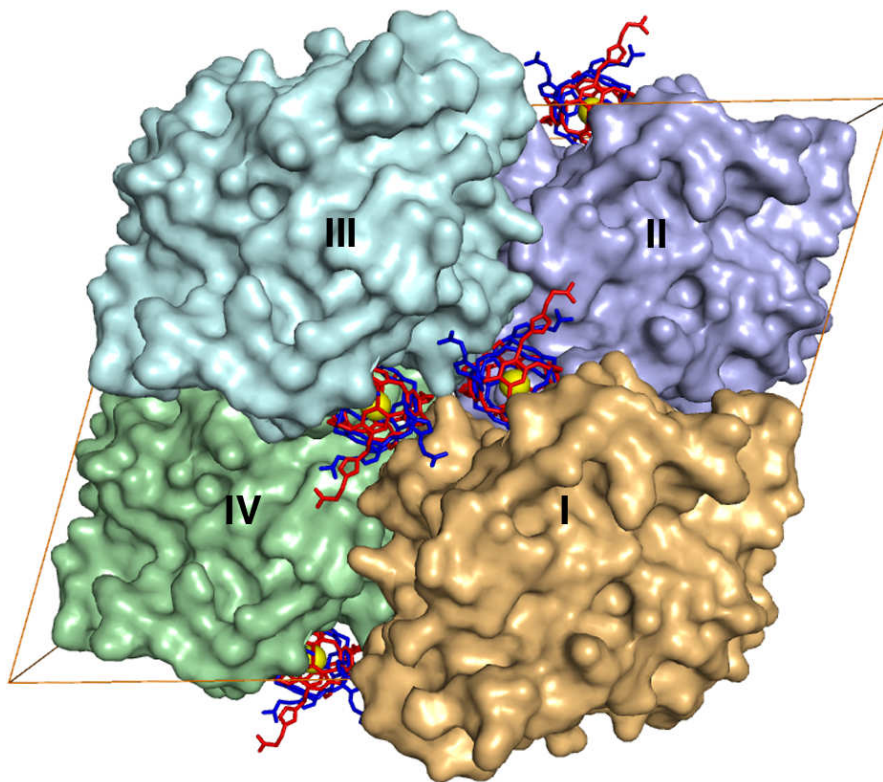


Figure 3.5. The unit cell of CAII-biosensor-9 complex. The crystals in space group C2 contains four molecules: I (x,y,z), II ($x+1/2,y+1/2,z$), III ($-x,y,-z$) and IV ($-x+1/2,y+1/2,-z$).

3.3 Future Applications of Cryptophane-based CA Biosensors

Although a very valuable model system, molecular imaging of human CA II is not relevant for disease diagnosis since CA II is found in many tissues in the body, and is not upregulated in connection with a particular disease. In comparison, CA IX and CA XII overexpression is induced by hypoxia, a pathological condition caused by oxygen deprivation, and associated with many types of cancer (Supuran, 2008; De Simone, 2010). CA IX and CA XII are two of only three transmembrane human CAs (the third is CA XIV). In addition to tumors, CA IX is also localized in gastrointestinal mucosa, while CA XII is localized in renal, intestinal, reproductive epithelial, and eye tissues (Kivela, 2005; Supuran, 2008). Structural analyses of human CA IX and XII offer useful insight in the design of a selective biosensor for imaging cancerous tissue.

Human CA IX contains four domains in addition to an N-terminal 37 amino acid signal peptide. Attached to the signal peptide is a proteoglycan-like (PG) domain, followed by a 257 residue catalytic domain, a transmembrane segment that transverses the membrane once, and finally a short intracytoplasmic tail (De Simone, 2010). The crystal structure of the catalytic domain of CA IX was recently determined (Alterio, 2009), and revealed several interesting structural observations, which may aid in rational drug design of selective CA IX inhibitors. Human CA IX forms a dimer (Hilvo, 2008), mediated by an intermolecular disulfide bond between C41 of each monomer, in addition to $\sim 1590 \text{ \AA}^2$ buried surface area (Alterio, 2009). The structure of the dimer suggests that the proteoglycan domains are located at the border of the active site, and are proposed to be involved in assisting the catalytic domain in catalysis. pH dependent activity profiles

indicate that the presence of the PG domains reduces the pKa of the zinc-bound water molecule from 7.01 to 6.49, making CA IX more active in solid and hypoxic tumors where the pH is typically slightly acidic, and coincidentally where CA IX is typically overexpressed (Alterio, 2009). Overall, the structure of the catalytic region of CA IX is highly conserved with respect to the other CA isozymes, the rmsd for the superposition of the backbone atoms of the catalytic domain of CA IX with human CA II is 1.4 Å. CA IX was co-crystallized with the acetazolamide, a sulfonamide inhibitor with a K_I of ~9-25 nM, depending upon the CA IX construct (Hilvo, 2008). However, some structural differences in the region of residues 125-137, namely residues 131, 132, 135, and 136 can be exploited for specific inhibitor design. Furthermore, the extracellular localization of CA IX allows for the design of positively charged, membrane-impermeable, inhibitors which can bind to membrane-associated CA isozymes without effecting cytosolic and mitochondrial CA isoforms (Alterio, 2009).

Since CA IX expression is highly tumor specific, and is primarily localized on the surface of solid tumors including gliomas/ependymomas, mesotheliomas, and many types of carcinomas (Potter, 2003), it is an ideal candidate for selective imaging for early cancer detection. Specifically, a functionalized cryptophane containing a CA IX targeting moiety coupled with positively-charged amine water-solubilizing groups are a starting point for design of a CA IX-specific ^{129}Xe MRI contrast agent (Taratula, 2009).

The crystal structure of the catalytic domain of human CA XII also indicated the formation of a biological isologous dimer (Whittington, 2001). The dimer interface

contains 19 hydrogen-bond interactions and buries $\sim 1,100 \text{ \AA}^2$ of surface area per monomer, while leaving the enzyme active site accessible. The residues that define the core of the active site are highly conserved with respect to CA II, however the structures diverge in the “130’s segment” of the active site. Specifically, human CA XII has an alanine residue in place of F131 in human CA II, creating a larger cavity in the active site and exposing S135 (Whittington, 2001). This region of the human CA XII active site can be exploited for CA XII-specific biosensors.

Recently, bioreductive nitro-containing sulfonamides have been investigated as carbonic anhydrase inhibitors with selectivity for tumor associated CA IX and XII. Several compounds have been identified with selectivity ratios for the inhibition of CA IX and XII over CA II up to 17 times, and for the inhibition of CA IX and XII over CA I up to 1400 times (D'Ambrosio, 2008). These compounds serve as excellent lead compounds for the design of a selective CA IX or CA XII biosensor.

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Part II: Structural and Functional Studies of the Sesquiterpene Cyclase *epi*-Isozizaene Synthase

Chapter 4: Introduction

4.1 Terpenes and Terpene Synthases

The terpenome is comprised of a family of more than 55,000 structurally and stereochemically diverse natural products, all of which ultimately derive from the universal 5 carbon precursors dimethylallyl diphosphate (DMAPP) and isopentenyl diphosphate (IPP) (Figure 4.1) (Christianson, 2007). Increasingly longer polyisoprenoids are formed by the coupling of DMAPP and IPP, in a head to tail fashion to form geranyldiphosphate, farnesyl diphosphate, gernarylgeranyl diphosphate, and geranylfarnesyl diphosphate, the linear precursors of the mono-, sesqui-, di-, and sester-terpenes, respectively (Figure 4.1) (Tholl, 2006). Terpenoids are ubiquitous throughout nature and serve a multitude of specific functions in plants, animals, insects, bacteria and fungi. For example, terpenoids are critical for plant survival and account for a large number of primary metabolites, including molecules involved in photosynthesis, respiration, and membrane structure. Terpenoids also account for a wide range of secondary metabolites in plants, where they bestow unique flavors and fragrances, provide chemical defense against pests, and facilitate interactions between plants and other organisms (Aharoni, 2005; Pichersky, 2006). From a medicinal perspective, terpenoids are of great interest because many of these natural products exhibit anti-cancer, anti-malarial, and anti-microbial activities (Aharoni, 2005). Furthermore, two molecules of farnesyldiphosphate can be coupled together via a head-to-head

condensation to form squalene, the triterpene linear precursor to the steroids (Abe, 1993).

A family of enzymes known as terpenoid synthases, are responsible for the tremendous structural diversity of the terpenoids. Terpenoid synthases can be divided into two categories: class I enzymes adopt the FPP synthase α -helical fold and initiate catalysis by metal triggered ionization of the substrate diphosphate group, and class II enzymes adopt an unrelated double α -barrel fold and initiate catalysis by protonation of an epoxide ring or carbon-carbon double bond (Figure. 4.2). Class I terpenoid synthases can be further subdivided into three categories: coupling enzymes that catalyze chain elongation reactions to form increasingly longer polyisoprenoid diphosphates, coupling enzymes that catalyze irregular (i.e., non-head-to-tail) isoprenoid condensation reactions such as cyclopropanation, cyclobutanation, or branching reactions (Thulasiram, 2007), and cyclization enzymes that catalyze the conversion of linear isoprenoid substrates into single and multi-ringed hydrocarbon products (Croteau, 1985; Cane, 1990; Lesburg, 1998; Wendt, 1998). The potential diversity of carbon-carbon bond formation afforded by the flexible linear isoprenoid substrate, and the chemical potential for subsequent biosynthetic functionalization of cyclic terpenoids by cytochrome P450, monooxygenases, etc., make terpenoid biosynthesis an attractive system for engineering novel compounds (Aharoni, 2005; Yoshikuni, 2006; Austin, 2008).

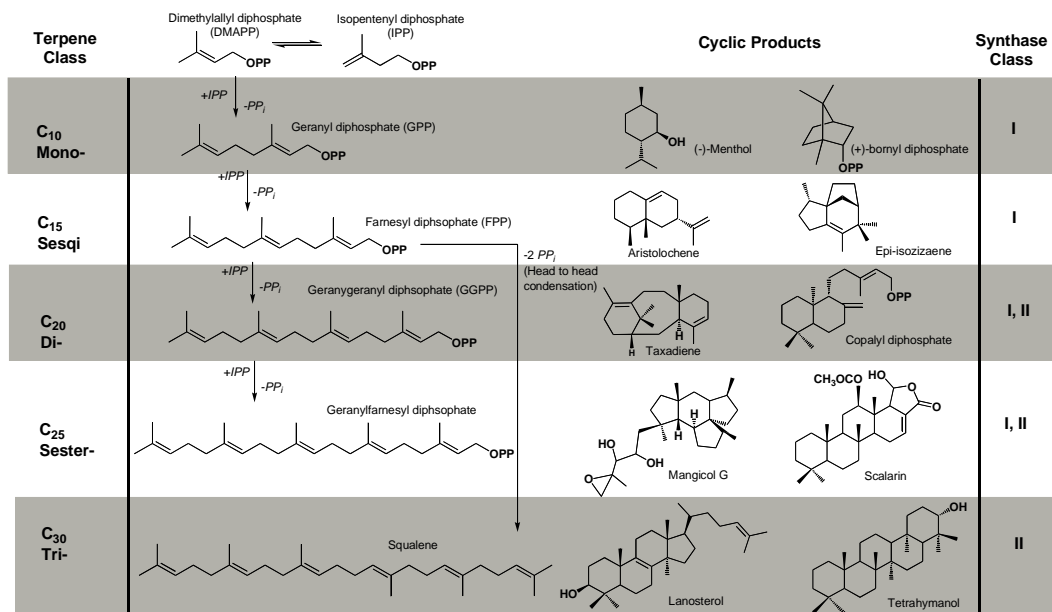


Figure 4.1. General scheme of terpenoid nomenclature and biosynthesis (OPP = diphosphate, PP_i = inorganic diphosphate).

The crystal structures of several class I terpenoid coupling and cyclization enzymes have been solved, revealing a conserved α -helical terpenoid synthase fold across the domains of life. Structures of enzyme complexes with substrates, inhibitors, and/or products have also revealed the universal conservation of a trinuclear metal cluster implicated in the molecular recognition of the substrate diphosphate group as well as the initiation of catalysis. Metal ions are coordinated by metal binding motifs on opposing helices near the mouth of the active site. The metal binding motifs are generally described as either “aspartate-rich” [**DDXX**(XX)**D/E**] or as a secondary metal binding motif “N**SE/DTE**” [(**N,D**)D(L,I,V)X(**S,T**)XXX**E**], in which boldface residues typically coordinate to catalytically obligatory Mg^{2+} or Mn^{2+} ions (where metal ligands are indicated in boldface) (Christianson, 2006). X-ray crystal structures have been instrumental in understanding the catalytic mechanisms of terpenoid synthases: the active site of each synthase provides a template that binds the flexible substrate(s) in the proper orientation and conformation so that, upon the departure of the diphosphate leaving group and resultant generation of a reactive carbocation, the active site template ensures a specific trajectory of intermolecular and intramolecular carbon-carbon bond formation in the ensuing cyclization cascade (Christianson, 2008).

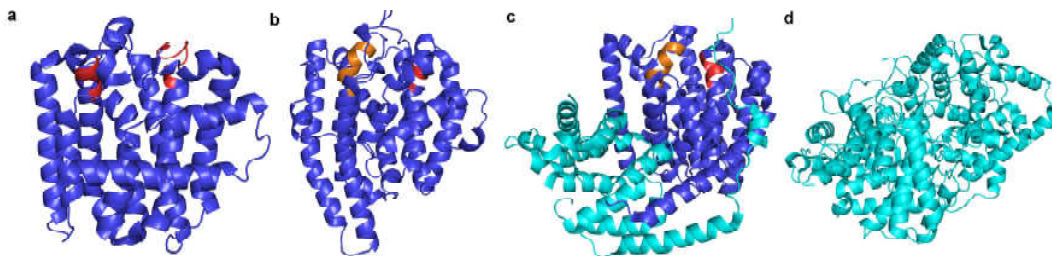


Figure 4.2. Structural similarities among various terpenoid synthases define the core class I terpenoid cyclase fold (blue). Conserved metal binding motifs are the aspartate-rich motifs (red) and “NSE/DTE” motifs (orange) highlighted in (a) *E. coli* FPP synthase (PDB code 1RQI), (b) *epi*-isozizaene synthase (PDB code 3KB9), and (c) (+)-bornyl diphosphate synthase (PDB code 1N22), which contains an additional N-terminal domain (cyan). This α -helical domain is topologically similar to the α -barrel fold of the class II terpenoid cyclases, which occurs in a double domain architecture in the triterpene cyclase (d) oxidosqualene cyclase (PDB code 1W6K).

4.2 *Streptomyces*

Streptomyces are gram-positive, filamentous, saprophytic, soil-dwelling bacteria. To-date, more than 500 species of *Streptomyces* have been identified. In addition to their central role in carbon recycling, *Streptomyces* are also characterized by their complex secondary metabolism (Challis, 2003). *Streptomyces* are a very abundant source of antibiotics, amazingly, over two-thirds of naturally derived antibiotics currently in use are produced by *Streptomyces* (Bentley, 2002). Of the thousands of secondary metabolites that are isolated from *Streptomyces* many are polyketides (ie. tetracycline) (Pickens, 2009) and aminoglycosides (ie. neomycin) (Kudo, 2009), however very few are cyclic terpenoids. One such example of a cyclic terpenoid is pentalenonelactone, an antibiotic derived from the sesquiterpene pentalenene, produced by a number of *Streptomyces* strains (Cane, 1994). Prior to this work, the X-ray crystal structure of pentalenene synthase (Lesburg, 1997) was the only known structure of a bacterial sesquiterpene cyclase. However, the structure of pentalenene synthase, determined at 2.6 Å resolution, was in an open, unliganded conformation, and therefore did not provide evidence of a conserved trinuclear metal cluster among bacterial sesquiterpene cyclases. Additional X-ray crystal structures of bacterial terpenoid cyclases are necessary to draw conclusions about the evolutionary relationships among these enzymes. Furthermore, comparative studies of the harmless *Streptomyces* genus with other members of the Actinomycetales order, for example disease causing *Mycobacterium tuberculosis* (tuberculosis) and *Mycobacterium leprae* (leprosy), may offer insight into the treatment of these pathogens (Bentley, 2002).

The complete genome of *Streptomyces coelicolor* A3(2) was sequenced in 2002, and contained a surprisingly high 7,825 predicted genes, compared with 4,289 genes in the Gram-negative bacterium *Escherichia coli*; and a predicted 31,780 in humans (Bentley, 2002). The genome contains many genes involved in secondary metabolism including polyketide synthases, chalcone synthases, non-ribosomal peptide synthetases as well as several gene clusters coding for terpene synthesis including geosmin, hopanoid, and albaflavenone biosynthesis (Bentley, 2002).

4.3 *epi-Isozizaene*

epi-Isozizaene is a member of a unique family of tricyclic sesquiterpene, like its parent hydrocarbon, zizaene (Coates, 1972), it has a highly strained ring system including a quaternary center (Figure 4.3). *epi-Isozizaene* was first observed as a natural product from the bacteria *Streptomyces coelicolor* A3(2), where it is formed by a novel terpene cyclase, *epi-isozizaene* synthase (EIZS). EIZS was first characterized by Lin and Cane (Lin, 2006), due to its 23.8% sequence identity with pentalenene synthase, a sesquiterpene cyclase isolated from *Streptomyces exfoliates* UC5319. Furthermore, the EIZS protein contained the two conserved Mg²⁺-binding motifs, an aspartate-rich motif (D⁹⁹DRHD) and the secondary metal binding motif (N²⁴⁰DLCSLPKE). EIZS catalyzes the Mg²⁺-dependant cyclization of the FPP (the linear precursor of the sesquiterpenes).

The proposed mechanism for the cyclization of farnesyl diphosphate to *epi-isozizaene* was elucidated by 1-D and 2-D NMR analysis of the products isolated from the incubation of EIZS with several isotopically labeled substrates, namely [1,1-²H₂]-

FPP, (1*R*)-[1-²H]-FPP and (1*S*)-[1-²H]-FPP (Figure 4.4) (Lin, 2006). Recently, new mechanistic insights into the *epi*-isozizaene folding pathway have been provided by computational quantum chemistry (Hong, 2009). Using the computational program GAUSSIAN03 (Frisch, 2003) to conduct a thorough analysis of carbocation intermediates and transition states from several sesquiterpene cyclization pathways, Hong and Tantillo suggest that the conformation of the bisabolyl cation attainable in the enzyme active site is a primary determinate of the structure and stereochemistry of the resultant sesquiterpenes. They report four unique bisabolyl cation conformers; each proposed to be involved in formation of a specific set of sesquiterpene products. Surprisingly, outwardly related products, for example *epi*-isozizaene and isozizaene, are proposed to be formed by different conformers of the bisabolyl cation, which vary in the orientation of the acyclic hydrocarbon chain. However, some folding pathways appear to be somewhat permissive, certain products, including *epi*-isozizaene, can be formed via more than one bisabolyl cation conformer.

Using an arbitrarily defined zero-energy bisabolyl conformer, **A0**, as a basis for their calculations, Hong and Tantillo present a detailed theoretical cyclization scheme for the formation of *epi*-isozizaene, consisting of 6 cationic intermediates, beginning from (3*R*)-nerolidyl diphosphate in the cisoid conformation (Figure 4.5). The first intermediate, **A1**, is bisabolyl cation conformer formed by 1,6-cyclization of cisoid (3*R*)-nerolidyl diphosphate. The **A1** conformer is in a productive conformation to undergo a [1,2]-hydride shift between carbons 6 and 7 via the exterior face of the acyclic chain, to form cationic intermediate **B1**. Intermediate **B1** is in an appropriate conformation to then

undergo a subsequent 6,10-cyclization via attack of the C10=C11 π -bond on the pro-*R* face of the C6 cationic center to form the acorenyl cation, **C1**. Next, a thermodynamically favorable rearrangement to the **C2** acorenyl cation conformer facilitates the direct conversion of **C2** to **E1** via a concerted cyclization and alkyl shift, circumventing the formation of a discrete secondary cation. The final intermediate, **F1**, is the result of a [1,2]-methyl shift of either C12 or C13, via a bridged nonclassical carbocation transition state. The final product, *epi*-isozizaene is formed by direct deprotonation of **F1** at C10. Although the total calculated energy change for the pathway depends on the density functional theory method chosen, the overall rearrangement beginning from the bisabolyll cation is considerably exothermic, and involves a minimal number of conformational changes between steps. Overall, the theoretical study of the *epi*-isozizaene folding pathway is consistent with that presented by Lin and Cane, and offers additional insight into several steps in the pathway, namely the concerted 3,11-cyclization and C4 alkyl shift going from **C2** to **E1**, and the approximately equal energy for the [1,2]-methyl shift of C12 or C13 to form **F1** (Hong, 2009). This study illustrates the utility of quantum calculations for studying terpenoid cyclase reaction coordinates. By expanding theoretical calculations to include contributions from an enzyme active site, we can continue to develop an understanding of terpenoid cyclase structure-function relationships and work towards engineering novel, efficient terpenoid cyclases.

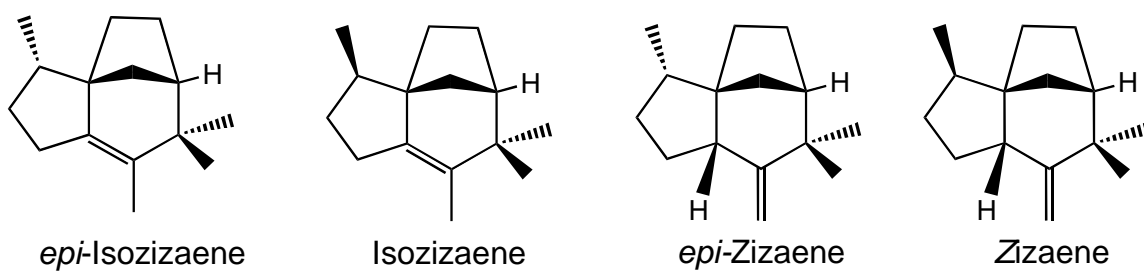


Figure 4.3. The structures of zizaene sesquiterpenes.

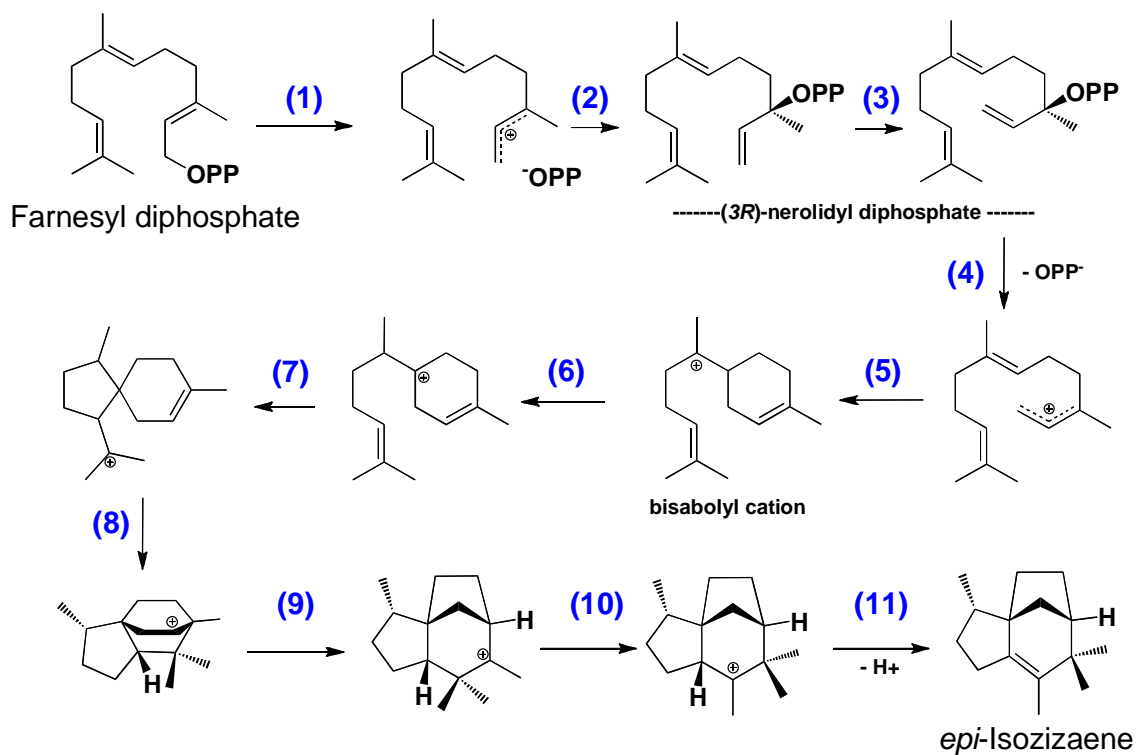


Figure 4.4. Proposed mechanism for the formation of *epi*-isozizaene from FPP by EIZS. **(1)** Ionization of FPP. **(2)** Isomerization will give *(3R)*-nerolidyl diphosphate. **(3)** Rotation about the newly generated C-2/C-3 bond generates the corresponding *cisoid* *(3R)*-nerolidyl diphosphate conformer. **(4)** Ionization of *(3R)*-nerolidyl diphosphate. **(5)** Cyclization to form bisaboyl cation. **(6)** A 1,2-hydride shift. **(7)** Spirocyclization. **(8)** Cyclization to form bisaboyl cation. **(9)** Ring contraction. **(10)** Methyl migration. **(11)** Deprotonation to yield (+)-*epi*-isozizaene.

