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Probing the Catalytic Mechanism of S-Ribosylhomocysteinase (LuxS) with Catalytic Intermediates and Substrate Analogues

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Abstract

S-Ribosylhomocysteinase (LuxS) cleaves the thioether bond in S-ribosylhomocysteine (SRH) to produce homocysteine (Hcys) and 4,5-dihydroxy-2,3-pentanedione (DPD), the precursor of the type II bacterial quorum sensing molecule (AI-2). The catalytic mechanism of LuxS comprises of three distinct reaction steps. The first step involves carbonyl migration from the C1 carbon of ribose to C2 and the formation of a 2-ketone intermediate. The second step shifts the C=O group from C2 to C3 position to produce a 3-ketone intermediate. In the final step, the 3-ketone intermediate undergoes a β -elimination reaction resulting in the cleavage of the thioether bond. In this work, the 3-ketone intermediate was chemically synthesized and shown to be chemically and kinetically competent in the LuxS catalytic pathway. Substrate analogues halogenated at the C3 position of ribose were synthesized and acted as time-dependent inhibitors of LuxS. The time dependence was caused by enzyme-catalyzed elimination of halide ions. Examination of the kinetics of halide release and decay of the 3-ketone intermediate catalyzed by wild-type and mutant LuxS enzymes revealed that Cys-84 is the general base responsible for proton abstraction in the three reaction steps, whereas Glu-57 likely facilitates substrate binding and proton transfer during catalysis.

Keywords

Quorum sensing; autoinducer 2; LuxS; S-ribosylhomocysteinase; catalytic mechanism; inhibitor

Introduction

Quorum sensing (QS) controls bacterial gene expression in response to cell density and is mediated by the production and detection of small signaling molecules known as autoinducers (AIs).¹ Type 1 QS is species-specific; each bacterial species uses a unique AI or a unique combination of AIs for intraspecies communication. Different bacterial species also share a common signal, AI-2, for interspecies communication. AI-2 is biosynthesized from S-adenosylhomocysteine (SAH) through the sequential action of nucleosidase Pfs and a lyase, S-ribosylhomocysteinase (LuxS).² Pfs converts SAH into adenine and S-ribosylhomocysteine (SRH). The latter is further processed by LuxS into homocysteine and 4,5-dihydroxy-2,3-

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pentanedione (DPD). DPD is the AI-2 precursor; it is unstable in aqueous solution and undergoes spontaneous cyclization to form a mixture of several furanones. Different bacteria apparently sense the different furanone forms. For example, *Salmonella typhimurium* responds to (2*R*,4*S*)-2-methyl-2,3,3,4-tetrahydroxytetrahydrofuran,³ whereas *Vibrio harveyi* specifically recognizes the borate diester of (2*S*,4*S*)-2-methyl-2,3,3,4-tetrahydroxytetrahydrofuran.⁴ Because QS regulates many bacterial behaviors such as virulence and biofilm formation, the enzymes and receptors involved in AI production and/or detection are currently being pursued as targets for designing novel antibacterial agents.⁵

LuxS is a metalloenzyme, containing a tetrahedrally coordinated Fe²⁺ ion in its active site.^{6,7} The proposed mechanism involves two sequential carbonyl migration steps followed by a β -elimination reaction (Scheme 1).⁷ Because SRH exists predominantly in the hemiacetal form under physiological conditions, the LuxS active site presumably binds the hemiacetal and catalyzes its ring opening. The resulting aldehyde binds to the metal ion, replacing a metal-bound water molecule. Coordination to the metal increases the acidity of the C2 proton, resulting in its abstraction by a general base, which has been proposed to be Cys-84 in *Bacillus subtilis* LuxS (BsLuxS).^{8–10} The enediolate **2** formed undergoes ligand exchange, shifting the metal from C1 to the C2 OH group, presumably assisted by a second base/acid (likely Glu-57), to give enediolate **3**. Reprotonation at the C1 position (presumably by Cys-84) generates a 2-keto intermediate **4**. Repetition of the above sequence shifts the carbonyl group to the C3 position to give a 3-keto intermediate **7**. Finally, intermediate **7** undergoes a β -elimination reaction to release homocysteine and DPD. Glu-57 has been proposed as the general base responsible for the elimination reaction.^{8–10}