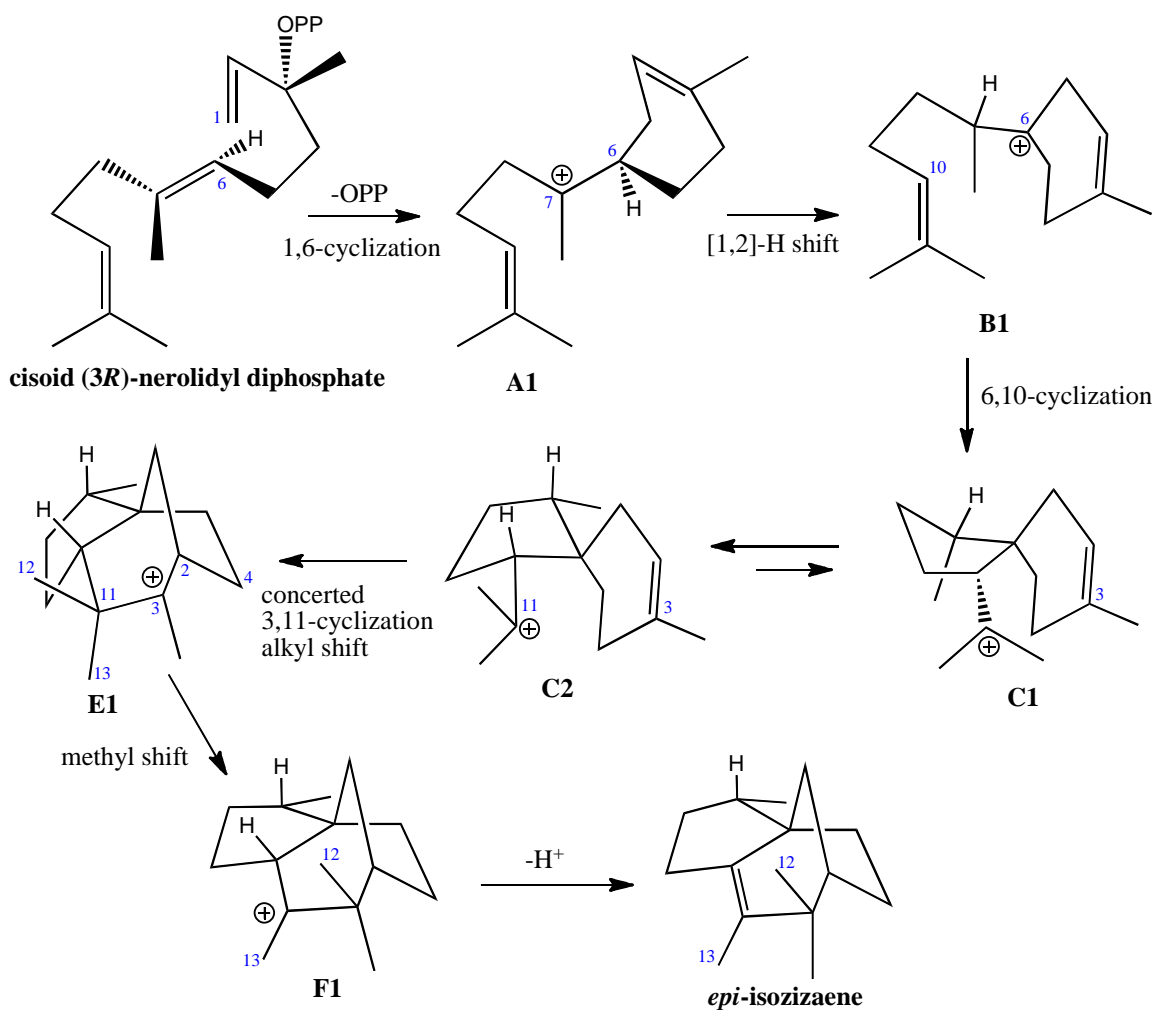


Figure 4.5. Proposed *epi*-isozizaene cyclization scheme based on quantum chemical calculations (Hong, 2009). The C12 1,2-methyl shift (**E1** to **F1** step) is preferred over a C13 1,2-methyl shift by ~2kcal/mol, however both are energetically accessible and could be affected by the active site of the enzyme.

In the *Streptomyces coelicolor* A3(2) genome, EIZS is transcriptionally coupled to cytochrome P450 170A1 (CYPA170 A1) (Zhao, 2008). Cytochrome P450 monooxygenases belong to a superfamily of heme-containing proteins that catalyze redox reactions. Specifically, P450 monooxygenases catalyze the oxidation of organic products using atmospheric dioxygen as the oxygen source and electrons from NAD(P)H, and producing a molecule of water as a side product (Bernhardt, 2006). CYP170A1 from *S. coelicolor* carries out a two-step allylic oxidation to convert *epi*-isozizaene to an epimeric mixture of albaflavenols and ultimately to the sesquiterpene antibiotic albaflavenone (Figure 4.6). Therefore, *epi*-isozizaene has been identified as an intermediate in albaflavenone synthesis, and is only detected in bacterial extracts in CYP170A1 knockout strains of *S. coelicolor*. The final product of the two-gene cluster, albaflavenone, has also been isolated from *S. albidoflavus*, and exhibits modest antibacterial activity against *Bacillus subtilis* (Zhao, 2008).

EIZS was chosen as a target for structure determination via protein X-ray crystallography in order to investigate the structural changes that occur upon the binding of the three Mg²⁺ ions and the substrate (FPP) or substrate analogues, triggering active site closure and substrate ionization. Previous work has indicated that the details of these structural changes generally differ between plant (Starks, 1997; Gennadios, 2009) and fungal (Rynkiewicz, 2001; Shishova, 2007) sesquiterpene cyclases. Until now, the mechanism of active site closure of a bacterial terpenoid cyclase has remained unknown since the only available crystal structure of a bacterial cyclase has been that of *S.*

exfoliatus UC5319 pentalenene synthase, which was determined only in an open, ligand-free conformation (Lesburg, 1997).

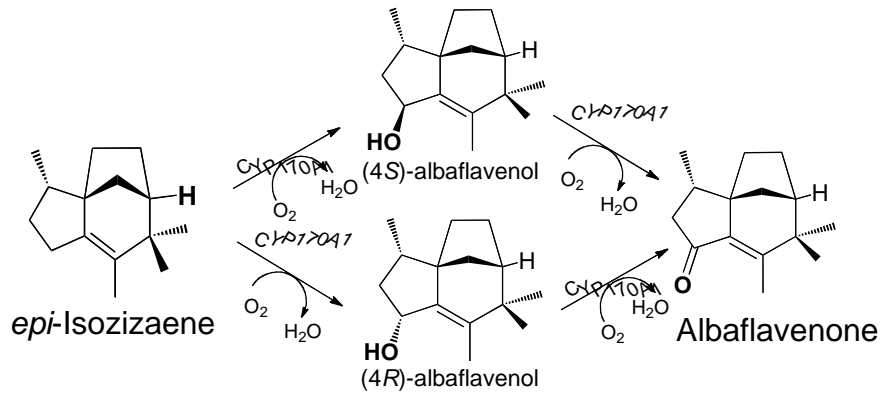


Figure 4.6. Albaflavenone biosynthetic pathway in *S. coelicolor*. EIZS catalyzes the Mg²⁺ dependent cyclization of FPP to form *epi*-isozizaene, then CYP170A1 catalyzes a two-step allylic oxidation to albaflavenone.

Chapter 5: X-Ray Crystal Structure of *epi*-Isozizaene Synthase from *Streptomyces coelicolor*

5.1 Expression and Purification

Recombinant EIZS from *Streptomyces coelicolor* A3(2) was expressed at high levels in *Escherichia coli* BL21(DE3) and purified as previously described (Lin, 2006) with minor modifications. Briefly, *E. coli* BL21(DE3) carrying pET28a(+)/SCO5222 was inoculated into Luria-Bertani (LB) medium containing kanamycin and grown overnight at 37 °C. A total of 4 L of LB/kanamycin medium was inoculated with 5 mL of the overnight seed culture, and *E. coli* was grown at 37 °C until the OD₆₀₀ reached 0.5. The temperature was reduced to 20 °C and the cells were induced with 0.2 mM isopropyl β-D-thiogalactopyranoside (IPTG) for 18 h. Cells were harvested, resuspended in 50 mL of Talon buffer A [50 mM sodium phosphate (pH 8.0), 300 mM NaCl, 20 % glycerol and 5 mM β-mercaptoethanol (BME)], supplemented with phenylmethylsulfonyl fluoride and DNase, and sonicated for 6 min using a 40 % duty cycle and power range 30 %. After three cycles of sonication, the cell lysate was clarified by centrifugation at 16000g and 4 °C for 75 min. The clarified lysate was loaded on a Talon (Clontech) Co²⁺ metal affinity resin (5 mL), and a step gradient from 0 to 200 mM imidazole in Talon buffer was applied to elute the enzyme. The fractions were analyzed using SDS-PAGE, and the most concentrated fractions were pooled and applied to a Superdex gel filtration column (HiLoad 26/60 Superdex, GE Healthcare) equilibrated in 20 mM Tris-HCl (pH 7.5), 300 mM NaCl, 10 % glycerol, 10 mM MgCl₂ and 2 mM Tris(2-carboxyethyl)phosphine (TCEP). The fractions were analyzed by SDS-PAGE and the fractions containing the

enzyme were pooled and concentrated to 8 mg/mL enzyme using a YM-10 centricon. The resulting protein preparation was >99 % pure on the basis of SDS-PAGE.

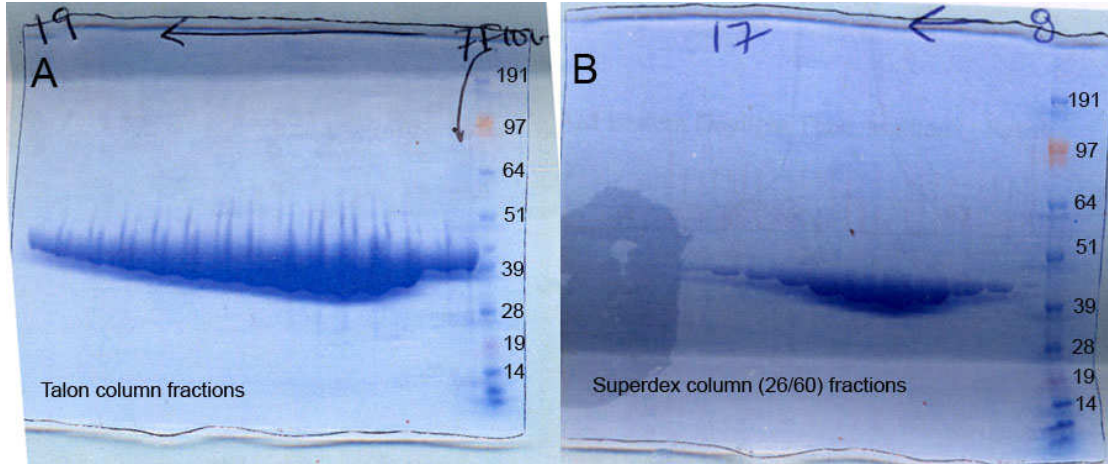


Figure 5.1. SDS-PAGE analysis of the purification of *epi-isozizaene* synthase. (A) Fractions from the Talon column: Molecular weight markers are shown on in the right lane, and the column flow-through is in the second lane from the right. The strong band is due to *epi-isozizaene* synthase (44 kDa). (B) Fractions from the Superdex (2660) column. Molecular weight markers are shown on in the right lane.

5.2 Crystallization

EIZS was crystallized by the sitting drop, and hanging drop vapor diffusion methods. Protein was freshly filtered using a 0.22 μm filter prior to setting up drops. Typically, a 4 μL drop of protein solution [8-10 mg/mL EIZS, 20 mM Tris-HCl (pH 7.5), 300 mM NaCl, 10 mM MgCl_2 , 10 % glycerol, 2 mM TCEP, 2 mM sodium pyrophosphate, 2 mM benzyltriethylammonium chloride] was added to 4 μL of precipitant solution [100 mM Bis-Tris (pH 5.5), 25-28 % polyethylene glycol 3350, 0.2 M $(\text{NH}_4)_2\text{SO}_4$] and equilibrated against a 1 mL well reservoir of precipitant solution. Initial crystallization conditions were identified using the commercially available Index screen from Hampton Research (condition #66). Crystals appeared within 2-3 days and grew to maximal dimensions of 100 μm \times 10 μm \times 10 μm . Initial crystals diffracted to 2.15 \AA at the home source, where the space group and unit cell parameters were first determined. Higher resolution data was then collected at the synchrotron. Crystals diffracted X-rays to 1.60 \AA resolution at the Advanced Photon Source, beamline NE-CAT 24-ID-C (Argonne, IL), and belonged to space group $P2_1$ with unit cell parameters $a = 53.185 \text{ \AA}$, $b = 47.374 \text{ \AA}$, $c = 75.376 \text{ \AA}$ and $\beta = 95.53^\circ$; with one monomer in the asymmetric unit, the Matthews coefficient $V_M = 2.1 \text{ \AA}^3/\text{Da}$, corresponding to a solvent content of 43 %.



Figure 5.2. Crystals of *epi*-isozizaene synthase.

5.3 Structure Determination with Heavy Atoms

5.3.1 Introduction

A crystal structure can be described as the best-fit model to a contour map of the electron density throughout the unit cell. The electron density, $\rho(x,y,z)$, can be represented by equation 5.1, a periodic function represented by a Fourier series using the

$$\rho(x,y,z) = 1/V \sum_h \sum_k \sum_l F_{hkl} e^{-2\pi i(hx+ky+lz)} \quad \mathbf{5.1}$$

structure factors, F_{hkl} . A structure factor is a complete description of a diffracted X-ray. Since an X-ray can be described as a wave, it has a frequency, amplitude, and phase. To calculate the electron density according to equation 5.1, and thus build a model of the protein structure, the frequency, amplitude and phase of each reflection, hkl , must be known. When an X-ray diffracts off a protein crystal, the frequency of the X-ray does not change, therefore the frequency is equal to that of the X-ray source. The amplitude of the diffracted X-ray is proportional to (I_{hkl}) , the square root of the measured intensity of reflection hkl (Rhodes, 2000). However, the phase of the diffracted X-ray is unknown, and is the missing piece of the crystallography puzzle that must be found to complete a crystal structure determination.

Protein crystallographers have developed several methods to obtain the “lost” phase information from the diffraction data. A very commonly used method is known as molecular replacement (MR), when the phases of a similar, known structure are used as the initial phase estimates for the new structure. This method works best when a homologous structure is known, typically the phasing model has >25 % sequence identity and an r.m.s. deviation of <2.0 Å between the C α atoms of the model and the new

structures (Taylor, 2003). When MR is not successful, other methods that rely upon the incorporation of heavy atoms to the crystal lattice are used. When heavy atoms are incorporated into a crystal lattice, measurable intensity changes are observed in the diffraction pattern, with respect to the native diffraction pattern. Upon incorporation of the heavy atoms, if the crystal lattices remain isomorphous, except for the presence of the heavy atom, the isomorphous difference between the reflection amplitudes of native and derivative crystals can be used as an estimate of the heavy-atom structure-factor amplitude, which can be used to determine the positions of the heavy-atoms. Once the phases of the heavy atoms are known, a first approximation of the phase of the native protein can be made. When a single heavy atom derivative is used, this method is known as single isomorphous replacement (SIR). This method can be extended, when necessary, to include information from more than one heavy atom derivative, and is known as multiple isomorphous replacement (MIR) (Taylor, 2003). SIR and MIR are typically very effective methods for structure determination, however in some cases the incorporation of heavy atoms into native crystals causes unexpected protein conformational changes. These conformational changes are sometime evident from changes in the unit cell parameters, however in other cases non-isomorphism is not apparent, yet critically hampers structure solution.

In addition to their mere presence, heavy atoms can additionally aid in structure solution due to their ability to anomalously scatter X-rays. Heavy atoms absorb X-rays at specific wavelengths, known as the element's absorption edge, resulting in unequal intensities between the hkl and $-h -k -l$ reflections. The anomalous difference can be

used to determine the location of the anomalous scattering atom in the unit cell, thus providing phase information. Single-wavelength anomalous diffraction (SAD), can be advantageous since it does not require an isomorphous native data set. Also, anomalous scattering methods can be extended include multiwavelength anomalous diffraction (MAD), where several data sets are collected from the same crystal at different wavelengths. Furthermore, isomorphous and anomalous heavy atom methods can be combined, for example in SIRAS (single isomorphous replacement with anomalous scattering), and MIRAS (multiple isomorphous heavy-atom replacement using anomalous scattering) (Taylor, 2003). Some of the most common heavy atoms used for phasing are mercury, platinum, gold, and uranium. These metals are typically incorporated into the crystal lattice by soaking preformed crystals in a precipitant solution supplemented with a millimolar concentration of the metal ion, where they typically interact with surface cysteine, histidine, and methionine residues (Rhodes, 2000). Selenium can also be used as a heavy atom for phasing, and can be incorporated into the protein itself by expressing recombinant protein in media supplemented with selenomethionine (Hendrickson, 1990). Recently, xenon has also been explored as a heavy atom for phasing. Protein crystals are briefly pressurized with xenon gas prior to flash cooling, causing xenon atoms to be trapped in hydrophobic pockets within the protein, which can be used for phasing (Evans, 2003).

5.3.2 Results

A number of heavy metal compounds were tested to identify a good heavy metal derivative for phasing. These heavy metal compounds included thimerosal

(C₉H₉HgNaO₂S), mercuric chloride (HgCl₂), mersalyl acid (C₁₃H₁₈HgNO₆), ethylmercuryl chloride (C₂H₅HgCl), methylmercuric acetate (C₃H₆HgO₂), ytterbium chloride (YbCl₃), PIP (di- μ -iodobis(ethylenediamine)diplatinum (II) nitrate) (C₄H₁₆I₂N₆O₆Pt₂), dipotassium platinum hexachloride (K₂PtCl₆), samarium chloride (SmCl₂), manganese chloride (MnCl₂), and sodium bromide (NaBr). Additionally, selenomethionine containing EIZS protein was expressed and crystallized.

Of the many heavy metal derivatives that were investigated, two mercury compounds were identified which were used for successful phasing. Mercury derivatives were prepared by soaking crystals in precipitant buffer supplemented with 1 mM ethyl mercury chloride or methylmercuric acetate for 1-2 days, followed by flash cooling. Diffraction data were collected at the absorption edge of mercury (1.0548 Å) on beamline X29 at the National Synchrotron Light Source (NSLS). A sample diffraction image of a methylmercuric acetate derivatized EIZS crystal is shown in Figure 5.3. Complete data sets of ethylmercury chloride, and methylmercuric acetate derivatized EIZS crystals were collected. Crystals diffracted X-rays to 1.90 Å and 2.10 Å, respectively and belonged to space group *P2*₁ with unit cell parameters in Table 5.1; data reduction was achieved with Denzo and Scalepack (Otwinowski, 1997), derivative data sets were scaled anomalously.

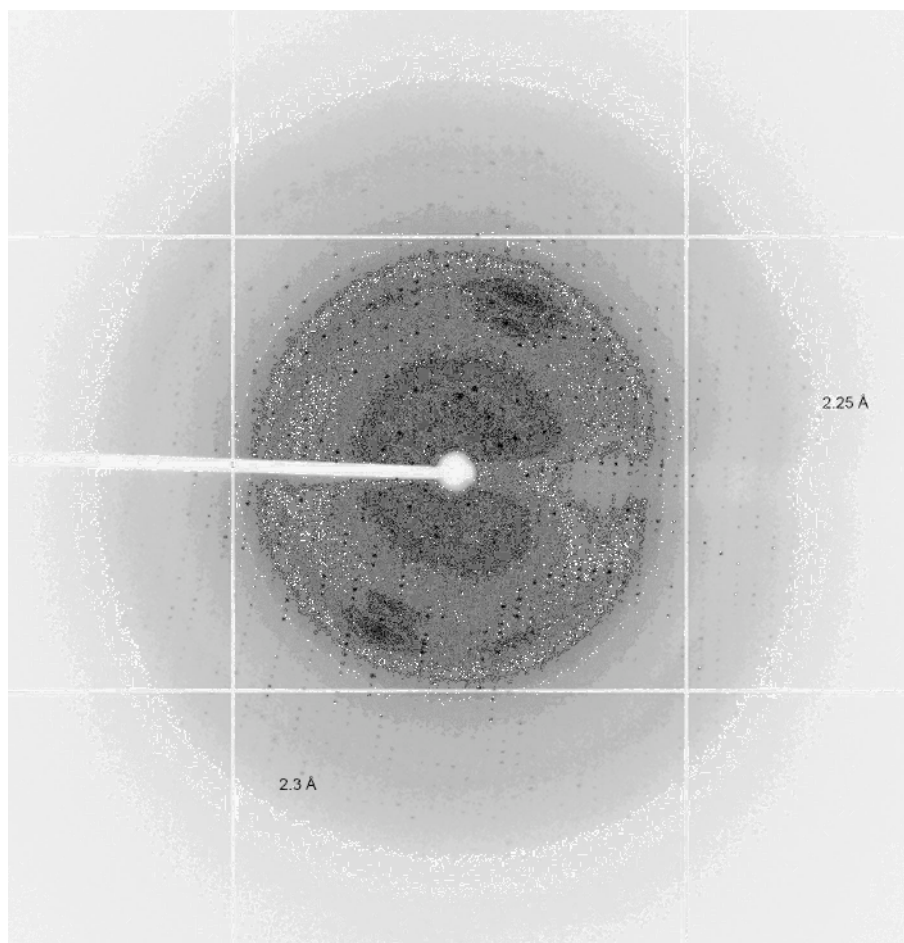


Figure 5.3. Diffraction pattern of methylmercuric acetate derivatized EIZS crystals, collected at beamline X29, National Synchrotron Light Source. The crystal diffracted to 2.1 Å resolution.

Table 5.1. Data collection statistics for native and mercury derviatized EIZS

crystals. The space group of all crystals is $P2_1$.

EIZS Structure	Native	Ethylmercury chloride	Methylmercuric acetate
Data			
Wavelength, Å	1.0080	1.0548	1.075
Resolution, Å	1.60	1.90	2.10
Completeness*, %	92.9 (100)	99.4 (98.6)	99.6 (97.8)
Redundancy*	3.6 (3.6)	5.8 (5.1)	3.5 (3.0)
$R_{\text{sym}}^{*\dagger}$	0.058 (0.208)	0.075 (0.325)	0.108 (0.390)
I/σ	34 (8.3)	18 (4.4)	10 (2.7)
Unit Cell Parameters			
a, Å	53.185	51.693	52.931
b, Å	47.374	46.549	46.462
c, Å	75.376	75.551	75.689
β , (°)	95.53	92.338	94.992

*Values in parentheses refer to the highest shell

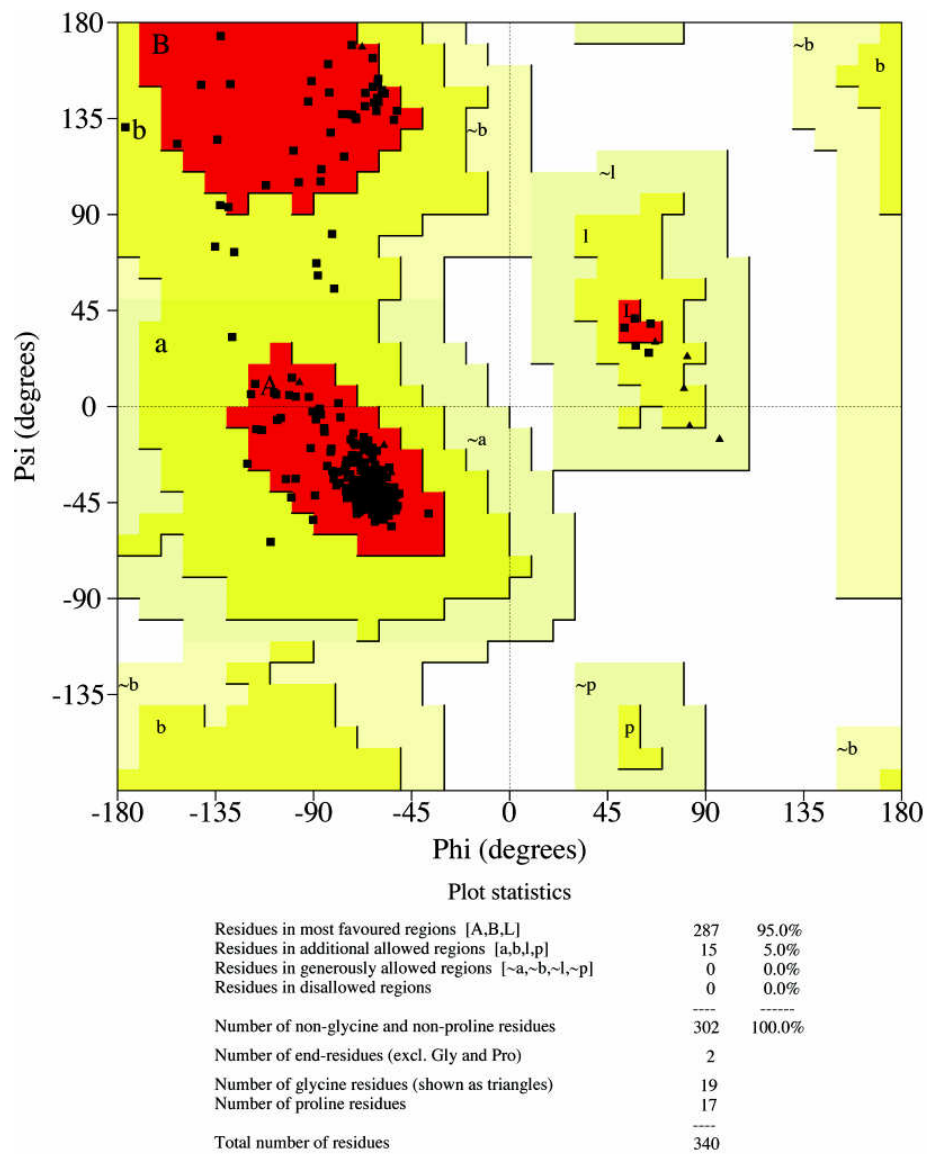
$\dagger R_{\text{sym}} = \sum |I_h - \langle I_h \rangle| / \sum I_h$, where $\langle I_h \rangle$ is the average intensity over symmetry equivalent reflections.