We have previously shown that the LuxS reaction in D₂O results in deuterium incorporation into product DPD, indicating the presence of a proton transfer step(s) during catalysis (instead of hydride transfer).⁷ Experiments with specifically deuterium-labeled substrates ([2-²H] and [3-²H]SRH) confirmed the stereochemical course of the proton transfer steps in the proposed mechanism.¹¹ Several lines of evidence have provided strong support for the intermediacy of 2-ketone **4**. First, NMR studies with [2-¹³C]SRH revealed a transient species with a chemical shift value of δ 213, suggesting a C=O group at the C2 position.⁸ Second, 2-ketone **4** has been chemically synthesized and shown to be chemically and kinetically competent against LuxS.⁸ Finally, the X-ray crystal structure of 2-ketone **4** bound to a catalytically inactive LuxS mutant (C84A) showed that the C2 carbonyl oxygen was directly coordinated with the catalytic metal ion.⁹ When [3-¹³C]SRH was used as a substrate, a peak at δ 214 was also observed, suggesting that 3-ketone **7** is likely also a bona fide intermediate on the catalytic pathway.⁸

Mutagenesis studies have shown that Glu-57 and Cys-84 are crucial residues for LuxS catalysis.⁷ Mutation of Cys-84 [or Cys-83 in *V. harveyi* LuxS (VhLuxS)] to alanine rendered the enzyme inactive toward SRH, whereas the E57A mutant had only trace amount of activity. In free *Escherichia coli* LuxS (EcLuxS), Cys-83 (corresponding to Cys-84 in BsLuxS) has a pK_a value of <6.¹⁰ These findings are consistent with the proposed role of Cys-84 and Glu-57 as general acids/bases. However, the precise roles of Glu-57 and Cys-84 in each catalytic step have been difficult to determine, due to their involvement in multiple catalytic steps. In this work, we have chemically synthesized the 3-ketone **7** and demonstrated its chemical and kinetic competence in the LuxS reaction. The availability of this intermediate has also provided a unique opportunity for us to assess the function of Glu-57 and Cys-84 during the last catalytic step (β -elimination reaction). In addition, we have synthesized two SRH analogues modified at the C3 position and used them as mechanistic probes to assess the function of Glu-57 and Cys-84 during the first catalytic step.

Results and Discussion

Synthesis of 3-Ketone Intermediate 7 (Scheme 2)

Synthesis of 3-ketone **7** started from D-xylofuranoside **12**,¹² which was prepared in two steps from the commercially available D-xylose. The hydroxyl group of **12** was protected with a p-methoxybenzyl (PMB) group to give xylofuranoside **13**, which was subsequently converted into hemiacetal **14** by treating with potassium tert-butoxide in DMSO followed by reaction with iodine in THF-H₂O.^{12, 13} Reduction of hemiacetal **14** with NaBH₄ gave diol **15**. The two hydroxyl groups of **15** were protected with PMB groups to give the fully protected xylitol **16**. Hydrolysis of the acetonide group was achieved by treatment with 60% AcOH and the resulting diol **17** was converted to cyclic sulfate diester **18** by reacting with thionyl chloride followed by oxidation of the resulting sulfite with RuCl₃ and NaIO₄. It is worth noting that formation of the cyclic sulfate diester is crucial for the subsequent nucleophilic substitution reaction. It serves the dual purpose of activating the C5-OH as a leaving group and protecting the C3-OH group (numbering based on the xylose structure). In our previous synthesis of the 2-keto intermediate (**4**),⁸ the cyclic ester was the only acceptor that reacted with the incoming nucleophile (homocysteine) to give appreciable amount of the desired product, while all other activation/protection strategies failed. Presumably, cyclization locks the densely functionalized xylitol into a conformation that helps expose the C5 carbon for nucleophilic attack. Thus, treatment of cyclic sulfate **18** with properly protected homocysteine⁸ in the presence of BuLi in DMF readily opened the ring at the primary carbon position (C5) to give the desired thioether **19**. The exposed C3 hydroxyl group was oxidized with Dess-Martin periodinane to give 3-ketone **20**. Global deprotection of ketone **20** with TFA produced the target compound (3-ketone **7**), with an overall yield of 18% (from **12**).