Difference Patterson maps generated with anomalous scattering data revealed large peaks corresponding to mercury. Initial attempts to solve the crystal structure using single or multiple isomorphous replacement were not successful due to apparent non-isomorphism between the native and mercury-derivatized crystals. The program autoSHARP (Vonrhein, 2007) was ultimately used for structure determination. AutoSHARP is an automated structure solution program built around the phasing program SHARP (Bricogne, 2003), the density modification program SOLOMON (Abrahams, 1996), and the ARP/wARP package (Langer, 2008) for automated model building and refinement using REFMAC (Skubak, 2004). The following data is required to begin an autoSHARP run, the type of heavy atom present in the crystal, the number of expected heavy atom sites, the space group and wavelength of data collection. The file format of the reflection data input can be either MTZ or SCALEPACK. Additionally, the contents of the asymmetric can be input as a molecular weight, number of residues or the amino acid sequence, the later enables automatic building of the structure.

The initial electron density map was phased by single-wavelength anomalous dispersion (SAD) of with the 1.90 Å data set of the ethylmercury chloride derivative using autoSHARP. Four mercury sites were identified, and the phase correlation coefficient calculated by autoSHARP for this data set was 0.466, with anomalous phasing power 0.447. Following density modification and automatic building, approximately 50% of the protein residues were built into the electron density map. At this point, the molecular model of EIZS was refined against the 1.60 Å resolution data set collected from the native crystal instead of the mercury derivative. Molecular replacement using

the partially built model was achieved with PHASER (Storoni, 2004). Iterative cycles of refinement and manual model building using CNS (Brunger, 2007), O (Jones, 1991), COOT (Emsely, 2004), and PHENIX (Adams, 2002) allowed for the assembly of the complete protein model. Individual atomic B-factors were utilized during refinement. Buffer molecules, ions, water, glycerol, the benzyl triethylammonium cation (BTAC), PP_i, and a sulfate ion (hydrogen bonded to R163, H164, R220 and R226) were included in later cycles of refinement. Data reduction and refinement statistics are recorded in Table 5.2. A total of 340 of 381 residues (A16-N355) are present in the final model, as the N- and C- termini are disordered. According to the program PROCHECK (Laskowski, 1993), 95 % of the residues were located in the most favorable regions of the Ramachandran plot, and no residues are in disallowed regions (Figure 5.4).

In order to investigate the mercury binding sites of the ethyl mercury chloride derivatized crystals used for structure determination, refinement of the 1.90 Å-resolution structure was completed. Four mercury binding sites were identified by strong peaks in a Bijvoet difference Fourier map adjacent to C68, C213, C243 and C283. Iterative cycles of refinement and manual model building were achieved with PHENIX and COOT, respectively. Ions and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized, and the mercury, chloride and sulfur atoms were refined anisotropically. Data reduction and refinement statistics are recorded in Table 5.2. A total of 323 of 381 residues are present in the final model, as there was no interpretable density for the A251-L267 loop as well as the N- and C- termini.



Based on an analysis of 118 structures of resolution of at least 2.0 Angstroms and R-factor no greater than 20%, a good quality model would be expected to have over 90% in the most favoured regions.

Figure 5.4. Ramachandran plot of the refined EIZS-Mg²⁺₃-PP_i-BTAC structure.

Table 5.2. Refinement Statistics for wild-type EIZS complexes

EIZS Structure:	Mg ²⁺ ₃ -PP ₁ - BTAC complex	Hg ²⁺ ₄ complex
Data		
Wavelength, Å	1.0080	1.0548
Resolution, Å	40 - 1.60	50 - 1.90
Unique reflections	46,113	28,446
Completeness*, %	92.9 (100)	99.4 (98.6)
Redundancy*	3.6 (3.6)	5.1 (5.1)
R _{sym} * [†]	0.058 (0.208)	0.075 (0.325)
Refinement		
R _{cryst} /R _{free} [‡]	0.158 / 0.189	0.169 / 0.202
r.m.s.d. bonds, Å	0.012	0.005
r.m.s.d. angles, °	1.4	0.8
r.m.s.d. dihedral angles, °	21	15
No. of atoms		
Protein atoms	2858	2512
Solvent atoms	431	335
Ligand atoms	55	13
Ramachandran plot		
Allowed, %	94.7	95.2
Additionally allowed, %	5.3	4.8

*Values in parentheses refer to the highest shell

[†] $R_{\text{sym}} = \sum |I_h - \langle I_h \rangle| / \sum I_h$, where $\langle I_h \rangle$ is the average intensity over symmetry equivalent reflections.

[‡] $R_{\text{cryst}} = \sum ||F_{\text{obs}}| - |F_{\text{calc}}|| / \sum |F_{\text{obs}}|$, where summation is over the data used for refinement. R_{free} was calculated as for R_{cryst} by using 5% of the data that was excluded from refinement.

5.4 Structure of EIZS-Mg²⁺-PP_i-BTAC Complex

At 1.60 Å resolution, the crystal structure of this complex is the highest resolution structure of any terpenoid cyclase determined to date (Aaron, 2010). EIZS adopts the class I terpenoid synthase α -helical fold and consists of a bundle of 10 α -helices, designated A-J (Figure 5.5), in which the 20 Å-deep active site is defined mainly by helices C, D, G, H, and J. Among the terpenoid cyclases of known structure, EIZS is structurally most similar to pentalenene synthase (r.m.s. deviation = 3.3 Å for 304 C α atoms). EIZS crystallizes as a monomer, consistent with dynamic light scattering measurements indicating that the protein is a monomer in solution (Figure 5.6).

The electron density map clearly reveals 3 Mg²⁺ ions, PP_i, and a BTAC molecule bound in the active site (Figure 5.7 (a)). The side chain of D99 in the aspartate-rich motif **D**⁹⁹DRHD¹⁰³ coordinates to Mg²⁺_A and Mg²⁺_C with *syn,syn*-bidentate geometry, while Mg²⁺_B is chelated by the “NSE” motif **N**²⁴⁰DLCSLP**K**E²⁴⁸ (boldface indicates Mg²⁺ ligands). Each Mg²⁺ ion is coordinated with octahedral geometry, with nonprotein coordination sites occupied by the oxygen atoms of PP_i and by water molecules. The PP_i anion also accepts hydrogen bonds from the side chains of R194, K247, R338, and Y339 (Figure 5.7 (b)). It is likely that similar metal coordination and hydrogen bond interactions are normally formed with the diphosphate group of the substrate FPP in the precatalytic Michaelis complex. These interactions stabilize the closed active site conformation that sequesters FPP from bulk solvent and triggers ionization to initiate the electrophilic cyclization cascade.

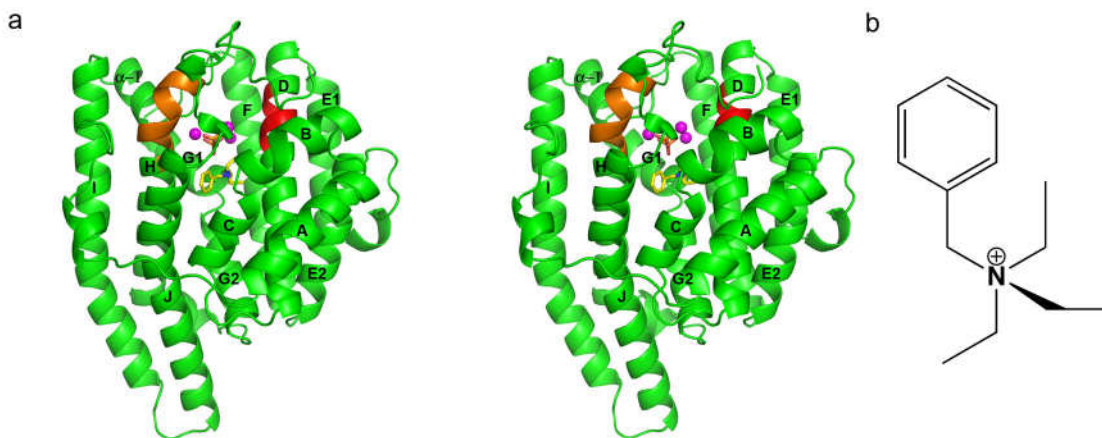


Figure 5.5. (a) Ribbon plot of the EIZS-Mg²⁺₃-PP_i-BTAC complex showing the aspartate-rich motif (red) and the NSE motif (orange) flanking the mouth of the active site. The Mg²⁺ ions are shown as magenta spheres, PP_i and BTAC molecules are color coded by atom [carbon (yellow), nitrogen (blue), oxygen (red), and phosphate (orange)]. Helices are labeled according to the convention first established for FPP synthase (Tarshis, 1994). (b) The quaternary ammonium group of the benzyltriethylammonium cation (BTAC) serves as a mimic of a carbocation intermediate in catalysis.

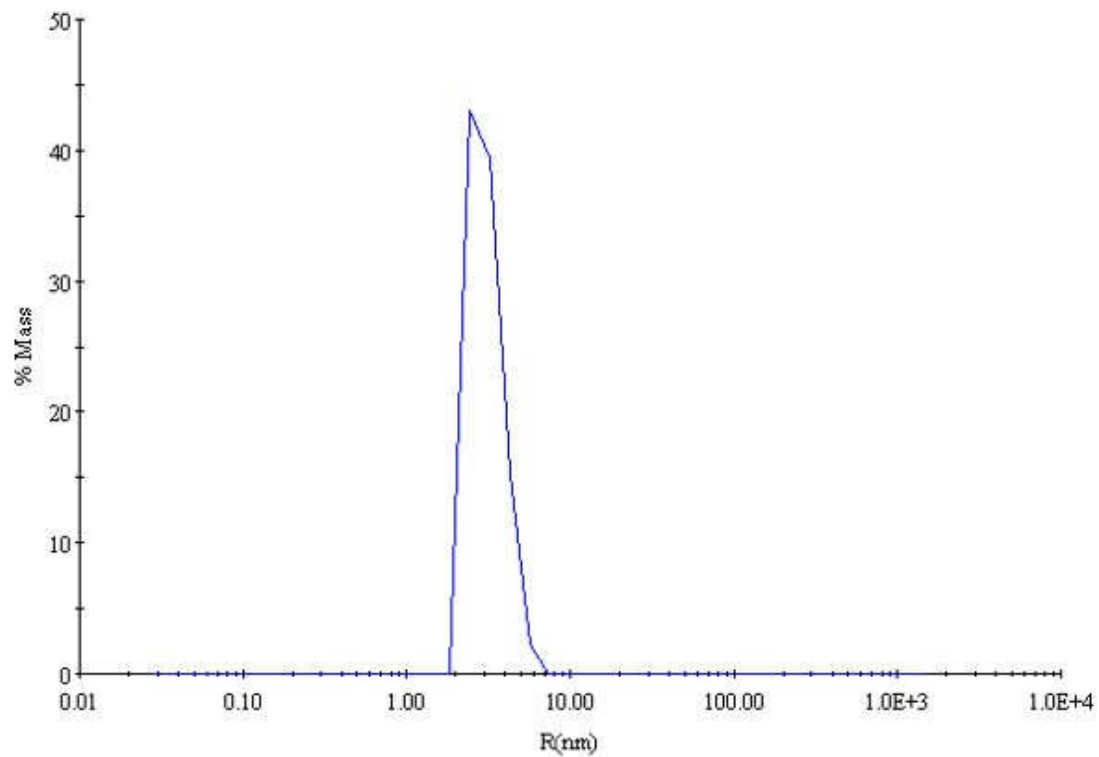


Figure 5.6. Particle size distribution of EIZS sample, measured by dynamic light scattering. Results indicated a monomeric sample, with an average particle radius of 3 nm.

Crystals of EIZS form upon addition of BTAC (Figure 5.5 (b)), commonly used in synthetic organic chemistry as a phase transfer catalyst. As a hydrophobic cation, BTAC mimics the bisabolyl carbocation intermediate proposed for the EIZS-catalyzed cyclization mechanism (Lin, 2006). The positively charged quaternary ammonium group of BTAC appears to be stabilized by long-range electrostatic interactions with the PP_i anion (the shortest N---O separation is 4.1 Å), as well as cation- π interactions with the aromatic side chains of F95, F96 and F198 (N---ring centroid separations range 4.7 – 5.4 Å) (Figure 5.6 (a)). Such cation- π interactions are proposed to play a critical role in stabilizing the highly reactive carbocation intermediates found in all enzyme-catalyzed terpenoid cyclizations (Jenson, 1997; Lesburg, 1997; Ma, 1997). The binding of BTAC in the active site of EIZS demonstrates that the bacterial terpenoid cyclase active site can accommodate and stabilize a positively charged ligand resembling the positively charged carbocation intermediates of the normal cyclization cascade. Although determined at a much lower resolution of 2.85 Å, the recently reported structure of the trichodiene synthase- Mg^{2+}_3 - PP_i -BTAC complex provides an example of the stabilization of positively charged ligands in the active site of a fungal terpenoid cyclase (Vedula, 2007).

5.5 Structure of EIZS- Hg^{2+}_4 Complex

To date, we have been unable to prepare crystals of wild-type EIZS in a completely ligand-free state, since the additives BTAC and PP_i required for crystallization are not readily dialyzed out of the crystals. However, soaking crystals of the EIZS- Mg^{2+}_3 - PP_i -BTAC complex with ethyl mercury chloride displaces 3 Mg^{2+} ions,

PP_i, and BTAC to yield the 1.90 Å resolution structure of the EIZS-Hg²⁺₄ complex in which all Hg²⁺ ions bind remotely from the active site (Figure 5.8 (a)). In this complex, Hg²⁺_A is coordinated by C68, Hg²⁺_B is coordinated by C283 (the electron density is best interpreted as 2 alternate positions), Hg²⁺_C is coordinated by C243, and Hg²⁺_D is coordinated by C213 (2 alternate positions). Surprisingly, however, the now ligand-free active site is not open and empty, as it is in all other ligand-free terpenoid cyclase structures. Instead, helix G, which forms one side of the active site (and contains R194, a residue that donates a hydrogen bond to PP_i in the ligand-bound structure), bends by ~110° to occupy the location formerly occupied by Mg²⁺₃-PP_i (Figure 5.8 (c)). Additionally, helix D moves ~1.5 Å outward, helix J moves ~4 Å outward, while helix H, which was bent in the ligand-bound structure to enable Mg²⁺_B chelation by the NSE motif, becomes straight and also moves outward (Figure 5.8 (c)). The positions and conformations of helices C, E, and F remain essentially unchanged, and there is no interpretable electron density for helix I or the loop (A251-L267) connecting it to helices H and J. Salt bridges between helix G and helices D and F stabilize the closed ligand-free structure: R194 makes a salt-bridge with E175 (helix F) and R195 makes a salt bridge with D103 (helix D) (Figure 5.8 (c)). Although these residues are found in pentalenene synthase as R173, F174, D84 and E152, a comparable closed ligand-free conformation was not observed by Lesburg and colleagues (Lesburg, 1997). Although the EIZS-Hg²⁺₄ structure illustrates a unique terpenoid cyclase structure, the role of the Hg²⁺ ions in dislodging the Mg²⁺₃-PP_i-BTAC complex to result in the closed ligand-free conformation of EIZS is unclear. However, this structure does not appear to be biologically relevant, as

this structural transition would result in an inactive enzyme and does not likely reflect the state of the wild type enzyme in the absence of substrate.

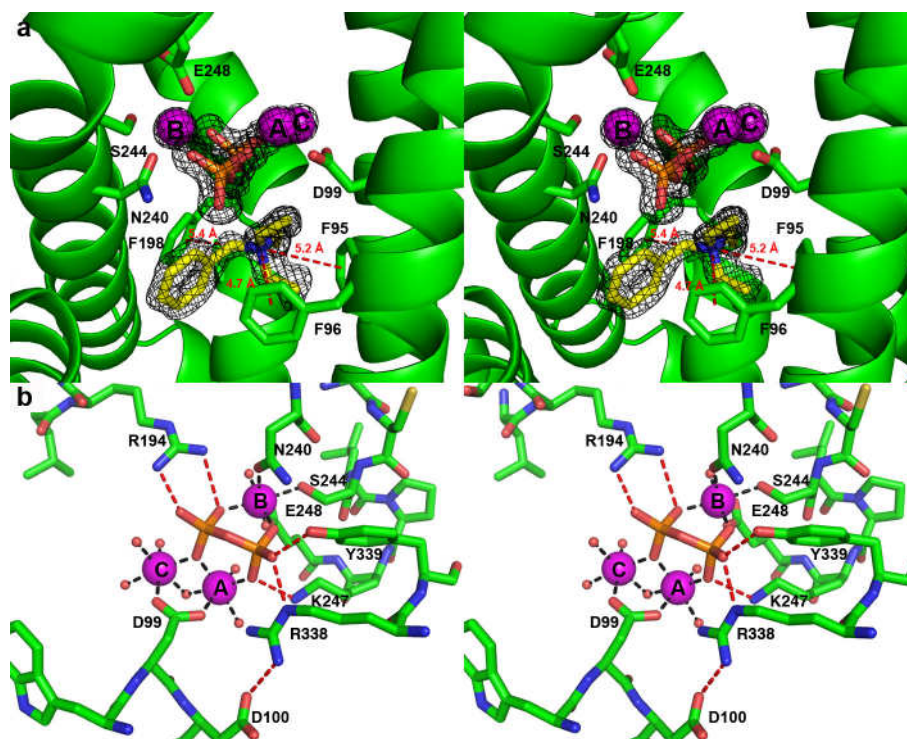


Figure 5.7. Active site of EIZS-Mg²⁺₃-PP_i-BTAC complex. (a) Simulated annealing omit maps (black) of the PP_i anion, Mg²⁺ ions, and BTAC, contoured at 5σ. Note the cation-π interactions between the positively charged quaternary ammonium group of BTAC and the aromatic rings of F95, F96 and F198 (red dashed lines). (b) Metal coordination interactions (black dashed lines) and hydrogen bond interactions (red dashed lines) in the EIZS-Mg²⁺₃-PP_i complex.

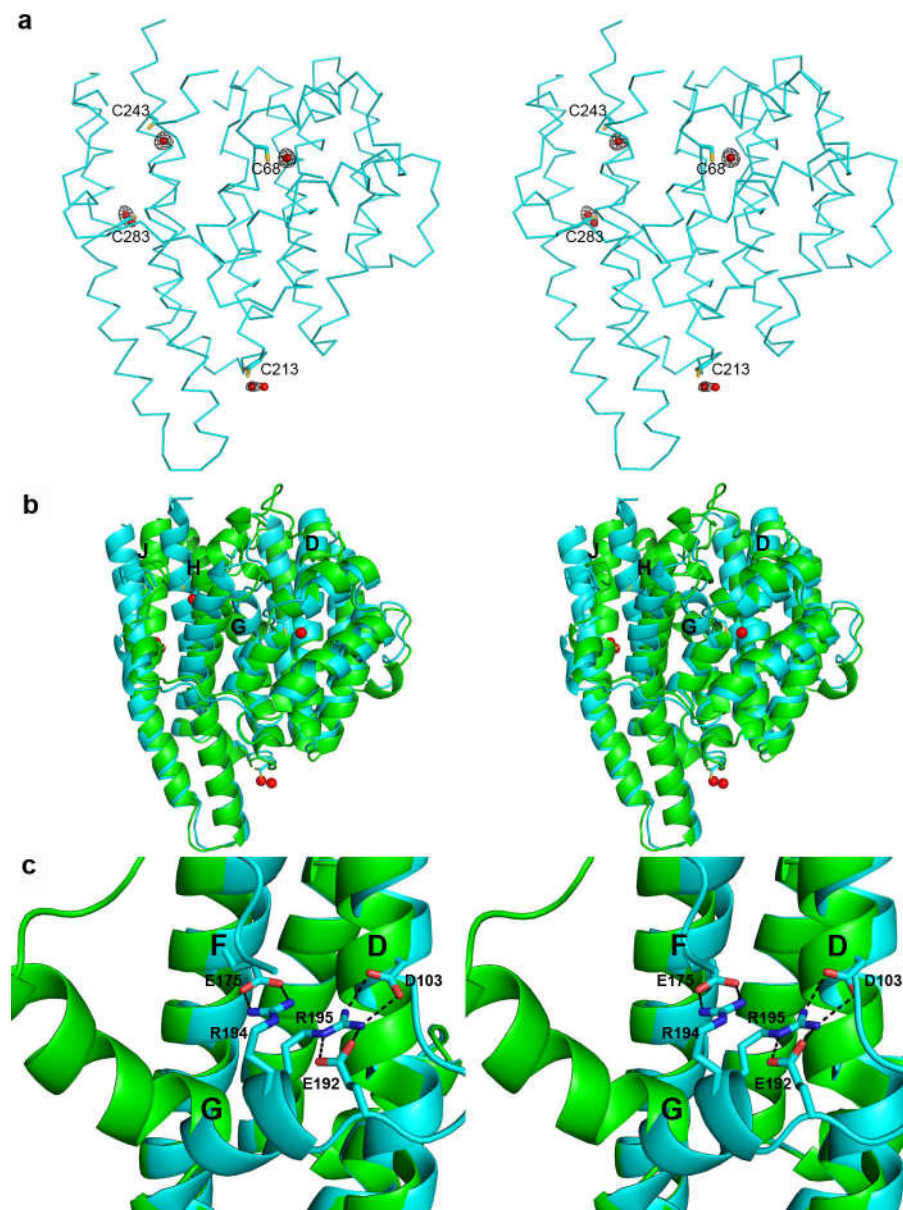


Figure 5.8. Structural changes between EIZS $\text{Mg}^{2+}_3\text{-PP}_i\text{-BTAC}$ and Hg^{2+}_4

Complexes. (a) Ribbon diagram of EIZS- Hg^{2+}_4 complex. Anomalous Fourier map (black) contoured to 6σ , indicates the location of Hg atoms (red spheres). (b) Superposition of ribbon plots of EIZS- $\text{Mg}^{2+}_3\text{-PP}_i\text{-BTAC}$ (green) and EIZS- Hg^{2+}_4 (cyan; Hg^{2+} ions appear as red spheres). Helices D, G, H and J, which undergo the largest changes, are labeled. (c)

Salt bridges (black dashes) between helix G and helices D and F stabilize the closed ligand-free structure.