Synthesis of S-[3-Bromo-3,5-dideoxy-D-ribofuranos-5-yl]-L-homocysteine 10 ([3-Br]SRH, Scheme 3)

Diol **21**,¹⁴ prepared from the commercially available D-xylose, was treated with TBDPS-Cl to protect its less hindered 5-OH group.¹⁵ The resulting alcohol **22** was treated with N-bromosuccinimide (NBS) and triphenylphosphine (TPP) in refluxing chlorobenzene to give bromide **23**.¹⁶ Treatment of **23** with tetrabutylammonium fluoride (TBAF) in THF removed the silyl group at C5 position and the free hydroxyl group was activated by conversion into tosylate **24**. Nucleophilic substitution of tosylate **24** with the protected homocysteine gave thioether **25**, which was deprotected with TFA to afford the target compound [3-Br]SRH (6.6% overall yield from xylose **21**).

Synthesis of S-[3-Fluoro-3,5-dideoxy-D-ribofuranos-5-yl]-L-homocysteine 11 ([3-F]SRH, Scheme 4)

Synthesis of [3-F]SRH started from 5-O-benzyl-1,2-O-isopropylidene- α -D-xylofuranose **26**.¹⁷ Our initial attempts to directly fluorinate **26** with DAST or Deoxo-Fluor¹⁸ failed to provide 3-deoxy-3-fluororibofuranose **29**. We then utilized Mikhailopulo's approach¹⁹ for the synthesis of methyl 3-deoxy-3-fluororiboside **28**. Thus, removal of the isopropylidene of **26** with excess HCl (AcCl/MeOH/CH₂Cl₂/H₂O)²⁰ gave xylofuranoside **27** as a mixture of two anomers (α : β 1:1.4, 88% yield). Fluorination of the α / β -anomers of **27** with DAST/pyridine afforded **28** in 21% yield in addition to the corresponding *lyxo*-epoxide.¹⁹ We found that treatment (72 h) of **27** (α : β , 1:1.4) with Deoxo-Fluor (1.5 equiv) in pyridine also produced **28** (α : β , 1:1.1; 33%) plus *lyxo*-epoxide (32%), unreacted **27** (14%, α -anomer only), and some olefinic byproduct (5%). Re-introduction of the isopropylidene protecting group gave compound **29** which, upon debenzoylation at C5 position, gave alcohol **30**. Alcohol **30** was activated by tosylation and coupled with homocysteine to give thioether **32**, which was then deprotected with TFA to afford the target molecule [3-F]SRH in 7.8% overall yield (from **27**).

Chemical and Kinetic Competence of 3-Ketone Intermediate **7**

Treatment of 3-ketone **7** with LuxS resulted in the rapid release of homocysteine as detected by reaction with 5,5'-dithio-bis-(2-nitrobenzoic acid) (DTNB). The formation of DPD was also confirmed by the addition of 1,2-phenylenediamine to the reaction and isolation/analysis of the quinoxaline derivative by HPLC, MS, and NMR spectroscopy.⁷ In the absence of LuxS, the 3-ketone intermediate was stable for several weeks at 4 °C without any sign of decomposition as judged by the DTNB assay or derivatization with 1,2-phenylenediamine. Thus, 3-ketone **7** is chemically competent in the LuxS catalytic pathway. To demonstrate its kinetic competence, 3-ketone **7** was assayed against various Co(II)-substituted LuxS forms and its kinetic constants were compared with those of SRH and 2-ketone intermediate **4** (Table 1). Our earlier studies have shown that Co-LuxS has virtually identical catalytic properties to the native Fe-LuxS but is much more stable.⁷ Surprisingly, although wild-type VhLuxS exhibited higher catalytic activity toward 3-ketone intermediate **7** ($k_{\text{cat}}/K_{\text{M}} = 2.4 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$) than that of SRH ($k_{\text{cat}}/K_{\text{M}} = 1.1 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$) or 2-ketone intermediate **4** ($k_{\text{cat}}/K_{\text{M}} = 1.0 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$), both BsLuxS and EcLuxS were less active toward 3-ketone **7** than SRH or 2-ketone **4** (Table 1). The reduced catalytic activities were due to an increase in the K_{M} values (e.g., EcLuxS has K_{M} values of 17 and 170 μM for SRH and 3-ketone **7**, respectively). This suggests that the 3-ketone intermediate has restricted access to BsLuxS and EcLuxS active sites (slower binding than SRH or ketone **4**). During normal catalysis (with SRH as substrate), the 3-ketone intermediate is generated in situ and thus not affected by the accessibility. To test this notion, we compared the kinetics of binding of intermediates **4** and **7** to C84A Co-BsLuxS. This mutant has no detectable activity toward SRH or ketone **4** and only a trace amount of activity to ketone **7** (vide infra). The Co^{2+} ion provides a sensitive probe to monitor the binding of these intermediates to the enzyme active site. As shown in our previous work,⁹ 2-ketone **4** bound to the enzyme and reached the final E•S complex immediately. On the other hand, the binding of 3-ketone **7** was time-dependent and the absorption spectra kept changing during the 210-min incubation period (Figure 1). The changes in the d-d transition bands at ~550, 630, and 680 nm indicated changes in the metal environment during this period. The simplest explanation is that the ketone intermediate rapidly forms an initial E•S' complex, which slowly converts into the final, productive E•S complex. Alternatively, one might argue that the time-dependent spectral change was caused by slow turnover of the bound intermediate. However, depletion of the 3-ketone intermediate would be expected to return the spectrum to that of the free enzyme, as previously observed for the wild-type enzyme.⁷