Chapter 6: X-ray Crystal Structure of D99N *epi*-Isozizaene Synthase and Implications for Substrate Recognition

6.1 Introduction

Interestingly, the detailed structural changes that occur between open and closed active site conformations differ between fungal and plant terpenoid cyclases. For example, the r.m.s. deviation between ligand-free and $\text{Mg}^{2+}_3\text{-PP}_i$ -complexed trichodiene synthase from *F. sporotrichioides* is 1.4 Å (Rynkiewicz, 2001), whereas that of bornyl diphosphate synthase from *S. officinalis* (culinary sage) is only 0.6 Å for the catalytic domain (Whittington, 2002). Until now, it had not been possible to study the structural changes between the open and closed active site conformations of bacterial terpenoid cyclases because pentalenene synthase had been determined only in the ligand-free state (Lesburg, 1997). Since the residues that interact with Mg^{2+} ions or PP_i in *epi*-isozizaene synthase (EIZS) are conserved in pentalenene synthase, superposition of the closed, ligand-bound structure of EIZS with ligand-free pentalenene synthase enables an approximated comparison of the open-to-closed structural transition in bacterial terpenoid cyclases. Alignment of the two structures reveals a very similar alignment of the metal binding motifs, with pentalenene synthase helices D (“aspartate-rich” motif) and H (“NSE” motif) being 1.5 Å further apart than in EIZS. We hypothesize that upon binding of Mg^{2+}_3 and substrate or PP_i , the active site of pentalenene synthase undergoes a change to a closed conformation comparable to that observed for the EIZS- $\text{Mg}^{2+}_3\text{-PP}_i\text{-BTAC}$

complex. To further explore structural changes resulting from the dissociation of Mg^{2+} - PP_i -BTAC from the active site of EIZS, we attempted to determine the structure of EIZS in the absence of Mg^{2+} and PP_i , however we were unable to obtain crystals of the WT enzyme without Mg^{2+} , PP_i and BTAC. Furthermore, we were not able to remove, via soaking, Mg^{2+} - PP_i -BTAC from crystals of the EIZS- Mg^{2+} - PP_i -BTAC complex.

Therefore, an alternative approach was taken to investigate the open structure of EIZS. The D99N amino acid substitution severely decreases catalytic efficiency (Lin, 2009) by compromising the *syn,syn*-bidentate coordination of Mg^{2+}_A and Mg^{2+}_C (Figure 5.7). Thus this single-site mutant enables the structural study of an unliganded conformation of EIZS.

6.2. Experimental Methods

6.2.1 Site-Directed Mutagenesis, Expression and Purification

The D99N single site-specific mutation was introduced into the EIZS wild-type plasmid using primers 1 and 2 as follows (lower case letters represent the mutant codon introduced): primer 1, 5'GGT TCT TCG TCT GGa acG ACC GTC ACG AC-3', primer 2, 5'-GTC GTG ACG GTC gtt CCA GAC GAA GAA CC-3'. The optimal reaction mixture for PCR amplification of the insert was 100 ng of each forward and reverse primer, 3 μ L of 10 mM dNTP mix, 100 ng plasmid, 5 μ L of *Pfu* turbo polymerase buffer, and 1 unit of *Pfu* turbo polymerase diluted with water to a final volume of 50 μ L. Optimal PCR conditions required initial denaturation of the reaction mixture at 95 °C for 5 min, addition of polymerase followed by thirty cycles (1 min denaturation at 95 °C, 1

min annealing at 60 °C, 8 min extension at 72 °C), and a final 10 min extension at 72 °C followed by a final hold at 4 °C. One μL of DpnI was added to the PCR mixture and incubated at 37 °C for 1 h to digest the template. PCR products were transformed into XL1-Blue cells for DNA isolation and sequencing. DNA was purified (Qiagen mini-prep kit) from cultures from single colonies, and DNA sequencing (DNA Sequencing Facility, University of Pennsylvania) confirmed incorporation of the mutation. Mutant proteins were expressed and purified using the same procedures as described for the wild-type enzyme (Section 5.1).

6.2.2 Crystallization and Structure Determination

The D99N EIZS mutant was crystallized by the hanging drop vapor diffusion method. Crystals formed with a precipitant solution at a slightly higher pH than the wild type crystals [100 mM Bis-Tris (pH 6.0), 25-28% polyethylene glycol 3350, 0.2 M $(\text{NH}_4)_2\text{SO}_4$], and were improved by successive rounds of micro-streak seeding using D99N crystals as the seed stock. Crystals diffracted to 1.9 Å resolution at APS beamline NE-CAT 24-ID-C and belonged to space group $P2_12_12_1$ with unit cell parameters $a = 41.144$ Å, $b = 81.952$ Å, $c = 106.693$ Å. The Matthew's coefficient (V_M) of 2.04 Å³/Da, and solvent content of 40 % suggest the presence of one monomer per asymmetric unit. Molecular replacement calculations were performed with PHASER (Storoni, 2004) using the atomic coordinates of native EIZS (less ligand and solvent atoms) as a search probe. Iterative cycles of refinement and manual model building were achieved with PHENIX (Adams, 2002) and COOT (Emsely, 2004), respectively. Sulfate and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized.

Data reduction and refinement statistics are recorded in Table 6.1. A total of 316 of 381 residues (P18-E335) are present in the final model, as the N- and C- termini are disordered.

Table 6.1. Data Collection and Refinement Statistics for D99N EIZS.

EIZS Structure:	D99N (ligand-free)
Data	
Wavelength, Å	0.9795
Resolution, Å	50 – 1.90
Unique reflections	28,694
Completeness*, %	97.6 (95.6)
Redundancy*	3.3 (3.0)
$R_{\text{sym}}^{*\dagger}$	0.089 (0.634)
Refinement	
$R_{\text{cryst}}/R_{\text{free}}^{\ddagger}$	0.162 / 0.207
r.m.s.d. bonds, Å	0.009
r.m.s.d. angles, °	1.1
r.m.s.d. dihedral angles, °	16
No. of atoms	
Protein atoms	2638
Solvent atoms	316
Ligand atoms	5
Ramachandran plot	
Allowed, %	94.4
Additionally allowed, %	5.6

*Values in parentheses refer to the highest shell

$\dagger R_{\text{sym}} = \sum |I_h - \langle I_h \rangle| / \sum I_h$, where $\langle I_h \rangle$ is the average intensity over symmetry equivalent reflections.

$\ddagger R_{\text{cryst}} = \sum |F_{\text{obs}}| - |F_{\text{calc}}| / \sum |F_{\text{obs}}|$, where summation is over the data used for refinement. R_{free} was calculated as for R_{cryst} by using 5% of the data that was excluded from refinement.

6.3. Structure of D99N EIZS and Implications for Substrate Recognition

Unexpectedly, D99N EIZS crystallizes in a unique packing arrangement with respect to wild-type EIZS, resulting in a different space group and unit cell parameters. Both crystal forms contain one monomer per asymmetric unit, however wild-type EIZS crystallized in space group $P2_1$, with two monomers in the unit cell (Figure 6.1 (a)), whereas D99N EIZS crystallized in space group $P2_12_12_1$, with four equivalent monomer positions in the unit cell (Figure 6.1. (b)).

In the closed, ligand-bound conformation of EIZS only the first aspartate of the D⁹⁹DRHD motif coordinates to Mg²⁺_A and Mg²⁺_C; substitution of a neutral asparagine residue for the negatively-charged aspartate residue at position 99 is sufficient to disrupt the assembly and stability of the trinuclear metal cluster required for substrate recognition and activation. The crystal structure of D99N EIZS reveals the complete absence of Mg²⁺₃-PP_i-BTAC in the active site. Moreover, the active site of D99N EIZS adopts an open conformation (Figure 6.2) without any conformational changes of helix G. Therefore, we conclude that the conformational change of helix G shown in Figure 5.8 is somehow caused by Hg²⁺ binding and is not biologically relevant.

Comparison of EIZS-Mg²⁺₃-PP_i-BTAC and D99N EIZS structures enables a structural understanding of the molecular recognition of substrate by EIZS, and the ensuing structural changes that result in closing of the active site upon binding of substrate and Mg²⁺ ions. The crystal structures are particularly informative in that they reveal two biologically relevant active site conformations: a closed, ligand-bound

conformation and an open, ligand-free conformation, respectively. Alignment of the two structures reveals ligand-induced conformational changes of helix H and loop H- α -1, as well as the J-K loop (which is completely disordered in D99N EIZS). The overall r.m.s. deviation between wild-type and D99N EIZS structures is 1.6 Å for 318 C α atoms. The D99N EIZS structure suggests that in the absence of PP_i, or a PP_i-containing substrate, the C-terminal residues, beginning at E335, become disordered, which results in an open active site. Upon Mg²⁺ and PP_i (or PP_i-containing substrate) binding, the C-terminus becomes ordered and closes the active site and blocks the solvent from entering the active site. Corresponding structural changes generally accompany active site closure in fungal (Shishova, 2008) and plant (Whittington, 2002) terpenoid cyclases. The template for FPP cyclization is fully formed in the closed active site conformation of EIZS as well as all other class I terpenoid cyclases.

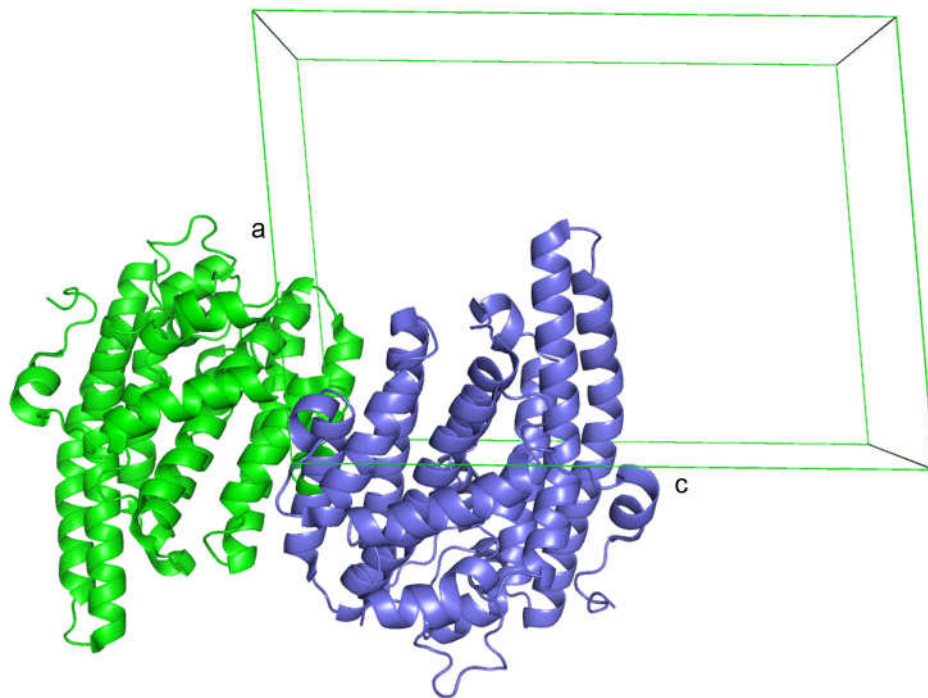


Figure 6.1. (a) The unit cell of EIZS-Mg²⁺₃-PP₁-BTAC crystals viewed along the b axis. The crystals belong to space group *P2*₁, the asymmetric unit is a monomer, and the two positions are related in the unit cell by a screw axis along b.

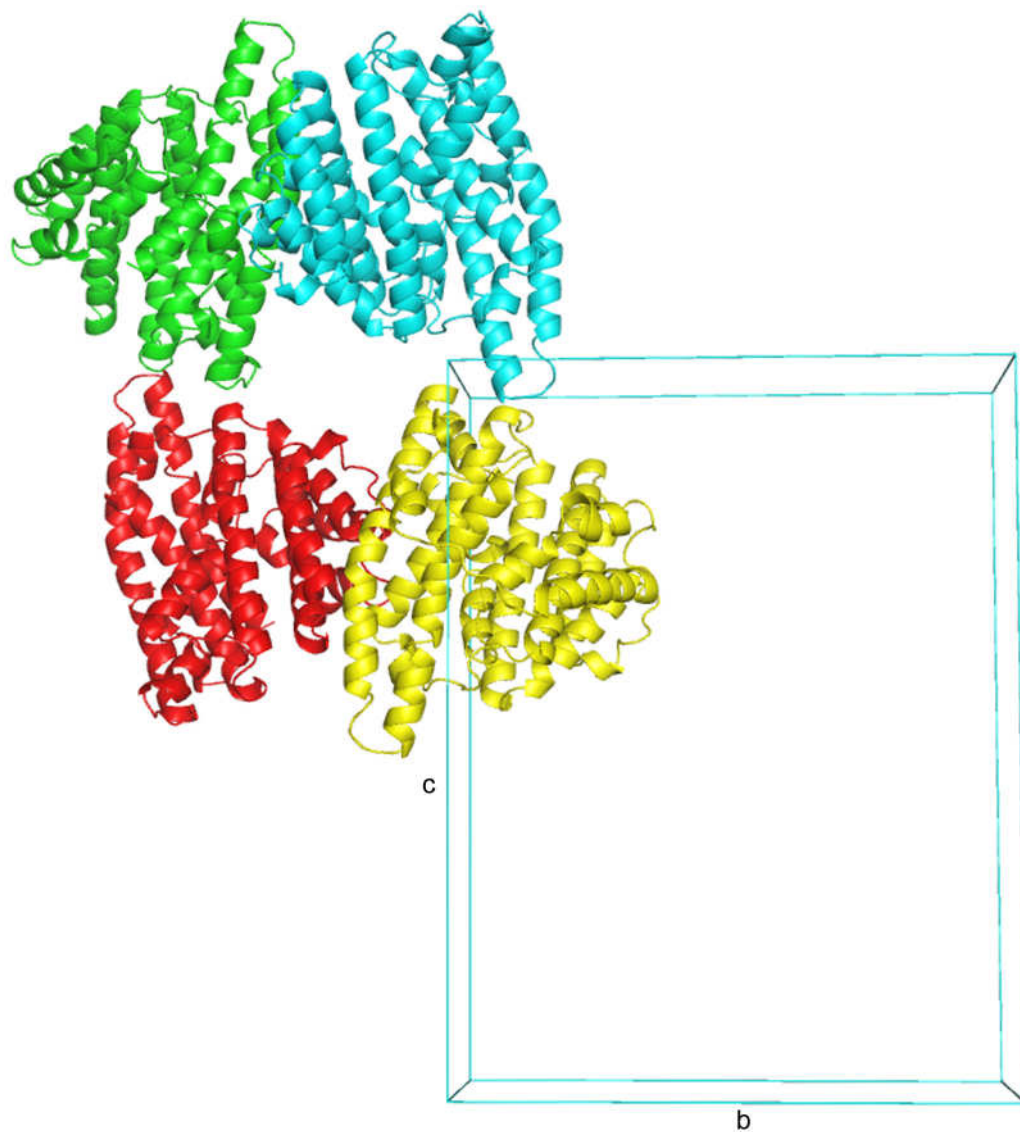


Figure 6.1. (b) The unit cell of D99N EIZS crystals viewed along the a axis. The crystals belong to space group $P2_12_12_1$, the asymmetric unit is a monomer. There are four equivalent positions in the unit cell, which possesses three perpendicular twofold screw axes.

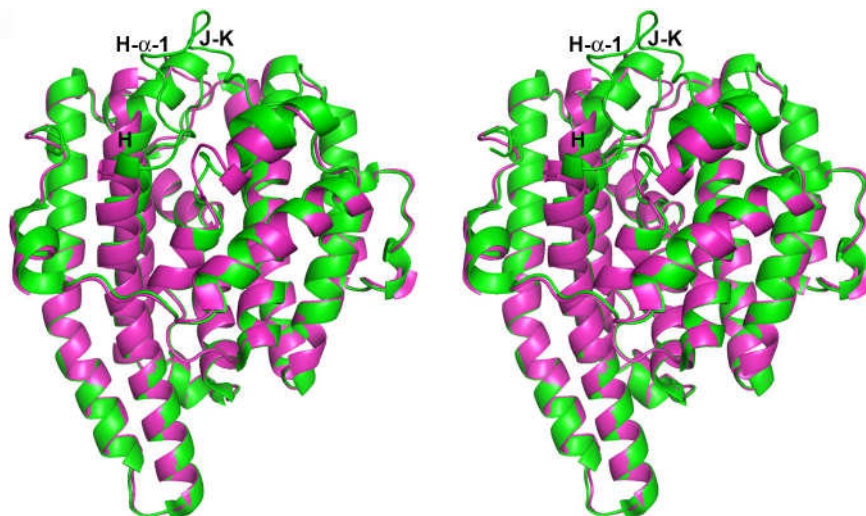


Figure 6.2. A stereoview of a superposition of the structures of EIZS-Mg²⁺₃-PP_i-BTAC complex (green) and D99N EIZS (purple), illustrating the structural changes in helix H and the H- α -1 and J-K loops that accompany active site closure.

The newly determined open and closed structures of EIZS enable a comparison of substrate recognition and active site closure amongst bacterial, fungal and plant terpenoid cyclases. Although the first and third aspartate residues in the aspartate-rich metal binding motif coordinate to Mg^{2+}_A and Mg^{2+}_C in the plant terpenoid cyclases containing a complete trinuclear metal cluster such as (+)-bornyl diphosphate synthase (Figure 6.3 (c)) (Whittington, 2002), limonene synthase (Hyatt, 2007), and (+)- δ -cadinene synthase (Gennadios, 2009), only the first aspartate residue in the aspartate-rich motif coordinates to Mg^{2+}_A and Mg^{2+}_C in Mg^{2+}_3 -PP_i complexes with the fungal cyclases trichodiene synthase (Rynkiewicz, 2001) and aristolochene synthase (Figure 6.3 (b)) (Shishova, 2007). EIZS is similar to the fungal cyclases in that only D99 coordinates to Mg^{2+}_A and Mg^{2+}_C . A critical role for D99 in metal complexation is reflected in the dramatic losses of catalytic activity measured for the D99N and D99E mutants (Lin, 2009). Although D100 of EIZS does not directly interact with the Mg^{2+} ions or PP_i, it does accept a hydrogen bond from R338, which also donates a hydrogen bond to PP_i (Figure 6.3 (a)). Site-directed mutagenesis reveals that the D100N mutant has lost >95 % activity compared to the native enzyme (Lin, 2009), suggesting that the D100N mutation disrupts the D100-R338-PP_i hydrogen bond network. In aristolochene synthase, the second aspartate in the aspartate-rich motif, D91, similarly stabilizes a hydrogen bond network with R314 and PP_i (Shishova, 2007). Surprisingly, in (+)-bornyl diphosphate synthase, the second aspartate, D352, is involved in a hydrogen bond network with R314 and PP_i (Whittington, 2002), illustrating the importance of the arginine residue in stabilizing the closed conformation. Furthermore, a second conserved arginine residue makes a hydrogen bond to the opposite end of the diphosphate moiety, R194 in EIZS, R175 in aristolochene

synthase and R493 in (+)-bornyl diphosphate synthase, suggesting that the bacterial, fungal, and plant cyclases share the same molecular strategy for linking the molecular recognition of the substrate diphosphate group with the mechanism of active site closure. Two additional PP_i coordinating interactions are conserved amongst EIZS and aristolochene synthase; K247 and Y339 in EIZS donate H-bonds to diphosphate oxygen atoms and are conserved as K226 and Y315 in aristolochene synthase. A higher degree of conservation amongst active site residues suggests bacterial and fungal terpenoid cyclases derive from a more recent common ancestor than plant terpenoid cyclases.

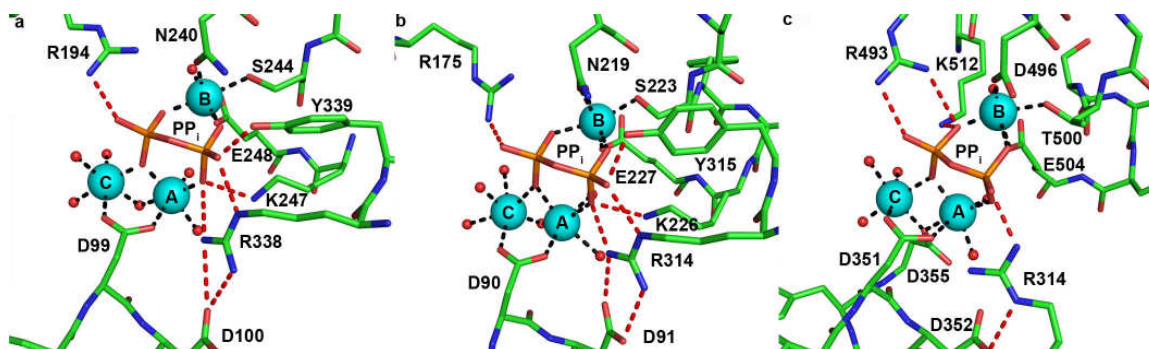


Figure 6.3. Conservation of Mg^{2+} - PP_i and -diphosphate binding motifs among bacterial and fungal terpenoid cyclases. Metal coordination (black) and hydrogen bond (red) interactions with phosphate(s) are indicated. (a) Bacterial sesquiterpene cyclase *S. coelicolor* epi-isozizaene synthase- Mg^{2+} - PP_i complex (PDB code 3KB9); (b) Fungal sesquiterpene cyclase *A. terreus* aristolochene synthase- Mg^{2+} - PP_i complex (PDB code 2OA6); (c) Plant monoterpene cyclase *S. officinalis* (+)-bornyl diphosphate synthase- Mg^{2+} - PP_i complex (PDB code 1N22; metal ions are labelled according to the convention first established for trichodiene synthase).

Chapter 7: Structural and Biochemical Studies of the Active Site of EIZS

7.1 Introduction

In general, the permissiveness and promiscuity of terpenoid cyclases vary, both in terms of the substrates they accept and the product(s) they generate. These properties are dictated by the three-dimensional contour of the fully formed template in the closed active site conformation. Many terpenoid cyclases, such as *A. terreus* aristolochene synthase (Felicetti, 2004), are high-fidelity cyclases that generate one product exclusively. However other cyclases, such as EIZS, generate one major product and minor quantities of one or more alternative products. Detailed gas chromatography-mass spectrometry (GC-MS) analysis of the organic products that result from incubation of WT EIZS with FPP reveal the promiscuity of the EIZS template. Specifically, 79 % of the total sesquiterpene product mixture is *epi*-isozizaene, and the remaining 21 % is identified as a mixture of β -farnesene (5 %), sesquisabinene-A (3 %), zizaene (9 %), α -cedrene (2 %), sesquiphellandrene (1 %), and 2 % of an unidentified sesquiterpene (Lin, 2009). The structural basis for such mechanistic promiscuity is presumably rooted in how well the active site contour enforces the correct regiochemistry and stereochemistry for cyclization and eventual quenching of the carbocation intermediates by chaperoning the conformations of reactive intermediates. Intriguingly, the conformations and orientations of such intermediates may not reflect the original conformation and orientation of the substrate if the template is somewhat permissive (Hong, 2009). A more permissive template allows alternative premature quenching of on-pathway intermediates or off-pathway conformations that lead to the formation of aberrant products.

Manipulation of the cyclization template by site-directed mutagenesis can redirect the biosynthetic trajectory of a terpenoid cyclase. This result can be achieved by modification of active site contour residues (Yoshikuni, 2006) or of residues that are more distant from the active site (O'Maille, 2008). With EIZS, two different strategies have been employed to manipulate the cyclization template: mutagenesis of metal-binding motif residues, and residues that directly contribute to forming the unique active-site contour.

Mutagenesis of the conserved Mg^{2+} -binding domains severely compromises catalytic efficiency. Single-site metal-binding motif mutants D99E, D100N, N240D, S244A, and E248D all retain less than 5% of WT activity, however GC-MS analysis of their respective organic products indicates these mutations have only a modest effect on the cyclization template, resulting in slightly altered relative proportions of *epi*-isozizaene and alternate sesquiterpene side-products. Specifically, *epi*-isozizaene accounts for 62 to 91 % of the relative sesquiterpene products of these mutants, and only one additional sesquiterpene side product is identified, α -neocallitropsene (3 % D99E, 3 % N240 D, <1 % E248D), thus the fidelity of *epi*-isozizaene biosynthesis is not significantly compromised. Indeed, certain amino acid substitutions involving the Mg^{2+} -binding residues, such as D100N, N240D, S244A, and E248D, actually lead to increased proportions of *epi*-isozizaene and lower levels of the alternative sesquiterpene products, although with significantly decreased overall catalytic efficiency (Lin, 2009).

It is hypothesized that the structural and stereochemical diversity achieved by the terpenoid cyclases is the result of an enzyme active site that merely chaperones a multi-step intermolecular cyclization cascade. As such, a terpenoid cyclase active site is primarily lined with hydrophobic and aromatic residues, which serve to guide the cyclization cascade by assisting in the stabilization of carbocation intermediates (Christianson, 2008). Aromatic residues are able to stabilize carbocation intermediates via charge-quadrupole (cation- π) interactions. The active site of EIZS has several aromatic residues, namely F95, F96, F198, W203, F332, H333, and W325. As a first step in exploring the importance of active site aromatic residues for catalysis and product diversity in EIZS, we have prepared the F96A, F198A, and W203F mutants. These residues were selected for mutation based on their proximity to the BTAC cation that was observed to coordinate in the active site of the WT EIZS-Mg²⁺-PP_i-BTAC complex presented in Chapter 5 (Figure 5.6). Furthermore, the effect of mutating aliphatic residues in the active site was also explored. Specifically L72, A236 and V329 were chosen for investigation, and each site was selectively mutated to investigate the following single-site EIZS mutants: L72V, L72I, A236G, A236V, V329A and V329L to explore the effects of subtle changes to the active site contour on enzyme activity and product distribution.