We next tested the 3-ketone intermediate against several catalytically impaired VhLuxS mutants. The mutations in these enzyme variants have greatly slowed down their chemical steps and therefore their overall activities should be less sensitive to the rate of substrate binding. Indeed, C83A VhLuxS exhibited a low but real activity toward 3-ketone **7** ($k_{\text{cat}}/K_{\text{M}} = 31 \text{ M}^{-1} \text{ s}^{-1}$), whereas it had no detectable activity toward SRH and 2-ketone **4**. The S6A mutant also had higher activity toward 3-ketone **7** than 2-ketone **4** or SRH. Several other mutants (E57D, R39M, and H11Q) are apparently still limited by the slow binding of the 3-ketone intermediate, as evidenced by their greatly increased K_{M} values (Table 1). However, all of the enzyme variants showed equal or substantially higher k_{cat} values toward 3-ketone **7** than either 2-ketone **4** or SRH (within the margin of experimental error, which is ~2-fold). Taken together, the kinetic data indicate that the 3-ketone intermediate is kinetically competent.

Time-Dependent Inhibition of LuxS by [3-Br]SRH and [3-F]SRH

[3-Br]SRH exhibited time-dependent inhibition of BsLuxS (Figure 2). Since the observed inhibition was reversible (vide infra), it ruled out the possibility of irreversible inactivation. The observed inhibition appears to involve the formation of an initial enzyme-inhibitor complex, E•I, which slowly turns into a tighter enzyme-inhibitor complex, E•I*. The inhibition kinetics can be described by the equation

where K_I is the equilibrium constant for the formation of the initial $E \cdot I$ complex, k is the rate constant for the conversion of the $E \cdot I$ complex to the tighter $E \cdot I^*$ complex, and K_I^* represents the dissociation constant of the $E \cdot I^*$ complex. The K_I and K_I^* values of [3-Br]SRH against BsLuxS were determined to be 7.9 μM and 1.4 μM , respectively (Table 2). Similar time-dependent inhibition of EcLuxS and VhLuxS was also observed. The rate of conversion from $E \cdot I$ to $E \cdot I^*$ (k value) was estimated to be $\sim 0.2 \text{ h}^{-1}$ (for VhLuxS). In order to gain insight into the mechanism of inhibition, we also synthesized [3-F]SRH and tested it against BsLuxS. Again, time-dependent inhibition was observed. For [3-F]SRH, the conversion of $E \cdot I$ into $E \cdot I^*$ occurred much faster than [3-Br]SRH (on the order of minutes instead of hours), although an accurate measurement of the k value was not possible. It should be noted that all of the values in Table 2 are only apparent values, as true time zero (for the measurement of K_I values) and time infinite (for the measurement of K_I^* values) are not possible.