7.2 Experimental Methods

7.2.1 Site-Directed Mutagenesis, Expression and Purification

The following EIZS single-site mutants were investigated: F96A, F198A, W203A, W203F, L72V, L72I, A236G, A236V, V329A and V329L. Single site-specific

mutations were introduced into the EIZS wild-type plasmid using forward and reverse primers for each respective mutant (Table 7.1). The optimal reaction mixture for PCR amplification of the insert was 100 ng of each forward and reverse primer, 3 μ L of 10 mM dNTP mix, 100 ng plasmid, 5 μ L of *Pfu* turbo polymerase buffer, and 1 unit of *Pfu* turbo polymerase diluted with water to a final volume of 50 μ L. Optimal PCR conditions required initial denaturation of the reaction mixture at 95 °C for 5 min, addition of polymerase followed by thirty cycles (1 min denaturation at 95 °C, 1 min annealing at 60 °C, 8 min extension at 72 °C), and a final 10 min extension at 72 °C followed by a final hold at 4 °C. One μ L of Dpn1 was added to the PCR mixture and incubated at 37 °C for 1 h to digest the template. PCR products were transformed into XL1-Blue cells for DNA isolation and sequencing. DNA was purified (Qiagen mini-prep kit) from cultures from single colonies, and DNA sequencing (DNA Sequencing Facility, University of Pennsylvania) confirmed incorporation of the mutations. Mutant proteins were expressed and purified using the same procedures as described for the wild-type enzyme (Section 5.1). The W203A mutant expressed poorly, and could not be successfully purified, therefore a more conservative W203F mutant was investigated. The L72I, A236V and V329L EIZS mutants did not express as well as their respective counterparts L72V, A236G and V329A, and were not investigated further.

Table 7.1. EIZS mutagenic primer sequences. Lower case letters represent the mutant codon introduced.

Protein		Primers
F96A	Forward	5'-CTA CAG CGC GTG GTT Cgc aGT CTG GGA CGA CCG TC-3'
	Reverse	5'-GAC GGT CGT CCC AGA Ctg cGA ACC ACG CGC TGT AG-3'
F198A	Forward	5'-GAA CTG CGC CGG CTC ACG gca GCG CAC TGG ATC TGG AC-3'
	Reverse	5'-GTC CAG ATC CAG TGC GcT gcC GTG AGC CGG CGC AGT TC-3'
W203A	Forward	5'-GTT CGC GCA CTG CAT Cgc aAC CGA CCT GCT GGA G-3'
	Reverse	5'-CTC CAG CAG GTC GGT tgc CAT CCA GTG CGC GAA C-3'
W203F	Forward	5'-GTT CGC GCA CTG GAT Ctt tAC CGA CCT GCT GG-3'
	Reverse	5'-GCT CCA GCA GGT CGG Taa aGA TCC AGT GCG CG-3'
L72I	Forward	5'-CTG TGC TAC ACG GAC att ATG GCG GGC TAC TAC C-3'
	Reverse	5'-GGT AGT AGC CCG CCA Taa tGT CCG TGT AGC ACA G-3'
L72V	Forward	5'-CTG TGC TAC ACG GAC gtg ATG GCG GGC TAC TAC-3'
	Reverse	5'-GTA GTA GCC CGC CAT cac GTC CGT GTA GCA CAG-3'
A236G	Forward	5'-GTC AGG AAT TCg gcG CCT GGT ACA AC-3'
	Reverse	5'-GTT GTA CCA GGC gcc GAA TTC CTG AC-3'
A236V	Forward	5'-GAG TCA GGA ATT Cgt gGC CTG GTA CAA CGA C-3'
	Reverse	5'-GTC GTT GTA CCA GGC cac GAA TTC CTG ACT C-3'
V329A	Forward	5'-CTG GTT CAG TTC Cgc gTA CTG GTT CCA CC-3'
	Reverse	5'-GGT GGA ACC AGT Acg cGG AAC TGA ACC AG-3'
V329L	Forward	5'-GAA CTG GTT CAG TTC Cct gTA CTG GTT CCA CCA CG-3'
	Reverse	5'-CGT GGT GGA ACC AGT Aca gGG AAC TGA ACC AGT TC-3'

7.2.2 Crystallization and Structure Determination of EIZS active site mutants

7.2.2.1 F198A EIZS-Mg²⁺₃-PP_i-BTAC complex.

The F198A EIZS mutant was crystallized by the hanging drop vapor diffusion method with the same conditions used to crystallize the wild-type enzyme, but with successive rounds of micro-streak seeding using native crystals as the seed stock. Crystals diffracted to 1.64 Å resolution at NSLS beamline X29 and belonged to space group $P2_1$ with unit cell parameters $a = 53.241$ Å, $b = 47.179$ Å, $c = 75.568$ Å and $\beta = 95.57^\circ$. Molecular replacement calculations were performed with PHASER (Storoni, 2004) using the atomic coordinates of native EIZS (less ligands and solvent atoms) as a search probe. The electron density clearly revealed the F198A mutation. Iterative cycles of refinement and manual model building were achieved with PHENIX and COOT, respectively. Ions, PP_i, BTAC, and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized. Data collection and refinement statistics for the F198A EIZS-Mg²⁺₃-PP_i-BTAC complex are listed in Table 7.2.

7.2.2.2 L72V EIZS-Mg²⁺₃-PP_i-BTAC complex.

The L72V EIZS mutant was crystallized by the hanging drop vapor diffusion method with the same conditions used to crystallize the wild-type enzyme. Crystals diffracted to 2.10 Å resolution at NSLS beamline X29 and belonged to space group $P2_1$ with unit cell parameters $a = 52.977$ Å, $b = 47.236$ Å, $c = 75.084$ Å and $\beta = 95.66^\circ$. Molecular replacement calculations were performed with PHASER (Storoni, 2004) using the atomic coordinates of native EIZS (less ligands and solvent atoms) as a search probe. The electron density clearly revealed the L72V mutation. Iterative cycles of refinement

and manual model building were achieved with PHENIX and COOT, respectively. Ions, PP_i, BTAC, and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized. Data collection and refinement statistics for the L72V EIZS-Mg²⁺₃-PP_i-BTAC complex are listed in Table 7.2.

7.2.2.3 A236G EIZS-Mg²⁺₃-PP_i-BTAC complex.

The A236G EIZS mutant was crystallized by the hanging drop vapor diffusion method with the same conditions used to crystallize the wild-type enzyme. Crystals diffracted to 1.76 Å resolution at APS beamline NE-CAT-ID-C and belonged to space group *P*2₁ with unit cell parameters *a* = 53.137 Å, *b* = 47.220 Å, *c* = 75.211 Å and β = 95.51°. Molecular replacement calculations were performed with PHASER (Storoni, 2004) using the atomic coordinates of native EIZS (less ligands and solvent atoms) as a search probe. The electron density clearly revealed the A236G mutation. Iterative cycles of refinement and manual model building were achieved with PHENIX and COOT, respectively. Ions, PP_i, BTAC, and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized. Data collection and refinement statistics for the A236G EIZS-Mg²⁺₃-PP_i-BTAC complex are listed in Table 7.2.

7.2.2.4 V329A EIZS-Mg²⁺₃-PP_i-BTAC complex.

The V329A EIZS mutant was crystallized by the hanging drop vapor diffusion method with the same conditions used to crystallize the wild-type enzyme. Crystals diffracted to 1.95 Å resolution at NSLS beamline X29 and belonged to space group *P*2₁ with unit cell parameters *a* = 53.214 Å, *b* = 47.485 Å, *c* = 75.283 Å and β = 95.50°.

Molecular replacement calculations were performed with PHASER (Storoni, 2004) using the atomic coordinates of native EIZS (less ligands and solvent atoms) as a search probe. The electron density clearly revealed the V329A mutation. Iterative cycles of refinement and manual model building were achieved with PHENIX and COOT, respectively. Ions, PP_i, BTAC, and water molecules were included in later cycles of refinement. Individual atomic B-factors were utilized. Data collection and refinement statistics for the V329A EIZS-Mg²⁺₃-PP_i-BTAC complex are listed in Table 7.2.

Table 7.2. EIZS active site mutant data collection and refinement statistics.

EIZS Structure:	F198A Mg ²⁺ ₃ -PP _i - BTAC complex	L72V Mg ²⁺ ₃ -PP _i - BTAC complex	A236G Mg ²⁺ ₃ -PP _i - BTAC complex	V329A Mg ²⁺ ₃ -PP _i - BTAC complex
Data				
Wavelength, Å	1.075	0.9795	0.9795	0.9795
Resolution, Å	50 – 1.64	50 – 2.10	50 – 1.76	50 – 1.95
Unique reflections	45,831	22, 011	37, 050	26,839
Completeness*, %	99.8 (100)	99.9 (99.2)	99.2 (98.2)	97.3 (97.4)
Redundancy*	3.6 (3.5)	3.6 (3.5)	3.3 (3.1)	4.2 (3.7)
R _{sym} * [†]	0.062 (0.238)	0.104 (0.301)	0.071 (0.339)	0.116 (0.432)
Refinement				
R _{cryst} /R _{free} [‡]	0.156/0.190	0.156/0.206	0.159/0.203	0.152/0.201
r.m.s.d. bonds, Å	0.015	0.007	0.009	0.007
r.m.s.d. angles, °	1.6	1.050	1.167	1.014
r.m.s.d. dihedral angles, °	18	15	17	17
No. of atoms				
Protein atoms	2858	2788	2801	2812
Solvent atoms	467	272	357	376
Ligand atoms	31	31	31	42
Ramachandran plot				
Allowed, %	95.0	94.0	94.7	94.0
Additionally allowed, %	5.0	6.0	5.3	6.0

*Values in parentheses refer to the highest shell

[†] $R_{\text{sym}} = \sum |I_h - \langle I_h \rangle| / \sum I_h$, where $\langle I_h \rangle$ is the average intensity over symmetry equivalent reflections.

[‡] $R_{\text{cryst}} = \sum |F_{\text{obs}} - F_{\text{calc}}| / \sum F_{\text{obs}}$, where summation is over the data used for refinement. R_{free} was calculated as for R_{cryst} by using 5% of the data that was excluded from refinement.

7.2.3 Radioactive substrate kinetic assay of EIZS mutants

EIZS mutants were assayed as previously described (Lin, 2006) in 50 mM piperazine-N,N'-bis(2-ethanesulfonic acid) (PIPES) (pH 6.5), 20 % glycerol, 100 mM NaCl, 10 mM MgCl₂, and 5 mM BME. Each series of assays was performed 2-3 times using concentrations of [1-³H]FPP (100 mCi/mmol) ranging from 0.025 to 50 μM. The optimal enzyme concentration for each mutant was determined where the dependence of product formation on enzyme concentration was linear and less than 10% of the substrate was turned over: wild-type (1 nM), F96A (20 nM), F198A (20 nM), W203F (20 nM), L72V (2.5 nM), A236G (2.5 nM), and V329L (2.5 nM). A 1-mL reaction mixture in a test-tube was overlaid with 1 mL hexane immediately after addition of substrate, covered with aluminum foil, and incubated for 15 min at 30 °C. The reaction was quenched by addition of 75 μL of 500 mM EDTA (pH 8.0) and vortexed for 20 s. The hexane extract was passed through a silica gel column directly into a scintillation vial containing 5 mL of scintillation fluid. The aqueous phase was extracted with an additional 2 x 1 mL hexane and passed through the same silica gel column. Finally, the column was washed with an additional 1 mL hexane. A Beckman scintillation counter was used to measure product formation, and the substrate concentration versus rate of product formation data was fit by nonlinear regression using the program Prism to determine k_{cat} based on the known total enzyme concentration. For the L72V, A236G and V329A EIZS mutants, the method was adapted to a 0.5 mL reaction volume format, however the volume of hexane used for the extraction was not changed.

7.2.4 GC-MS Analysis of Product Arrays Generated by EIZS Mutants.

The substrate, farnesyl diphosphate (60 μM), was incubated with 40 μM mutant EIZS (F96A, F198A or W203F) in 6 mL buffer (50 mM PIPES, 15 mM MgCl_2 , 100 mM NaCl, 20% glycerol, 5 mM BME) and overlaid with 3 mL HPLC-grade *n*-pentane in a glass test tube at 30 °C for 18 h. Reaction products were extracted with *n*-pentane 3 times, dried with anhydrous MgSO_4 , and concentrated on an ice-water mixture under reduced pressure until the volume was reduced to 100 μL . The products were analyzed using an Agilent 6890 GC/JEOL JMS-600H mass spectrometer, using a 30 m x 0.25 mm HP5MS capillary column (Department of Chemistry, Brown University) in EI (positive) mode using a temperature program of 60-280 °C, with a gradient of 20 °C/min and a solvent delay of 3.5 min. Analysis of the organic extracts resulting from the incubation of FPP with the mutant cyclases by GC-MS reveals the formation of mixtures of sesquiterpene hydrocarbons with $m/z = 204$. Compounds were identified by comparison of their individual mass spectra and chromatographic retention indices with those of authentic compounds in the MassFinder 3.0 Database (Harangi, 2003).

7.3 Results

7.3.1 Radioactive Substrate Kinetic Assay

The steady-state kinetic parameters, K_M and k_{cat} , of the WT and mutant enzymes were measured by a radioactive substrate (^3H -FPP) assay. Scintillation counting was used to monitor the amount of tritium labeled organic products generated by the enzyme during a fixed time. For WT and mutant EIZS, a plot of initial velocity versus substrate concentration (Figures 7.1 and 7.2) was used to determine the steady-state kinetic

parameters using non-linear regression. The resulting kinetic parameters, summarized in Table 7.3, indicate that the mutations only modestly affect K_M , with respect to WT EIZS. This result is expected since the residues required for recognition of the substrate diphosphate (R194, K247, R338, and Y339) are not affected by the mutations. However, the rate of the reaction is affected by some of the mutations. Specifically, the three aromatic mutants (F98A, F198A, and W203F), cause a dramatic decrease in k_{cat} of two orders of magnitude, resulting in overall catalytic efficiencies (k_{cat}/K_M) decreased 205- to 275-fold. The aliphatic mutations (L72V, A236G, and V329A), however, do not result in decreased reaction rates, and thus retain the catalytic efficiency of the WT protein. It is important to note however that this assay simply measures tritium-labeled products extracted in hexanes, therefore the total product yield of the WT or mutant EIZS contains a mixture of sesquiterpene products. Furthermore, the activity of the enzymes may be underestimated, since hydroxylated products may not be extracted with this procedure. This is particularly important in the case of the aromatic mutants, which may be able to accommodate solvent molecules in their active sites (Section 7.4.1), which may increase the proportion of hydroxylated products.

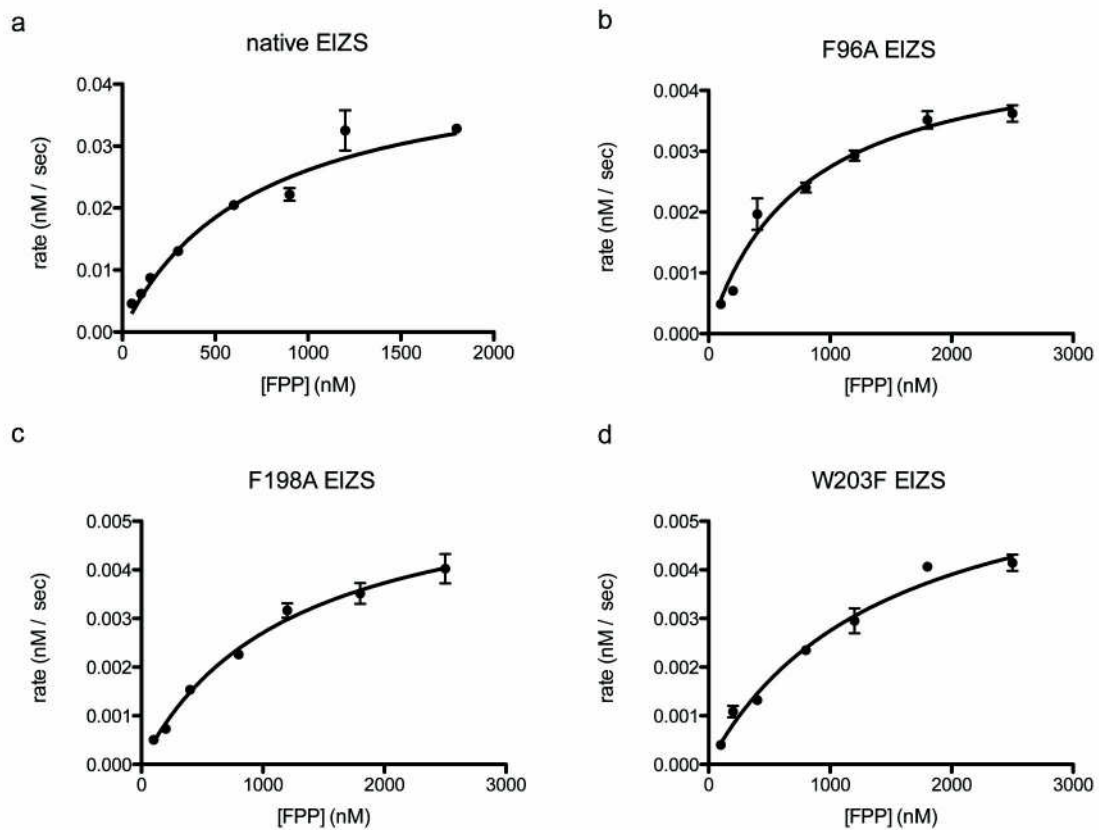


Figure 7.1. Initial rate versus substrate concentration for the reaction of WT and aromatic mutant EIZS with FPP. (a) WT EIZS. (b) F96A EIZS. (c) F198A EIZS. (d) W203F EIZS.

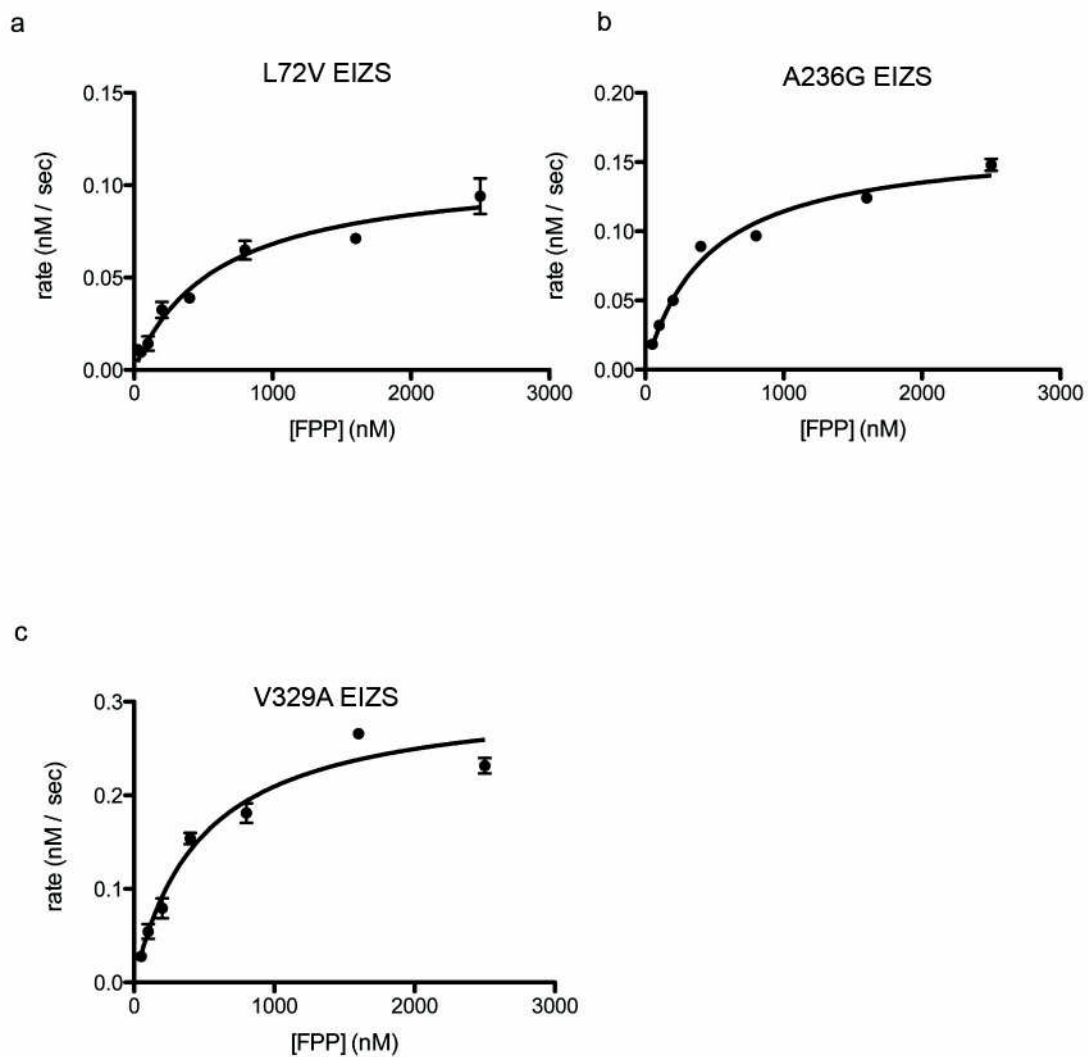


Figure 7.2. Initial rate versus substrate concentration for the reaction of aliphatic mutant EIZS with FPP. (a) L72V EIZS. (b) A236G EIZS. (c) V329A EIZS.

Table 7.3. Steady-state kinetic parameters for wild-type EIZS and site-specific mutants.

Protein	k_{cat} (s^{-1})	K_M (nM)	k_{cat}/K_M ($M^{-1}s^{-1}$)
WT	0.045 ± 0.003	710 ± 100	$6.3 \pm 0.1 \times 10^4$
F96A	0.00024 ± 0.00002	770 ± 130	310 ± 60
F198A	0.00030 ± 0.00002	1200 ± 200	250 ± 45
W203F	0.00034 ± 0.00003	1450 ± 200	250 ± 45
L72V	0.044 ± 0.003	600 ± 120	$7.3 \pm 1.5 \times 10^4$
A236G	0.066 ± 0.002	450 ± 50	$1.5 \pm 0.2 \times 10^5$
V329A	0.126 ± 0.007	470 ± 80	$2.7 \pm 0.5 \times 10^5$

7.3.2 GC-MS Analysis

The sesquiterpene products of the F98A, F198A, and W203F mutants were separated and identified by gas chromatography-mass spectrometry (GC-MS) as GC peaks with corresponding MS parent ions having m/z of 204 ($C_{15}H_{24}$) in the MS. The gas chromatograms of the products of the three aromatic mutants, shown in Figure 7.3, illustrate very unique product arrays for each mutant. The identity of each peak was confirmed by comparison of the observed mass spectra from the MassFinder 3.0 database (Figure 7.4). Several products could not be identified from the available MS data, and are recorded as unknown sesquiterpene products (Figure 7.5). Results are summarized in Table 7.4.

As expected, the product array is severely altered in the aromatic site-specific EIZS mutants. However, the formation of each sesquiterpene product identified can be reasoned to be a result of derailing the proposed EIZS cyclization pathway (Figure 7.6). Notably, none of the three active site aromatic mutants generate *epi*-isozizaene as a major product; indeed, no *epi*-isozizaene whatsoever is generated by the F198A mutant, and the active site contour of this variant as observed in the crystal structure of the F198A EIZS- Mg^{2+}_3 -PP₁-BTAC complex is more complementary in shape to bisabolene-derived cyclization products such as β -acoradiene, as illustrated in Figure 7.7. Notably, β -acoradiene is not generated by wild-type EIZS, so the appearance of this spiroterpenoid represents a new catalytic activity introduced by a single amino acid substitution.

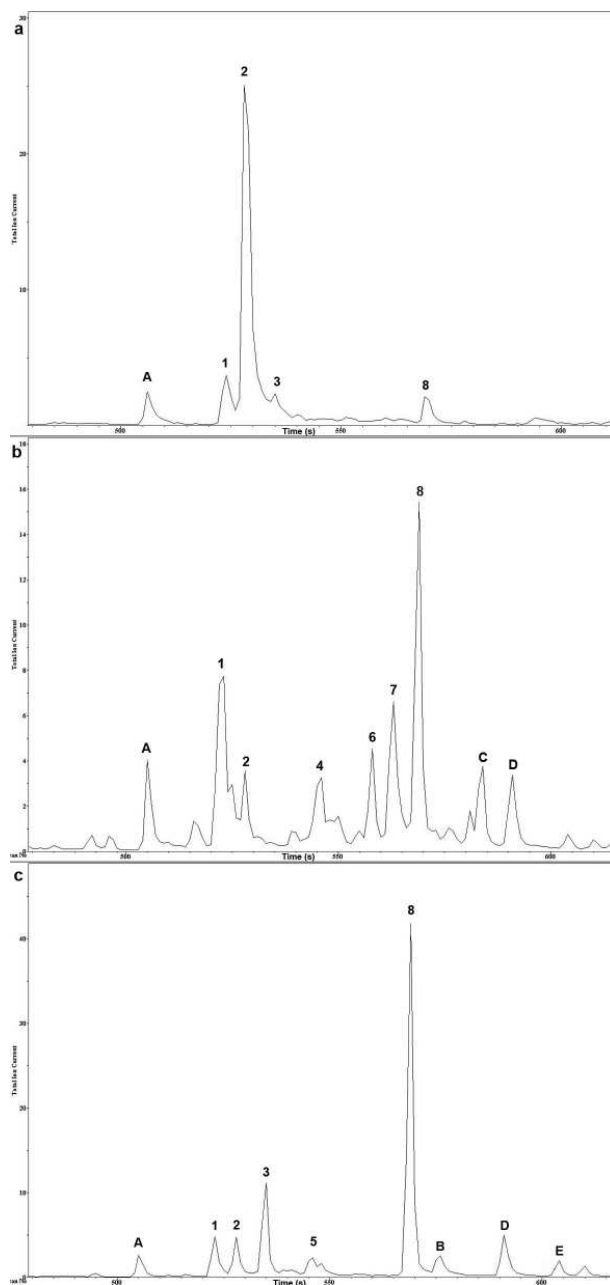
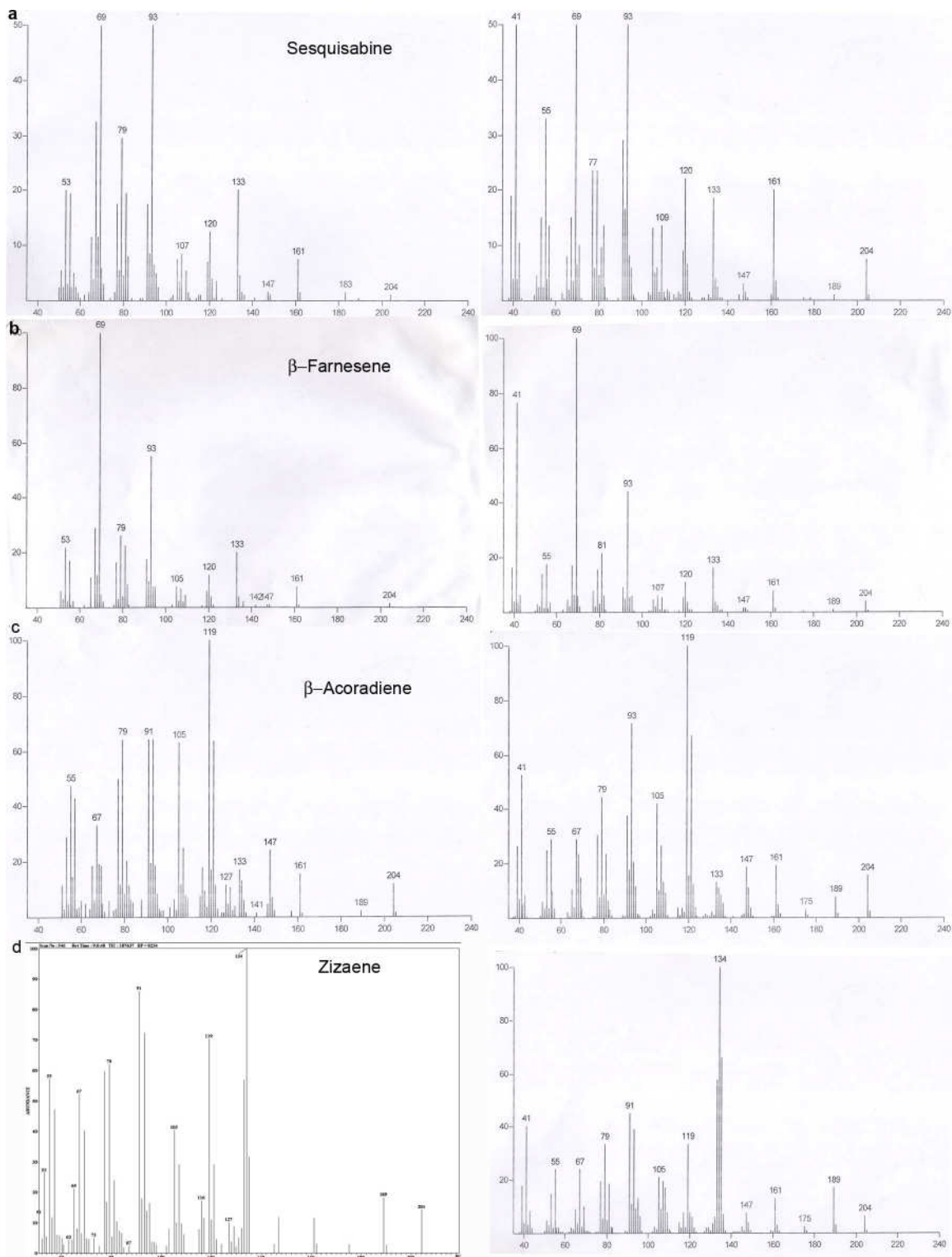
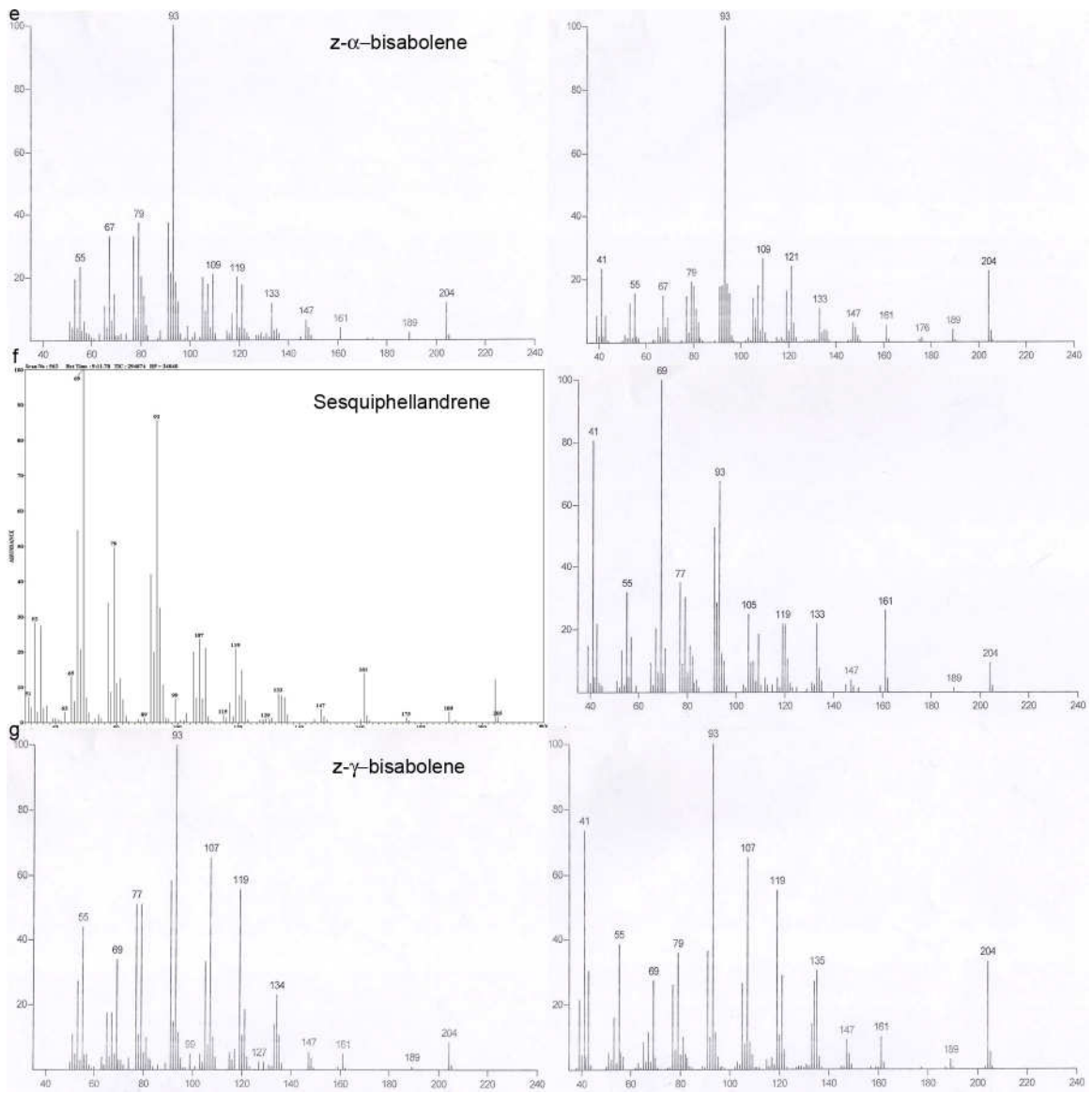


Figure 7.3. Gas chromatographs of hexane extracts of reaction of mutant EIZS with FPP. Unidentified products are labeled as **A-E**, and identified products are labeled as follows: Sesquisabine (**1**), β -farnesene (**2**), *epi*-Isozizaene (**3**), Zizaene (**4**), β -acoradiene (**5**), *Z*- α -bisabolene (**6**), Sesquiphellandrene (**7**), *Z*- γ -bisabolene (**8**). (a) F96A EIZS. (b) F198A EIZS. (c) W203F EIZS.

Figure 7.4 (Following pages). Mass spectra of identified sesquiterpene products of F96A, F198A, and W203F EIZS. For each product the experimental MS is shown on the left and the reference MS from the MassFinder 3.0 database is shown on the right. (a) Sesquisabine. (b) β -farnesene. (c) β -acoradiene. (d) Zizaene. (e) Z- α -bisabolene. (f) Sesquiphellandrene. (g) Z- γ -bisabolene.





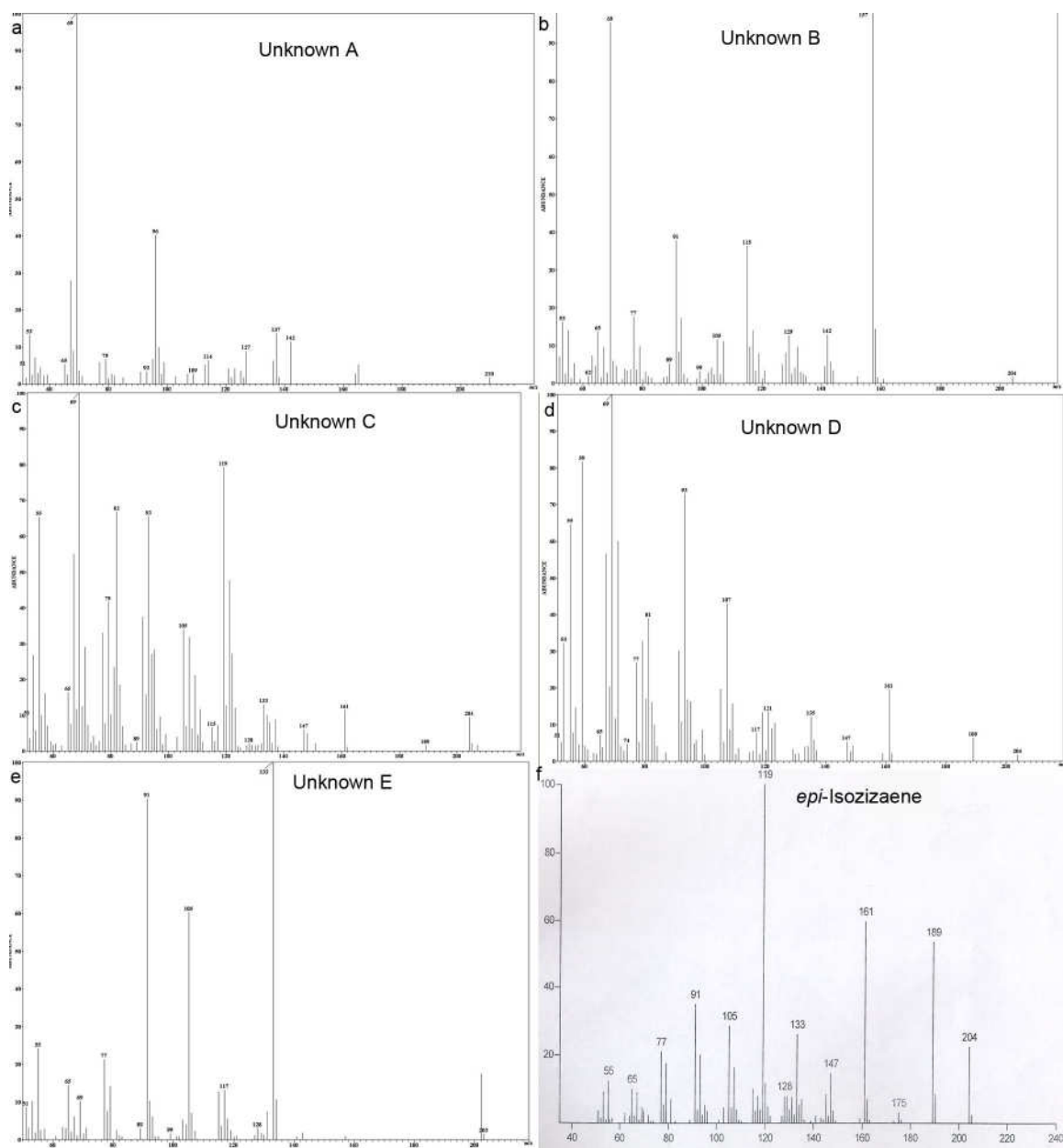


Figure 7.5. Mass spectra of unidentified sesquiterpene products (a-e) of F96A, F198A, and W203F EIZS, and (f) *epi*-Isozizaene.

Table 7.4. Distribution of sesquiterpene products from wild-type EIZS and site specific mutants. Relative product percentages were calculated from the relative peak areas of the sesquiterpene products. The relative product proportions are based on the assumptions that the relative areas of the peaks in the gas chromatogram represent the relative proportion of each sesquiterpene analyte, and the concentration of each sesquiterpene analyte is within the linear range for detection by the instrument.

Protein	Relative product percentage (%)													
	Unknown A	Sesquisabine	β -Farnesene	<i>epi</i> -Isoizaene	Zizaene	β -acoradiene	Z- α -bisabolene	Sesquiphellandrene	Z- γ -bisabolene	Unknown B	Unknown C	Unknown D	Unknown E	α -cedrene
WT	-	2	5	79	9	-	-	1	-	-	-	-	-	2
F96A	7	9	70	8	-	-	-	-	7	-	-	-	-	-
F198A	6	20	5	-	-	12	6	13	24	-	7	7	-	-
W203F	4	7	6	14	7	-	-	-	47	6	-	8	3	-

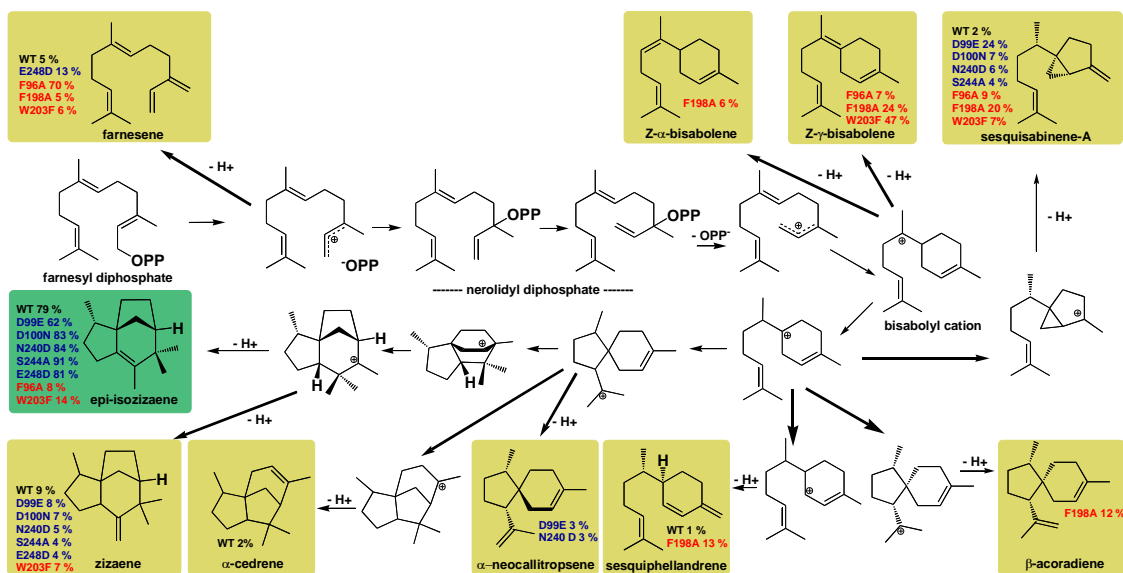


Figure 7.6. Proposed cyclization cascade for observed products of WT and mutant EIZS. Biosynthetic versatility of EIZS can be manipulated by site-directed mutagenesis, as illustrated for sesquiterpene products identified for wild-type (WT) and mutant cyclases. In general, more diverse sesquiterpene product arrays result from the substitution of aromatic residues defining the active site contour (red labels) than from substitution of residues that coordinate the Mg^{2+} ions required for catalysis (blue labels) (Lin, 2009). For example, F198A EIZS does not generate *epi*-isozizaene at all, but instead generates a mixture of sesquisabinene-A, Z- α - and Z- γ -bisabolenes, sesquiphellandrene, and β -acoradiene as its major cyclization products. Remolding the active site contour permits the generation of alternative products as long as they can be accommodated within the remolded template, as illustrated for β -acoradiene in Figure 7.7.

Altering the active site contour of EIZS alters the conformations and cyclization trajectories of FPP and reactive carbocation intermediates, and the formation of alternative products appears to depend on how well a particular alternative product fits the remolded active site contour in a mutant cyclase. Additionally, the F96A and F198A substitutions could compromise the potential stabilization of carbocation intermediates by cation- π interactions. Some of the sesquiterpenes generated by these aromatic mutants have previously been observed as side products generated by both wild-type and mutant EIZS enzymes (Lin, 2009), while three sesquiterpene products have not previously been observed with this cyclase.

(*E*)- β -Farnesene, the major product (70%) formed by F96A EIZS, results from deprotonation at C-3¹ of the allylic cation that results from the initial ionization of FPP; other reaction products include sesquisabinene-A, *Z*- γ -bisabolene, and an unidentified hydrocarbon presumed to be a sesquiterpene based on mass spectrometry, with $m/z = 204$. F198A EIZS generates sesquisabinene-A, (*E*)- β -farnesene, zizaene, β -acoradiene, *Z*- α -bisabolene, sesquiphellandrene, *Z*- γ -bisabolene and 3 unidentified sesquiterpenes. Interestingly, F198A EIZS generates no *epi*-isozizaene. The predominant product of W203F EIZS is *Z*- γ -bisabolene, generated by the abstraction of a proton from the intermediate bisaboly l cation (Figure 7.6). *epi*-Isozizaene accounts for 14% of the products of the W203F mutant; the remaining products include sesquisabinene, (*E*)- β -farnesene, zizaene, and 4 unidentified sesquiterpenes. It is notable that *epi*-isozizaene biosynthetic activity is preserved, if only partially so, in the EIZS mutant with the most

conservative aromatic-aromatic substitution, which presumably preserves more of the general contour and electrostatic profile of the active site.

7.4 Crystal Structures of Mutant EIZS

7.4.1 F198A EIZS-Mg²⁺₃-PP_i-BTAC complex

In order to investigate structural changes in the active site resulting from mutagenesis of aromatic residues, the X-ray crystal structure of F198A EIZS was determined at 1.64 Å resolution. This mutant was selected for X-ray crystallographic study because it retains some catalytic activity (Table 7.3) and exhibits a remarkably altered product array (Table 7.4). Overall, the F198A mutation results in minimal structural perturbations, and the r.m.s. deviation between the structure of wild-type EIZS and F198A EIZS is 0.10 Å for 340 C α atoms. In the active site, the largest structural changes resulting from the F198A substitution are a ~30° rotation of the side chain of F95 and an alternative rotamer of M73. The binding mode of Mg²⁺₃-PP_i is identical to that observed in the wild-type enzyme; however, the BTAC molecule occupies an alternative position such that its benzyl ring makes quadrupole-quadrupole interactions with the aromatic rings of F95, F96, W203, and W325 (Figure 7.8). Surprisingly, 4 solvent molecules are observed in the active site, forming a hydrogen bonding network with N233 and the backbone carbonyl of A236. Comparison of the contours of WT and F198A EIZS helps to explain the production of side products such as β -acoradiene by F198A EIZS, due to the higher complementarity in shape to bisabolene-derived cyclization products (Figure 7.7).

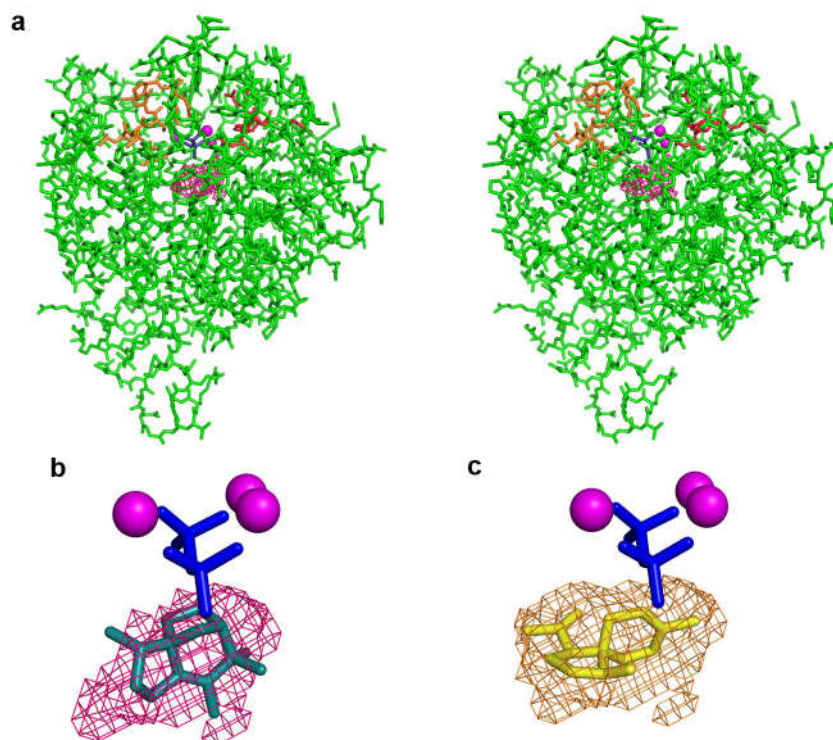


Figure 7.7. (a) A stereoview of the active site surface contour encapsulated by the closed conformation of EIZS is shown as magenta meshwork. The aspartate-rich motif (red) and the NSE motif (orange) are oriented as in Figure 1. (b) The cyclization product, *epi*-isozizaene, is modeled into the enclosed active site contour of EIZS (magenta meshwork), and the location of the $Mg^{2+}_3-PP_i$ cluster is shown as a visual reference. (c) The enclosed active site contour of F198A EIZS (light brown meshwork) into which the new cyclization product β -acoradiene is modeled. The remodeled active site contour in this mutant prevents *epi*-isozizaene formation but permits the formation of new or alternative sesquiterpene products predominantly derived from the bisabolyll carbocation intermediate.

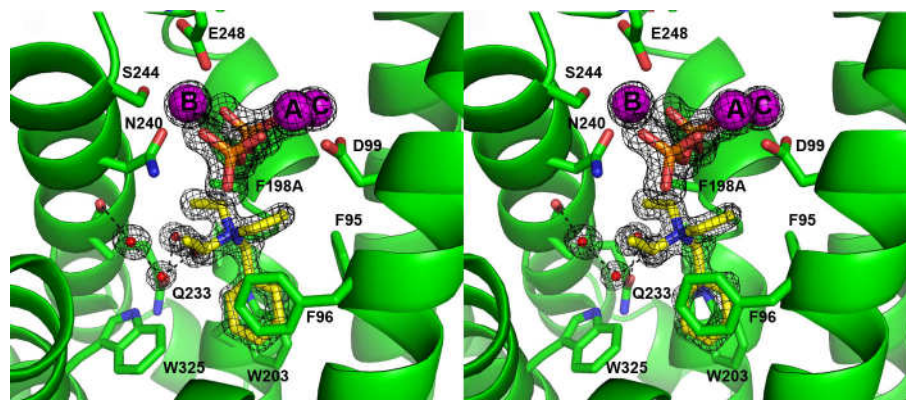


Figure 7.8. Stereoview of the active site of F198A EIZS- Mg^{2+}_3 - PP_i -BTAC complex. Simulated annealing omit maps (black) of the PP_i anion, Mg^{2+} ions, and BTAC in the active site of F198A EIZS, contoured at 5σ . Note the alternative position of BTAC resulting from the F198A mutation.