Mechanism of Inhibition by [3-Br]SRH and [3-F]SRH

We originally designed [3-Br]SRH as a mechanism-based inactivator (suicide inhibitor) of LuxS. We envisioned that LuxS would bind to [3-Br]SRH, catalyze its ring opening, and shift the C=O group from the C1 to C2 position (Scheme 5, pathway a). The resulting α -bromoketone (**34**) would undergo nucleophilic attack by Cys-84, forming a covalent, irreversible adduct between the enzyme and the inhibitor (**35**). To our surprise, several observations indicated that inhibition of LuxS by [3-Br]SRH did not follow this mechanism. Preincubation of LuxS with [3-Br]SRH (or [3-F]SRH) followed by high dilution into the assay solution resulted in nearly full recovery of the LuxS activity (data not shown). Electrospray ionization mass spectrometric (ESI-MS) analysis of the enzyme/inhibitor complex showed only unmodified LuxS (no enzyme-inhibitor adduct). [3-F]SRH, which was expected to undergo slower S_N2 reaction than [3-Br]SRH, showed faster $E \cdot I$ to $E \cdot I^*$ conversion.

To determine the mechanism of inhibition, several additional experiments were carried out. First, we examined whether Br^- (or F^-) was released during enzyme inhibition. Several BsLuxS variants were incubated with [3-Br]SRH and the reactions were monitored by paper chromatography in acetone/ H_2O (7:2). The Br^- ion was stained with AgNO_3 (enhanced by FITC) to appear as a brown spot with an R_f value of ~ 0.60 . Incubation of [3-Br]SRH with wild-type or E57D BsLuxS for 10 h resulted in the distinctive brown spot (R_f of 0.59) (Figure 3). However, Br^- release was not detected for C84S BsLuxS or in the absence of any enzyme, even after prolonged incubation (20 h). Next, the BsLuxS variants were incubated with [3-F]SRH and the release of F^- ion was monitored by proton-decoupled ^{19}F NMR spectroscopy (Figure 4). In the absence of enzyme, [3-F]SRH exhibited two peaks at δ -194.03 and -201.85, due to the presence of two hemiacetal anomers (α and β). Upon the addition of a catalytic amount of wild-type BsLuxS [40:1 (mol/mol) [3-F]SRH/LuxS], a new peak slowly developed at δ -120.40, as the substrate signals gradually decayed (over 60 min). When more LuxS (2:1

[3-F]SRH/LuxS) was used, the signal at δ -120.40 appeared immediately, with concomitant disappearance of the peaks at δ -194.03 and -201.85. The peak at δ -120.40 was assigned to the F^- ion by using NaF as the standard (δ -120.07). Similar results were obtained with E57D BsLuxS (data not shown). However, no F^- signal was detected when C84S BsLuxS was incubated with [3-F]SRH. A control experiment with [3-F]SRH alone (no LuxS) showed no release of F^- during the 2 h incubation period.

The above observations led us to propose an alternative mechanism (Scheme 5 pathway b). Again, LuxS opens the ribose ring to form aldehyde **33**. However, instead of the carbonyl migration in pathway a, the β -haloaldehyde undergoes a β -elimination reaction, presumably catalyzed by Cys-84, to form an α,β -unsaturated aldehyde **36**. Since [3-F]SRH reacted much faster than [3-Br]SRH, the elimination reaction must occur through the E1cb mechanism (i.e., rate-limiting abstraction of the C2 proton to form a carbanion intermediate first, followed by the fast release of halide ion). Aldehyde **36** would tautomerize into 1,2-dione **37**, and/or undergo addition of a water molecule to give the 2-ketone intermediate (**4**). To test the formation of **37**, [3-F]SRH was incubated with wild-type BsLuxS at 4 °C overnight and then treated with excess O-benzylhydroxylamine. ESI MS analysis of the resulting solution revealed the presence of several species with m/z ratios of 377, 397, and 482 (data not shown). The peak at m/z 397 is consistent with the structure of oxime **38** (plus Na^+), formed by the condensation of O-benzylhydroxylamine with unreacted [3-F]SRH (Scheme 6). The species at m/z 377 and 482 are most likely oximes **39** and **40**, respectively. The presence of oximes **39** and **40** suggests that 1,2-dione **37** was indeed formed during LuxS treatment of [3-F]SRH and subsequently converted into **39** and **40** by reacting with one and two molecules of O-benzylhydroxylamine, respectively. A similar set of products was observed when [3-Br]SRH was subjected to the same experimentation. In either experiment, no signal was observed for the oxime product that would be expected from the reaction of 2-ketone intermediate **4** and O-benzylhydroxylamine.