7.4.2 L72V, A236G and V329A EIZS-Mg²⁺₃-PP_i-BTAC complexes

Crystal structures of the L72V, A236G and V329A EIZS- Mg²⁺₃-PP_i-BTAC complexes were determined at 2.10 Å, 1.76 Å, and 1.95 Å respectively. These mutants were chosen for crystallography because they expressed well and, once purified, were stable and well behaved in solution. Notably, each of these three mutations replaced a larger residue with a smaller residue, resulting in slight changes to the active site cavity contour without altering the polarity of the cavity. All three mutants crystallized under the same conditions as the WT enzyme, including the addition of pyrophosphate and BTAC, resulting in the respective mutant EIZS-Mg²⁺₃-PP_i-BTAC complexes. Overall, each single mutation caused minimal structural perturbations; the rmsd between the structure of WT EIZS and the L72V, A236G and V329A mutants is 0.13 Å, 0.096 Å, and 0.125 Å respectively for 340 C α atoms. The binding mode of the trinuclear Mg²⁺ cluster and pyrophosphate was unchanged in the three mutants, and furthermore the position and orientation of BTAC was also unchanged, except for the L72V mutant in which BTAC moved ~0.5 Å deeper into the active site and rotated by ~20° (Figure 7.9). Minimal changes in the active site are confirmed by active site volume calculations using the program VOIDOO (Kleywegt, 1994). After removing BTAC's coordinates from the active site, the calculated volume of the WT cavity is 62.22 Å³, compared with 62.71 Å³ for L72V, 62.86 Å³ for A236G and 61.71 Å³ for V329A. To determine whether these seemingly trivial volume changes result in altered products arrays, future experiments include GC-MS analysis of the products of these mutants.

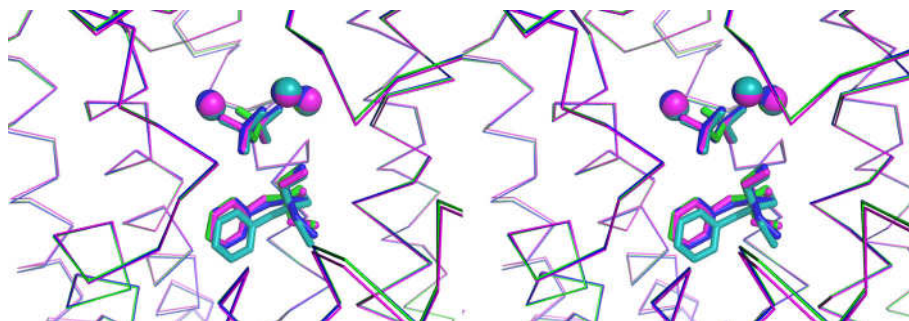


Figure 7.9. Stereoview of an overlay of the active sites of WT (green), L72V (cyan), A236G (blue) and V329L (purple) EIZS-Mg²⁺₃-PP_i-BTAC complexes.

7.5 Discussion

Mutagenesis of aromatic residues that contribute directly to the active site contour significantly compromises the fidelity of *epi*-isozizaene biosynthesis (Figure 7.6), with the F198A substitution completely suppressing *epi*-isozizaene formation and redirecting the cyclization cascade toward the generation of alternative acyclic, monocyclic, and bicyclic sesquiterpenes. Thus, remolding the active site contour by mutagenesis opens up new cyclization trajectories while closing off old ones.

The appearance of low levels of new or alternative cyclization products resulting from mutagenesis of the active site contour in a terpenoid cyclase may reflect past or future evolutionary potential, i.e., catalytic promiscuity in enzyme function may provide a "toehold of evolution" (Petsko, 1993). The evolution of biosynthetic diversity in this family of enzymes is achieved by simply remolding the active site contour to promote one cyclization pathway while suppressing hundreds of others, and it is notable that this is readily achieved by only a handful of amino acid substitutions. The current work represents the first step in deciphering the relationship between the structure of the EIZS active site and its biosynthetic specificity as a product-like template for terpenoid cyclization reactions: the three-dimensional contour of the active site can be remolded to better fit another product and disfavor others, even to the point of excluding *epi*-isozizaene formation. That the biosynthetic specificity of a terpenoid cyclase is so sensitive to and so readily manipulated by minimal mutagenesis in nature or in the laboratory will likely contribute to the growing structural and stereochemical diversity of the terpenome.

Chapter 8: Trinuclear Metal Clusters in Catalysis by Terpenoid Synthases

8.1 Introduction

Terpenoid synthases are ubiquitous enzymes that catalyze the formation of structurally and stereochemically diverse isoprenoid natural products. Many isoprenoid coupling enzymes and terpenoid cyclases from bacteria, fungi, protists, plants, and animals share the class I terpenoid synthase fold. This family of enzymes, which is responsible for such a diverse range of products, is of great interest medicinally since many terpenoid natural products exhibit anti-cancer, anti-malarial, and anti-microbial activities (Aharoni, 2005). Furthermore, in humans, the 15 carbon linear isoprenoid farnesyl diphosphate (FPP) is a precursor in the biosynthesis of steroids and is also utilized for posttranslational prenylation of Ras in GTPase signaling (Rondeau, 2006; Agrawal, 2009). Recently, human farnesyl diphosphate synthase (FPP synthase) has been identified as the target of nitrogen-containing bisphosphonate drugs used for the treatment of bone diseases such as osteoporosis, hypercalcemia, and metastatic bone disease (Ebetino, 2005; Licata, 2005). Moreover, protozoan FPP synthase homologues have recently been identified as targets for the treatment of parasitic infections, including Chagas disease and African sleeping sickness (Ferella, 2008).

Despite generally low amino acid sequence identity class I terpenoid synthases, which adopt the FPP synthase α -helical fold, contain conserved metal binding motifs that coordinate to a trinuclear metal cluster. This cluster not only serves to bind and orient the flexible isoprenoid substrate in the precatalytic Michaelis complex, but it also triggers the

departure of the diphosphate leaving group to generate a carbocation that initiates catalysis. Review of the available crystal structures of class I terpenoid synthases complexed with trinuclear metal clusters and either isoprenoid diphosphates or inorganic pyrophosphate highlights the conserved structural aspects of the trinuclear metal cluster and the additional conserved hydrogen bond donors that are required for catalysis.

8.2 Isoprenoid Coupling Enzymes

8.2.1 Farnesyl diphosphate synthase

Farnesyl diphosphate synthase, the archetypical prenyltransferase, catalyzes the formation of farnesyl diphosphate (FPP), the linear isoprenoid precursor of sesquiterpene natural products. Chain elongation to form FPP proceeds in two distinct steps (Figure 4.1): first, isopentenyl diphosphate (IPP) and dimethylallyl diphosphate (DMAPP) are coupled to form geranyl diphosphate (GPP), and then a second molecule of IPP is coupled to GPP to form FPP. The first crystal structure of FPP synthase was that of the avian enzyme (Tarshis, 1994), which revealed a novel α -helical fold. The structure revealed two conserved aspartate-rich (DDXXD) sequences (Ashby, 1990) on helices D and H, which flank the mouth of the active site cavity. Additionally, a single Sm^{3+} ion, used for heavy metal derivatization for MIR phasing, was bound by each DDXXD motif.

A decade later, the crystal structure of *E. coli* FPP synthase was the first to reveal the binding of a trinuclear magnesium cluster in the active site of an isoprenoid coupling enzyme (Hosfield, 2004), similar to the trinuclear magnesium clusters previously observed in fungal and plant terpenoid cyclases (Rynkiewicz, 2001; Whittington, 2002).

The structure of *E. coli* FPP synthase was solved as the enzyme-substrate ternary complex with the noncleavable DMAPP analogue, dimethylallyl S-thiolodiphosphate (DMSPP), and a molecule of IPP. Applying the Mg^{2+}_A , Mg^{2+}_B , and Mg^{2+}_C nomenclature first established for the trinuclear magnesium cluster of trichodiene synthase (Rynkiewicz, 2001), the crystal structure of the *E. coli* FPP synthase- Mg^{2+}_3 -DMSPP-IPP complex reveals octahedral coordination of all three metal ions (Figure 8.1 (a)): Mg^{2+}_A is coordinated by D105 and D111 of the first aspartate-rich motif on helix D, two diphosphate oxygen atoms, and two water molecules; Mg^{2+}_C is coordinated by the side chains of D105, and D111, as well as one diphosphate oxygen and three water molecules; and Mg^{2+}_B is coordinated by D244 of the second aspartate-rich motif, two diphosphate oxygen atoms, and three water molecules. The diphosphate group of DMSPP also accepts hydrogen bonds from R116, K202, and K258.

More recently, the structure of the human FPP synthase- Mg^{2+}_3 -zoledronate-IPP complex (Rondeau, 2006) reveals complete conservation of Mg^{2+}_3 -diphosphate recognition between *E. coli* and human FPP synthases (Figure 8.1 (b)). In human FPP synthase, two DDXXD motifs coordinate to the Mg^{2+}_3 cluster: the first aspartate of the **D¹⁰³DXXD¹⁰⁷** (hereafter boldface residues indicate Mg^{2+} coordinating residues) motif coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry, and one oxygen atom of D107 bridges Mg^{2+}_A and Mg^{2+}_C with *syn,anti*-coordination stereochemistry; the first aspartate of the second **D²⁴³DXXD** motif coordinates to Mg^{2+}_B . The diphosphate moiety additionally accepts hydrogen bonds from R112, K200, and K257. Interestingly, the closed active site conformation is also stabilized by newly formed hydrogen bonds

between K266 and D107 and D174. The r.m.s. deviation between the unliganded enzyme and the closed conformation of the Mg^{2+}_3 -zoledronate-IPP complex is 1.3 Å (341 C α atoms). Analysis of X-ray crystal structures of several human FPP synthase- Mg^{2+}_3 -bisphosphonate complexes suggests a two-step mechanism for substrate binding (Rondeau, 2006). First, the binding of DMAPP and 3 Mg^{2+} ions brings together the two DDXXD motifs, and loops D-E and H-I come together to form a hydrogen bond between T111 and the backbone of I258. These structural changes close the entrance to the allylic binding site and complete the formation of the IPP binding site. Secondly, as IPP binds, and as the basic C-terminal tail of the enzyme becomes ordered and closes the IPP binding site, IPP and DMAPP are properly oriented for catalysis.

The binding of a trinuclear magnesium cluster is similarly conserved in FPP synthases from parasitic organisms. The flagellated protozoan *T. cruzi* causes Chagas disease, primarily in Latin America (Tanowitz, 2009). Bisphosphonates have emerged as a potential treatment for Chagas disease by inhibiting *T. cruzi* FPP synthase (Garzoni, 2004). The crystal structures of *T. cruzi* FPP synthase- Mg^{2+}_3 -inhibitor complexes (Gabelli, 2006) suggest conservation of the trinuclear magnesium cluster for substrate binding and catalysis. In the complex with risedronate (Figure 8.1 (c)), the first carboxylate of the **D⁹⁸DXXD¹⁰²** motif on helix D coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry, and one oxygen atom of D102 bridges Mg^{2+}_A and Mg^{2+}_C with *syn,anti*-coordination stereochemistry. The first carboxylate of the **D²⁵⁰DXXD** motif on helix H is the only residue that directly coordinates to Mg^{2+}_B ; however, D251 and D254 indirectly interact with Mg^{2+}_B via bridging water molecules. Each Mg^{2+} ion is

coordinated with octahedral geometry, with non-protein coordination sites occupied by oxygen atoms of the inhibitor phosphonate groups and water molecules. Oxygen atoms of the two risedronate phosphonate groups accept hydrogen bonds from the side chains of R107, K207, and K264.

Trypanosoma brucei is an African parasitic protist, and its FPP synthase is related to that of *T. cruzi* by 70 % amino acid sequence identity. The crystal structure of *T. brucei* FPP synthase complexed with 3 Mg^{2+} ions and the bisphosphonate inhibitor BPH-721 (Zhang, 2009) reveals conservation of the trinuclear magnesium cluster for substrate binding and catalysis (Figure 8.1 (d)). The first carboxylate of the **D¹⁰³DXXD¹⁰⁷** motif on helix D coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry, and one oxygen atom of D107 bridges Mg^{2+}_A and Mg^{2+}_C with *syn,anti*-coordination stereochemistry. The first aspartate in the **D²⁵⁵DXXD** motif on helix H is the only residue that directly coordinates to Mg^{2+}_B ; however, D256 and D259 indirectly interact with Mg^{2+}_B via bridging water molecules. Oxygen atoms of the two phosphonate groups of the inhibitor BPH-721 also accept hydrogen bonds from the side chains of R112, K212, and K269 (Mao, 2006).

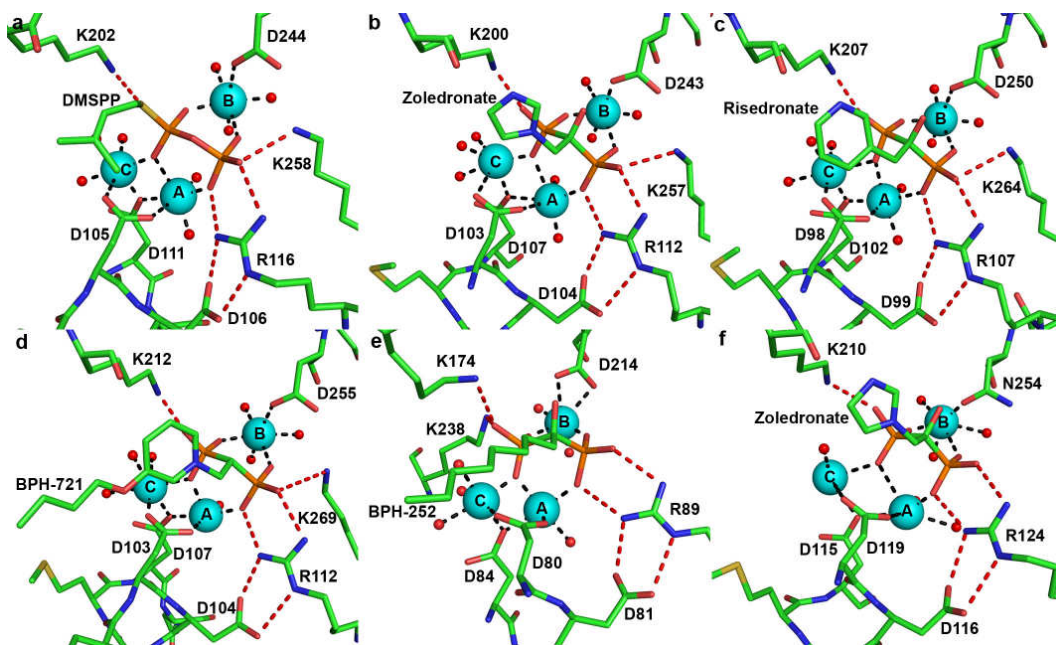


Figure 8.1. Conservation of Mg^{2+} - PP_1 and -diphosphate binding motifs among isoprenoid coupling enzymes. Metal coordination (black) and hydrogen bond (red) interactions with phosphate(s) are indicated. (a) *E. coli* FPP synthase- Mg^{2+} -DMSPP-IPP complex (PDB code 2EGW); (b) human FPP synthase- Mg^{2+} -zoledronate complex (PDB code 2F8Z); (c) *T. cruzi* FPP synthase- Mg^{2+} -risedronate complex (PDB code 1YHL); (d) *T. brucei* FPP synthase- Mg^{2+} -BPH-721 complex (PDB code 3DYH); (e) *S. cerevisiae* GGPP synthase- Mg^{2+} -BPH-252 complex (PDB code 2ZAX); (f) *C. parvum* nonspecific prenyl synthase- Mg^{2+} -zoledronate complex (PDB code 2Q58).

8.2.2 Geranylgeranyl diphosphate synthase

Geranylgeranyl diphosphate synthase (GGPP synthase) catalyses the condensation of IPP and FPP to form GGPP (Figure 4.1). Recently, GGPP synthase has emerged as a pharmaceutical target for the treatment of cancer since geranylgeranylation is involved in Rac, Rap and Rho signaling pathways (Russell, 2006). The crystal structures of GGPP synthases from *Thermus thermophilus* (Nishio, 2004), *Sinapis alba* (Kloer, 2006), and *Saccharomyces cerevisiae* (Chang, 2006) have been determined in addition to that of human GGPP synthase (Kavanagh, 2006). However, a crystal structure containing a complete trinuclear magnesium cluster has only been observed in the active site of monomer B of the *S. cerevisiae* GGPP synthase-Mg²⁺₃-BPH-252 complex (Figure 8.1 (e)) (Chen, 2008). The first aspartate of the **D⁸⁰DIED⁸⁴** motif coordinates to Mg²⁺_A and Mg²⁺_C with *syn,syn*-bidentate geometry, and one oxygen atom of D84 bridges Mg²⁺_A and Mg²⁺_C with *syn,anti*-coordination stereochemistry; the first aspartate of the second **D²¹⁴DYLN** motif coordinates to Mg²⁺_B. Each Mg²⁺ ion is coordinated with octahedral geometry, with non-protein coordination sites occupied by oxygen atoms of the inhibitor phosphonate groups and water molecules. Oxygen atoms of the two phosphonate groups of the inhibitor BPH-252 also accept hydrogen bonds from R89, K174 and K238.

8.2.3 Nonspecific prenyl synthase

Recently, the crystal structure of a nonspecific prenyl synthase from *Cryptosporidium parvum* has been determined (Artz, 2008). *C. parvum* causes livestock infections and is classified as a bioterrorism threat by the Centers for Disease Control and Prevention (Hashsham, 2004). The enzyme has a unique ability to catalyze chain

elongation reactions with isoprenoid substrates of various lengths to generate C₂₀-C₄₅ linear isoprenoids products. The crystal structure of the enzyme reveals conservation of the classic α -helical terpenoid synthase fold, and its complex with the inhibitor risedronate reveals that a complete trinuclear magnesium cluster is coordinated by DDXXD and NDXXD motifs (Figure 8.1 (f)) (Artz, 2008). The first carboxylate of the D¹¹⁵DXXD¹¹⁹ motif on helix D is oriented for coordination to Mg²⁺_A and Mg²⁺_C with *syn,syn*-bidentate geometry; however, the distance between D115 and Mg²⁺_C is 3.14 Å, thus too long to be considered an inner-sphere metal coordination interaction. One oxygen atom of the third aspartate, D119, bridges Mg²⁺_A and Mg²⁺_C with *syn,anti*-coordination stereochemistry. N254 of the N²⁵⁴DXXD motif on helix H coordinates to Mg²⁺_B, and D255 and D258 indirectly interact with Mg²⁺_B via bridging waters. The diphosphate moiety additionally accepts hydrogen bonds from K210 and G251, and the closed active site conformation is stabilized by hydrogen bonds between D116 of the DDXXD motif and R124, and D255 of the NDXXD motif and Q251.

8.3 Isoprenoid Cyclization Enzymes

8.3.1 Fungal cyclases

The sesquiterpene cyclase trichodiene synthase from *Fusarium sporotrichioides* catalyzes the first committed step in the biosynthesis of nearly 100 different trichothecene mycotoxins. Trichodiene synthase is one of the most thoroughly studied terpenoid cyclases, and enzymological and crystallographic studies have illuminated important features in the cyclization mechanism (recently reviewed in (Christianson, 2006)). Recent computational studies have also provided new insight on the catalytic mechanism (Hong,

2009). The structures of unliganded trichodiene synthase and the trichodiene synthase- $\text{Mg}^{2+}_3\text{-PP}_i$ complex were the first to reveal the binding of a trinuclear magnesium cluster in the active site of a terpenoid synthase (Figure 8.2 (a)) (Rynkiewicz, 2001). The first aspartate of the $\mathbf{D}^{100}\text{DXXD}$ motif on helix D coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry. The second metal binding motif $\mathbf{N}^{225}\text{DLMS}^{229}\text{FYKE}^{333}$ is located on helix H and coordinates to Mg^{2+}_B . All three metal ions are additionally coordinated by PP_i and solvent molecules to complete octahedral coordination polyhedra.

Superposition of the unliganded and $\text{Mg}^{2+}_3\text{-PP}_i$ complexed trichodiene synthase structures reveals conformational changes that cap the active site upon ligand binding. Overall, the r.m.s. deviation between the native and liganded structures is 1.4 Å for 349 C α atoms. Interestingly, upon ligand binding, D101 in the aspartate-rich motif forms a salt bridge with R304, which donates a hydrogen bond to PP_i . In addition to Mg^{2+} coordination interactions, the PP_i anion also accepts hydrogen bonds from R182, K232, and Y305. The D101-R304- PP_i hydrogen bond network appears to link substrate binding with the transition between the open and closed active site conformations. Assuming that the diphosphate group of FPP triggers the same structural changes as observed for PP_i , the substrate is sequestered from bulk solvent and the complete trinuclear magnesium cluster triggers departure of the diphosphate leaving group to generate the carbocation that initiates the cyclization cascade. The seemingly conservative D100E mutation results in a 22-fold loss in catalytic activity (measured by k_{cat}/K_M) and structural studies indicate that the additional methylene group of E100 perturbs the $\text{Mg}^{2+}_3\text{-PP}_i$ complex such that Mg^{2+}_A binding is weakened, E233 breaks its coordination interaction with Mg^{2+}_B , and

Mg^{2+} is dissociated; additionally, hydrogen bond interactions between PP_i and R182 and R304 are broken (Rynkiewicz, 2002; Vedula, 2005b).

The role of the D101-R304 salt bridge in closing the trichodiene synthase active site has been explored in mutagenesis studies. The D101E mutation results in a moderate 5-fold decrease in catalytic activity; however, there are no crystal structures of this mutant available for study (Cane, 1996). In contrast, the R304K mutant results in a 5000 fold decrease in catalytic activity, and the crystal structure of R304K trichodiene synthase complexed with $Mg^{2+}_3-PP_i-(R)$ -azabisabolene reveals the loss of the expected hydrogen bond between K304 and D101 (Vedula, 2005a). Although the PP_i binding motif remains intact in the R304K mutant, it is evident that the R304-D101 hydrogen bond is critical for properly activating the substrate diphosphate group. In contrast, while Y305 donates a hydrogen bond to PP_i in the wild-type enzyme complex with $Mg^{2+}_3-PP_i$, catalytic activity and PP_i binding are not significantly affected in the Y305F mutant (Cane, 1995; Vedula, 2005b).

Another fungal cyclase that has been the subject of extensive structural and functional study is aristolochene synthase, which is a sesquiterpene cyclase that catalyzes the cyclization of FPP to form (+)-aristolochene. Structures of aristolochene synthases from *Penicillium roqueforti* (Caruthers, 2000) and *Aspergillus terreus* (Shishova, 2007) have been solved; these enzymes are related by 61 % amino acid sequence identity. Although there is no crystal structure of *P. roqueforti* aristolochene synthase complexed with $Mg^{2+}_3-PP_i$, the structure of the *A. terreus* aristolochene synthase $Mg^{2+}_3-PP_i$ complex

(Shishova, 2007) (Figure 8.2 (b)) indicates conservation of the Mg^{2+}_3 -PP_i binding motif first observed in trichodiene synthase (Rynkiewicz, 2001).

The aspartate-rich motif of aristolochene synthase on helix D appears as **D⁹⁰DXXE**. The carboxylate side chain of D90 coordinates to Mg^{2+}_A and Mg^{2+}_B with *syn,syn*-bidentate geometry, and is the only residue in the aspartate motif that coordinates to the Mg^{2+} ions. The carboxylate group of D91 makes a salt bridge with R304, and the final carboxylate in the motif is E119, which accepts a hydrogen bond from a water molecule coordinated to Mg^{2+}_A . The second metal binding motif **N²¹⁹DIYS²²³YEKE²²⁷** is located on helix H and chelates Mg^{2+}_B (Shishova, 2007), consistent with the structures of terpenoid cyclases from plants, bacteria, and fungi (Starks, 1997; Rynkiewicz, 2001; Whittington, 2002; Hyatt, 2007; Aaron, 2010). As found in the active sites of trichodiene synthase (Rynkiewicz, 2001) and *epi*-isozizaene synthase (Aaron, 2010), PP_i binding in aristolochene synthase is similarly accommodated by hydrogen bonds donated from two arginines (R175 and F314), one lysine (K226), and one tyrosine (Y315) (Figure 8.2 (b)).