In order to gain further insight into the structural basis of LuxS inhibition by the SRH analogues, we co-crystallized Co-BsLuxS with [3-Br]SRH and determined the structure of the complex to 1.9 Å resolution. After an initial round of refinement without bound ligand, the F_o-F_c electron density map showed clear and strong density for a compound in the active site (Figure 5), which was originally modeled as the ring-opened 2-ketone **34** (Scheme 5). However, this compound did not refine well in the structure; there was not enough room to accommodate the larger Br atom at the C3 position, as it would clash with the Co^{2+} ion. When refinement was attempted with **34** in the active site, the rest of the ribosyl structure was pushed out of the observed density. Instead, the observed density is most consistent with the presence of 2-ketone intermediate **4**, which bears a hydroxyl group at the C3 position. In fact, the observed density and the refined conformation of the 2-ketone intermediate in the present structure are essentially indistinguishable from the previously reported structure of C84A BsLuxS bound with the same 2-ketone intermediate (**4**).⁹ None of the other structures that could potentially be derived from intermediate **36**, including the C3 epimer of intermediate **4** (*S* configuration at C3, which would be formed by the addition of a water molecule to the *si* face of intermediate **36**), were able to fit well to the observed electron densities. We therefore tentatively conclude that the species bound in the LuxS active site is most likely the 2-ketone intermediate (**4**), presumably formed by the addition of a water molecule to the *re* face of intermediate **36**. Why was the intermediate (**4**) not converted into DPD and homocysteine by the wild-type LuxS? According to the proposed mechanism (Scheme 5), the β -elimination reaction is accompanied by the protonation of Cys-84. Subsequent addition of a water molecule should not change the ionization state of Cys-84. Thus, in the resulting enzyme-intermediate complex, Cys-84 would remain in the protonated form, which cannot act as a general base to catalyze the subsequent reaction steps. We have previously shown that the active-site residues of LuxS do not freely exchange protons with the bulk solvent.^{7,11} When the LuxS reaction was carried out with [3-²H]SRH as substrate, ~90% of the deuterium was transferred to the C2 position of DPD product, with only

minor loss to solvent. In the current co-crystal structure, Cys-84 sits at the bottom of a deep pocket and is completely shielded from the solvent by the bound ligand. Since treatment of [3-Br]SRH or [3-F]SRH with active LuxS in the solution phase did not result in the accumulation of 2-ketone intermediate **4**, we hypothesize that the accumulation of intermediate **4** in the crystals might be a result of crystal contacts, which could limit conformational changes and slow down/prevent the release of bound ligands. In any rate, the uncertainty in the ligand identity does not detract from our main conclusion that LuxS catalyzes the release of halides from [3-Br]SRH and [3-F]SRH.