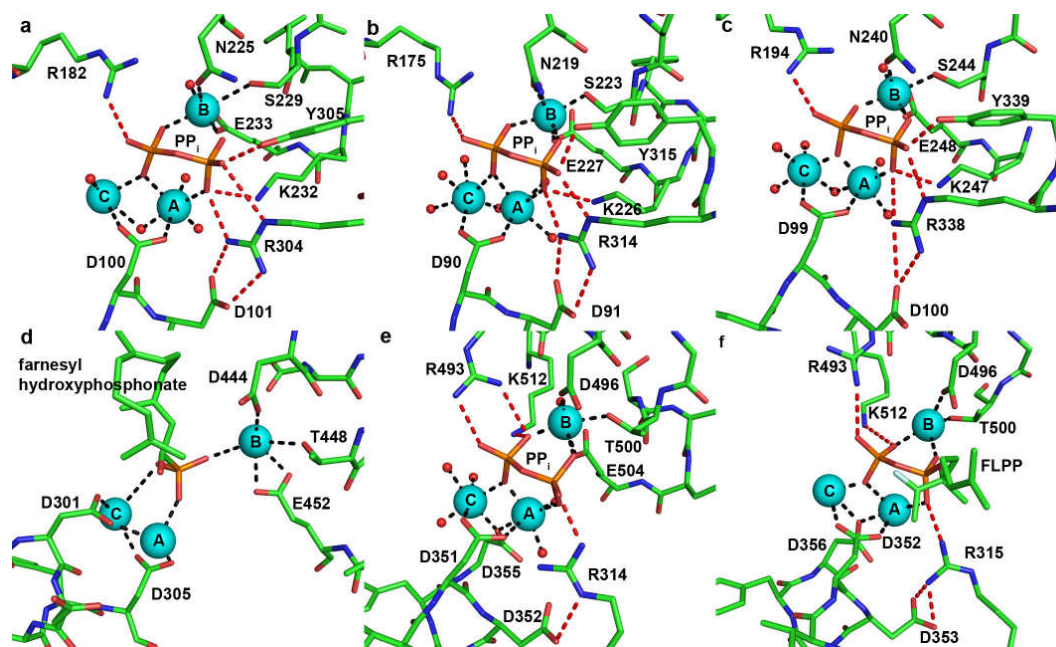


Figure 8.2. Conservation of Mg^{2+} - PP_i and -diphosphate binding motifs among terpenoid cyclases. Metal coordination (black) and hydrogen bond (red) interactions with phosphate(s) are indicated. (a) *F. sporotrichioides* trichodiene synthase- Mg^{2+} - PP_i complex (PDB code 1JFG); (b) *A. terreus* aristolochene synthase- Mg^{2+} - PP_i complex (PDB code 2OA6); (c) *S. coelicolor* *epi*-isozizaene synthase- Mg^{2+} - PP_i complex (PDB code 3KB9); (d) *N. tabacum* 5-*epi*-aristolochene synthase- Mg^{2+} -farnesyl hydroxyphosphonate complex (PDB code 5EAT; note that many of the metal-phosphate interactions indicated are too long to be considered inner-sphere metal coordination interactions); (e) *S. officinalis* (+)-bornyl diphosphate synthase- Mg^{2+} - PP_i complex (PDB code 1N22; metal ions are labeled according to the convention first established for trichodiene synthase); (f) *M. spicata* limonene synthase- Mn^{2+} -FLPP complex (PDB code 2ONG; conserved hydrogen bonding is indicated between D353 and R315 despite poor geometry).

8.3.2 Bacterial cyclases

In recent years, prokaryotes have emerged as sources of diverse isoprenoids. Specifically, a large number of novel isoprenoids have been isolated from organisms belonging to the taxonomical order *Actinomycetales* (Daum, 2009). The crystal structures of two bacterial sesquiterpene cyclases have been solved, and both derive from *Actinomycetales*: pentalenene synthase from *Streptomyces* UC5319 (Lesburg, 1997) and *epi*-isozizaene synthase from *Streptomyces coelicolor* (Aaron, 2010). Pentalenene synthase catalyzes the cyclization of FPP to form the tricyclic sesquiterpene pentalenene in the first committed step in the biosynthesis of the pentalenolactone family of antibiotics (Seto, 1980). Although the structure of pentalenene synthase was the first to be reported of a terpenoid cyclase and demonstrated that the terpenoid cyclase shared the FPP synthase fold first observed for avian FPP synthase (Tarshis, 1994), no structure of this bacterial terpenoid cyclase complexed with metal ions is available.

However, the crystal structure of *epi*-isozizaene synthase complexed with Mg^{2+} - PP_i -BTAC (BTAC is the benzyltriethylammonium cation, a crystallization additive) reveals that Mg^{2+} - PP_i binding motifs are conserved between fungal and bacterial terpenoid cyclases (Figure 8.2 (c)) (Aaron, 2010). The first aspartate of the aspartate-rich motif **D⁹⁹**DRHD coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry, and Mg^{2+}_B is chelated by **N²⁴⁰**DLCS²⁴⁴LPKE²⁴⁸. Each Mg^{2+} ion is coordinated with octahedral geometry and nonprotein coordination sites are occupied by oxygen atoms of PP_i and water molecules. The PP_i anion also accepts hydrogen bonds from the side chains of R194, K247, R338, and Y339 which correspond to R182, K232, R304 and Y305 of

trichodiene synthase (Rynkiewicz, 2001) and R175, K226, R314 and Y315 of aristolochene synthase (Shishova, 2007). Pentalenene synthase and *epi*-isozizaene synthase share 24 % amino acid sequence identity, and residues that interact with Mg^{2+} ions or PP_i in *epi*-isozizaene synthase are conserved in pentalenene synthase. Superposition of the liganded closed structure of *epi*-isozizaene synthase with unliganded pentalenene synthase reveals a very similar alignment of the metal binding motifs, with pentalenene synthase helices D and H 1.5 Å further apart than in *epi*-isozizaene synthase. Accordingly, upon substrate or PP_i binding the active site of pentalenene synthase presumably undergoes a conformational change to a closed conformation comparable to that observed for the *epi*-isozizaene synthase Mg^{2+}_3 - PP_i -BTAC complex.

Although the second aspartate of the *epi*-isozizaene synthase aspartate-rich motif, D100, does not directly interact with the Mg^{2+} ions or PP_i , it does accept a hydrogen bond from R338, which also donates a hydrogen bond to PP_i . Site-directed mutagenesis reveals that the D100N mutation causes a >95 % loss of activity with respect to the native enzyme (Lin, 2009), suggesting that the D100N mutation disrupts the D100-R338- PP_i hydrogen bond network presumed to be important for substrate recognition. As previously discussed, the second aspartate in the aspartate-rich motifs of trichodiene synthase and aristolochene synthase similarly stabilizes a hydrogen bond network with R304 and PP_i (Rynkiewicz, 2001; Shishova, 2007), so it appears that the bacterial and fungal cyclases share the same molecular strategy for linking the molecular recognition of the substrate diphosphate group with the active site closure mechanism (Aaron, 2010). The third aspartate in the aspartate rich motif of *epi*-isozizaene synthase, D103, points

away from the active site and makes no hydrogen bond interactions that are involved in substrate binding, as also observed in $\text{Mg}^{2+}_3\text{-PP}_i$ complexes of trichodiene synthase. The absence of a structural or catalytic role for the terminal aspartate in the aspartate-rich motif of bacterial terpenoid cyclases is supported by mutagenesis of the corresponding residue in pentalenene synthase: the D84E mutation results in a mere 3-fold loss of catalytic activity (as measured by k_{cat}/K_M), whereas the D80E and D81E mutations yield 3500- and 400-fold reductions in activity, respectively (Seemann, 2002).

8.3.3 Plant Cyclases

5-epi-Aristolochene synthase from *Nicotiana tabacum* catalyzes the cyclization of FPP to form *5-epi-aristolochene* in the first committed step in the biosynthesis of the antifungal phytoalexin capsidiol (Starks, 1997). As the first crystal structure determined of a plant terpenoid cyclase and the second terpenoid cyclase structure to be reported, the structure of *5-epi-aristolochene synthase* reveals the presence of 2 domains (Starks, 1997): a catalytically active C-terminal domain that adopts the α -helical class I terpenoid synthase fold, and an N-terminal domain of unknown function that exhibits an α -helical fold similar to that of a class II terpenoid synthase (Wendt, 1998). Two metal-binding motifs are identified: an aspartate-rich motif $\mathbf{D}^{301}\mathbf{DXXD}^{305}$, and a $\mathbf{D}^{444}\mathbf{DTAT}^{448}\mathbf{YEVE}^{452}$ motif. While the binding of a trinuclear magnesium cluster was identified in the *5-epi-aristolochene synthase* farnesyl hydroxyphosphonate complex (Figure 8.2 (d)), analysis of the structure reveals that many of the coordination interactions with Mg^{2+} ions range 2.2 Å – 3.7 Å, longer than expected for ideal Mg^{2+} coordination (Zheng, 2008). This could suggest that the structure is that of a partially closed conformation. Nonetheless,

D301 and D305 of the aspartate-rich motif coordinate to Mg^{2+}_A and Mg^{2+}_C , while the “DTE” motif chelates Mg^{2+}_B .

Interestingly, metal binding motifs are shared between sesquiterpene cyclases and monoterpene cyclases from plants. The monoterpene cyclase (+)-bornyl diphosphate synthase catalyzes the cyclization of geranyl diphosphate (GPP) to form (+)-bornyl diphosphate. This cyclization is unusual in that the substrate diphosphate group is reincorporated into the product. The structure of (+)-bornyl diphosphate synthase from *Salvia officinalis* was the first of a monoterpene cyclase (Whittington, 2002), and remains the only monoterpene cyclase for which structures have been solved in unliganded and liganded states. The crystal structure of (+)-bornyl diphosphate synthase reveals the two-domain α -helical architecture first observed for the plant sesquiterpene synthase 5-*epi*-aristolochene synthase: a catalytically active C-terminal domain adopting the class I terpenoid synthase fold, and an N-terminal domain adopting the class II terpenoid synthase fold (however, the N-terminal polypeptide caps the active site of the C-terminal domain in ligand complexes) (Whittington, 2002). The (+)-bornyl diphosphate synthase- $\text{Mg}^{2+}_3\text{-PP}_i$ complex reveals that conserved metal-binding motifs and the PP_i anion (or the diphosphate group of the product itself, (+)-bornyl diphosphate) coordinate to 3 Mg^{2+} ions (Figure 8.2 (e)). The first carboxylate of the **D³⁵¹DXXD³⁵⁵** motif coordinates to Mg^{2+}_A and Mg^{2+}_C with *syn,syn*-bidentate geometry, and D355 bridges Mg^{2+}_A and Mg^{2+}_C with *syn,anti*-coordination stereochemistry. Interestingly, unlike metal binding in the active sites of bacterial and fungal cyclases, both the first *and* third aspartates in the DDXXD motif of plant terpenoid cyclases coordinate to the catalytic metal ions. The

second metal binding motif, **D⁴⁹⁶DKGT⁵⁰⁰SYFE⁵⁰⁴**, chelates Mg²⁺_B (Whittington, 2002). In addition to metal ion coordination interactions, the PP_i anion accepts hydrogen bonds from R314, R493, and K512. Comparison of the structures of unliganded and Mg²⁺₃-PP_i complexed (+)-bornyl diphosphate synthase reveals several Mg²⁺₃-PP_i induced conformational changes; however, the r.m.s. deviation of 306 C α atoms in the catalytic C-terminal domain is only 0.6 Å (Whittington, 2002), significantly lower than observed for ligand-induced conformational changes in trichodiene synthase (1.4 Å) (Rynkiewicz, 2001) and aristolochene synthase (1.8 Å) (Shishova, 2007).

The recent structure determination of another plant monoterpene cyclase, limonene synthase from *Mentha spicata* (Hyatt, 2007), similarly reveals conservation of a trinuclear metal cluster in a cyclization reaction that generates 94% (-)-(4S)-limonene, and ~2% myrcene, (-)- α -pinene, and (-)- β -pinene (Williams, 1998). Limonene synthase shares the 2-domain α -helical fold common to plant terpenoid cyclases. Limonene synthase displays similar activity with Mg²⁺ or Mn²⁺ (a common feature of some terpenoid cyclases), and the structure of the enzyme has been determined in complex with 3 Mn²⁺ ions and the intermediate analogue 2-fluorolinalyl diphosphate (FLPP) (Figure 8.2 (f)) (Hyatt, 2007). Metal coordination interactions are similar to those observed in (+)-bornyl diphosphate synthase (Whittington, 2002). In limonene synthase, the first carboxylate of the **D³⁵²DXXD³⁵⁶** motif coordinates to Mn²⁺_A and Mn²⁺_C with *syn,syn*-bidentate geometry, and one oxygen atom of D356 bridges Mn²⁺_A and Mn²⁺_C with *syn,anti*-coordination stereochemistry. Two out of three residues in the second metal binding motif, **D⁴⁹⁶DLGT⁵⁰⁰SVEE⁵⁰⁴**, chelate Mg²⁺_B; the position of the side chain of

E504 is not indicated and is presumably disordered. Additionally, the γ -hydroxyl of T500 is 3.2 Å away from $\text{Mg}^{2+}_{\text{B}}$, which is too long to be considered an inner sphere coordination interaction. The diphosphate group of the bound intermediate analogue FLPP accepts hydrogen bonds from R315, R493, and K512 (Hyatt, 2007).

Finally, the sesquiterpene cyclase (+)- δ -cadinene synthase from *Gossypium arboreum* (tree cotton) catalyzes the first committed step in the biosynthesis of the triterpene phytoalexin gossypol, a major defense metabolite synthesized by cotton plants (Chen, 1995). The recently determined structure of the unliganded enzyme and its complex with 2-fluorofarnesyl diphosphate (2F-FPP) reveals that minimal structural deviations result from ligand binding (the r.m.s. deviations are 0.28 Å and 0.50 Å between unliganded and liganded monomers A (514 C α atoms) and B (494 C α atoms), respectively) (Gennadios, 2009). In contrast with the plant terpenoid cyclases previously discussed (Starks, 1997; Whittington, 2002; Hyatt, 2007), (+)- δ -cadinene synthase contains a second aspartate-rich motif in place of the DTE motif on helix H. As previously discussed, this motif on helix H is common to chain elongation enzymes such as farnesyl diphosphate synthase, and (+)- δ -cadinene synthase is unique among known class I terpenoid cyclases in that it contains two aspartate-rich motifs for metal coordination. The structure of the liganded enzyme reveals a putative Mg^{2+}_3 cluster (weak electron density characterizes the three Mg^{2+} ions); $\text{Mg}^{2+}_{\text{A}}$ and $\text{Mg}^{2+}_{\text{C}}$ are coordinated by D307 and D311 of the first **D³⁰⁷DXXD³¹¹** motif, and $\text{Mg}^{2+}_{\text{B}}$ is coordinated by D451 and E455 of the second aspartate-rich motif, **D⁴⁵¹DVAE⁴⁵⁵** (Figure 8.3). However, many of the carboxylate- Mg^{2+} distances observed are too long for inner sphere

metal coordination interactions; therefore, the structure may reflect an incomplete transition between the “open” and “closed” active site conformations. The diphosphate moiety of 2F-FPP accepts one hydrogen bond from a nearby basic residue, R448.

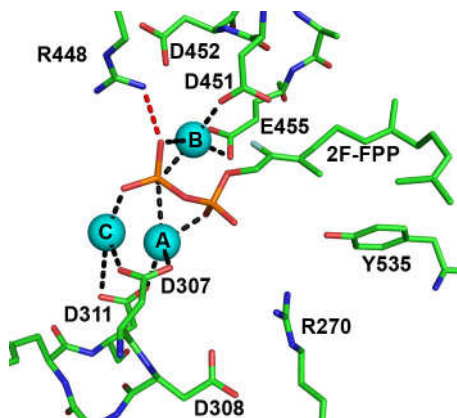


Figure 8.3. The diphosphate binding site of (+)- δ -cadinene synthase from *G. arboreum* (PDB code 3G4F) with a putative Mg^{2+}_3 cluster and 2F-FPP bound. Metal ions are labeled according to convention established for trichodiene synthase. Some metal-phosphate interactions are too long to be considered inner-sphere metal coordination interactions, which could be a consequence of the nonproductive binding mode observed for 2F-FPP.

8.4 Discussion

Although the metal-dependence of catalysis by class I terpenoid synthases has been known for decades (Robinson, 1970), it was not until 2001 that the crystal structure of a terpenoid cyclase-Mg²⁺₃-PP_i complex (trichodiene synthase) revealed that a trinuclear metal cluster accommodates PP_i binding; this trinuclear metal cluster is similarly implicated in binding and activating substrate farnesyl diphosphate for catalysis (Rynkiewicz, 2001). Since then, many X-ray crystal structures of isoprenoid coupling enzymes and terpenoid cyclases have been determined containing Mg²⁺₃ (or Mn²⁺₃) clusters. Comparisons of these structures reveal significant conservation in the constellation of metal ions and the residues that coordinate to these metal ions (Figures 8.1 and 8.2) despite generally insignificant amino acid sequence identity among these enzymes.

Trinuclear metal cluster coordination in FPP synthases is conserved among humans, bacteria and protozoans. Two aspartate-rich DDXXD binding motifs coordinate to 3 Mg²⁺ ions, which are also coordinated by the substrate diphosphate group. The first and last aspartate in the first DDXXD motif coordinate to Mg²⁺_A and Mg²⁺_C, and the first aspartate of the second DDXXD motif coordinates to Mg²⁺_B. Also conserved are one arginine and two lysine residues that donate hydrogen bonds to diphosphate oxygens; the conserved arginine residue also donates hydrogen bond(s) to the second aspartate in the first DDXXD motif (Figure 8.1 (a)-(d)). The crystal structures of other isoprenoid coupling enzymes, GGPP synthase and nonspecific prenyl synthase, similarly reveal conservation of Mg²⁺₃ binding motifs. Hydrogen bond interactions with PP_i are also

conserved (Figure 8.1 (e)-(f)).

It is notable that the constellation of three metal ions and hydrogen bond donors is also conserved, with minor variations, in terpenoid cyclases from plants, bacteria, and fungi (Figure 8.2). First, Mg^{2+}_A and Mg^{2+}_C are coordinated by the first DDXXD motif: bacterial and fungal cyclases utilize only the first aspartate of this motif, whereas plant cyclases utilize the first and third aspartates of this motif (analogous to isoprenoid coupling enzymes). Second, the second aspartate-rich motif is usually replaced by an **NDXXSXXXE** motif in bacterial and fungal terpenoid cyclases and a **DXXTXXXE** motif in plant terpenoid cyclases, in which boldface residues chelate Mg^{2+}_B (although there can be some variations in this sequence, e.g., see (Zhou, 2009)). One exception, however, is (+)- δ -cadinene synthase, in which two aspartate-rich motifs coordinate to the trinuclear metal cluster. Third, residues that donate hydrogen bonds to PP_i are conserved in terpenoid cyclases across different domains of life. Specifically, two arginines donate hydrogen bonds to diphosphate oxygens: one appears to replace a conserved lysine serving this function in the isoprenoid coupling enzymes, and the other also donates a hydrogen bond to the second aspartate of the first DDXXD motif (as observed in the isoprenoid coupling enzymes). In bacterial and fungal terpenoid cyclases, conserved lysine and tyrosine residues additionally donate hydrogen bonds to PP_i .

In all cases in which a complete Mg^{2+}_3 - PP_i cluster is bound, two 6-membered ring chelates are formed with Mg^{2+}_A and Mg^{2+}_B (Figure 8.4). The conformations of these 6-membered rings can vary, e.g., sofa, half-chair, etc. Such 6-membered ring chelates are

occasionally observed in metal-diphosphate binding interactions, e.g., in the binding of the substrate analogue imidodiphosphate to inorganic pyrophosphatase (Fabrichniy, 2007).

In summary, conservation of a trinuclear metal cluster is critical for catalysis by class I terpenoid synthases. This cluster not only serves to bind and orient the flexible isoprenoid substrate in the precatalytic Michaelis complex, but it also triggers leaving group departure and initial carbocation formation. Conserved hydrogen bond donors in the terpenoid synthase active site assist the metal cluster in this function. That the trinuclear metal cluster is conserved for catalysis by terpenoid synthases from many domains of life suggests a common ancestry for this family of enzymes in the evolution of terpenoid biosynthesis.

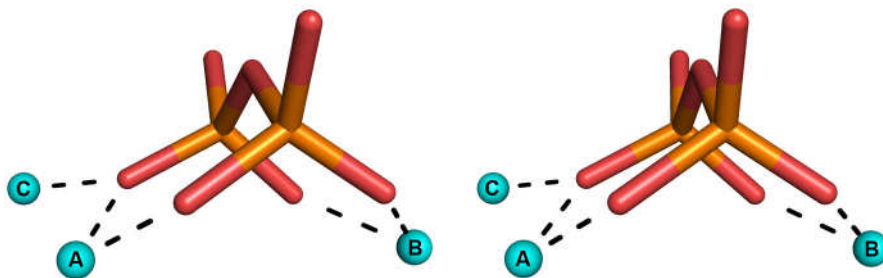


Figure 8.4. Stereoview of the $\text{Mg}^{2+}_3\text{-PP}_i$ cluster from *epi*-isozizaene synthase. Dashed lines (black) represent metal-coordination interactions. The PP_i anion forms 6-membered ring chelates with Mg^{2+}_A and Mg^{2+}_B , both of which adopt distorted sofa conformations.

Chapter 9: Future Directions

Terpenoid cyclases initiate and chaperone cyclization reactions to generate a multitude of structurally complex terpenoid products with precise regio- and stereo-specificity. The striking diversity of the terpenome is a direct result of the plasticity of the terpenoid synthases (Segura, 2003). It has been shown that the active site of a terpenoid synthase is predominantly lined with inert amino acids, which play a minimal role in the chemistry of catalysis beyond serving as a template and chaperone for the reaction (Christianson, 2008). The plasticity of the terpenoid cyclase active site has been studied in many systems (Greenhagen, 2006; Yoshikuni, 2006; Aaron, 2010).

A study of the sesquiterpene cyclase γ -humulene synthase provides an excellent example of the engineering potential of terpenoid cyclases. *Abies grandis* γ -humulene synthase, a promiscuous sesquiterpene cyclase, produces a mixture of 52 different terpenoid products. In the absence of a crystal structure of γ -humulene synthase, a homology model based on the known 5-*epi*-arisolochene synthase structure was used to identify “plasticity” residues in the γ -humulene synthase active site. The altered product profiles of a library of single-site mutants of the “plasticity” residues were determined and used to develop an algorithm to rationally design mutants using a combination of single-site mutations, based on the hypothesis that each plasticity residue is independent, meaning that the effect of a single mutation on the reaction mechanism is the same for the WT or any mutant form of the enzyme. Using their rational design algorithm, two to five

mutations were combined to create mutant enzymes with up to 13 times greater relative yields of the preferentially desired sesquiterpene product (Yoshikuni, 2006).

In this work, *epi*-isozizaene synthase (EIZS) has been identified as an excellent model system for studying the structure-function relationships of sesquiterpene cyclases. EIZS is a stable, monomeric enzyme that readily forms crystals which diffract to $\sim 1.6 \text{ \AA}$ resolution, and accommodates single amino acids mutations to active site residues, facilitating its potential use as a template for the rational design of novel terpenoid cyclases. Proposed experiments to continue this work include completing a GC-MS analysis of the products of the aliphatic active site mutants (L72V, A236G, and V329A) discussed in Chapter 7, to determine whether these small modifications to the contour of the active site result in perturbed product ratios. In addition, determining the kinetic activity, product-arrays by GC-MS, and crystal structures of several additional EIZS active site mutants, namely F95A, F332A, H333A, and W325F, would provide a thorough understanding of which residues directly affect the chaperoned cyclization cascade. Furthermore, crystal structures of the active site mutants provide an accurate picture of the enzymatic template, which can be used for modeling and quantum chemical calculations. To test the hypothesis that the product of a terpenoid cyclase can be predicted by how well the contour of the active site complements the shape of the product, modeling software, such as AutoDock (Morris, 2009), will be tested to determine a matching score for the respective enzymatic products of each EIZS mutant. These scores will be used to predict and test EIZS double and triple mutants, and to facilitate engineering new terpenoid cyclization templates.

It is important to remember that protein crystal structures provide a static picture of a dynamic system. Therefore, the orientation of the residues that form the active site cyclization template may occupy alternative conformations when the enzyme binds FPP in the closed conformation, with respect to the observed positions of the side chains in the Mg^{2+} - PP_1 -BTAC complexes determined. Additional proposed crystallography experiments include determining crystal structures of WT and mutant EIZS with substrate, or intermediate, analogues in order to observe the position of a partially folded substrate in the active site. A crystal structure containing a partially folded intermediate would offer insight into the role of the active site aromatic residues in stabilizing the cationic intermediates via cation- π interactions.

The ultimate goal of the structure-function studies of the terpenoid cyclases is to increase our understanding of these enzymes to the point where it is possible to systematically alter the function of a terpene cyclase using a rational design strategy. The potential terpenoid rational design has also recently led to the launch of Allylix, a start-up company aiming to exploit the versatility and plasticity of these enzymes to cost effectively produce useful commercial quantities of useful and novel terpenoids (Allylix, 2010). Exploiting the specificity and efficiency of these enzymes may have profound effects on the large-scale production of terpenoid products useful in the food, cosmetics and pharmaceutical industries.

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