Function of Cys-84 and Glu-57

The LuxS mechanism can be divided into three distinct chemical steps; the first step effects the migration of the C=O group from the C1 to C2 position (from **1** to **4**), the second step shifts the C=O group from the C2 to C3 position (from **4** to **7**), and the third step completes the β -elimination reaction (from **7** to **9**) (Scheme 1). Because Glu-57 and Cys-84 participate in all three steps, their precise roles in each catalytic step have been difficult to assess. [3-F]SRH and [3-Br]SRH, while disappointing as suicide inhibitors, provide alternative substrates to examine the function of active-site residues during the first catalytic step, since the E1cb elimination reaction involves initial deprotonation at the C2 position, which closely mimics the deprotonation event in the first catalytic step. Because E57D BsLuxS catalyzes the elimination of Br⁻ and F⁻ with similar rates as the wild-type enzyme (Figure 3) whereas the C84S mutant is inactive, we conclude that Cys-84 is the GA/GB responsible for the transfer of the C2 proton to the C1 position during the first C=O migration step. We have previously shown that Cys-83 of EcLuxS has a pKa value of <6 and therefore exists in the thiolate form under physiological conditions.¹⁰ E57D VhLuxS has higher activity toward 2-ketone intermediate (**4**) than SRH ($k_{\text{cat}}/K_{\text{M}}$ values of 280 and 118 M⁻¹ s⁻¹, respectively), suggesting that Glu-57 also plays a role in the first step. Presumably, Glu-57 facilitates substrate binding and catalysis by initially forming a hydrogen bond with the C2-OH and subsequently transferring the proton from the C2-OH to the newly formed C1-OH group (Scheme 1). For the elimination reaction of [3-F]SRH and [3-Br]SRH, there is no proton transfer from the C2-OH to the C1-OH, and therefore Glu-57 has minimal effect on the reaction rate. We originally proposed that Glu-57 effected proton transfer by directly hydrogen bonding to the C1-OH.⁸ More recent X-ray structural studies reveal that Glu-57 is too far from the C1-OH to allow a direct transfer of the proton originally derived from the C2-OH to the C1 oxyanion.^{5e, 9} We now suggest that the proton transfer is relayed through a series of active-site residues and/or solvent molecules, eventually to His-11.⁹ His-11 acts a general acid donating the proton to the C1 oxyanion as it departs from the metal ion. Consistent with this proposal, His-11 is directly hydrogen bonded to the C1-OH of the bound 2-ketone intermediate (**4**),⁹ and mutation of His-11 results in ~1000-fold reduction in the LuxS activity (Table 1). Tyr-89 is likely a component of this proton relay system, as it is physically proximal to the bound substrate/intermediates (Figure 5) and essential for catalysis (JZ and DP, unpublished data). Since the second step of the LuxS mechanism (from **4** to **7**) is simply a repetition of the first step, we propose that His-11, Glu-57, Cys-84, and Tyr-89 play similar roles in the second catalytic step.

The function of the active-site residues during the final step of the LuxS reaction can be examined by using the 3-ketone intermediate (**7**) as substrate. We previously proposed Glu-57 as the general base responsible for proton abstraction from the C4 position during the β -elimination reaction. However, mutation of Cys-83 in VhLuxS to alanine resulted in a ~30-fold decrease in the k_{cat} value toward ketone **7**, indicating that Cys-83 plays a crucial role during the elimination reaction. We now suggest that Cys-83 is most likely the general base for C4 proton abstraction, on the basis of two considerations. First, Cys-83 is actually the more logical candidate than Glu-57, as proton abstraction at the C4 position is the same reaction as the C2 and C3 deprotonation reactions. Second, the elimination reaction presumably involves the

coordination of the C=O group of intermediate **7** to the metal ion (Scheme 1). This would place the C4 proton next to the thiolate group of Cys-83 and away from the side chain of Glu-57, as is the case in structures **1** and **4**. Since mutation of Glu-57 greatly reduces the activity of LuxS toward the 3-ketone intermediate (Table 1), Glu-57 is also a critical residue for the elimination reaction. We propose that, as in the first and second catalytic steps, Glu-57 facilitates the reaction by hydrogen bonding to the C4-OH and anchoring the substrate for the elimination reaction. Both E57A and C83A mutants retain small amounts of catalytic activities (Table 1). Therefore, while these two residues are important for catalysis, they are not essential for the elimination reaction. The metal ion and the active-site environment can substantially accelerate the elimination reaction. In this regard, serine protease mutants with the catalytic triad completely removed still possess significant protease activities.²¹

Conclusion

Through chemical synthesis and kinetic analysis of catalytic intermediates and substrate analogues, we have demonstrated that 3-ketone **7** is a bona fide intermediate in the LuxS reaction mechanism. Kinetic analysis of wild-type and mutant LuxS toward ketone **7** suggests that Cys-84 is most likely the general base during the final β -elimination reaction. Substrate analogues [3-F]SRH and [3-Br]SRH act as time-dependent inhibitors of LuxS, due to enzyme catalyzed elimination of the halide ions via an E1cb mechanism. By using these alternative substrates, we have demonstrated that Cys-84 is the general base for proton abstraction during the first C=O migration step and likely the second step as well.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1.

Absorption spectra showing the binding kinetics of 3-ketone intermediate **7** (0 or 1.0 mM) to C84A Co-BsLuxS (186 μ M). The complex was incubated at room temperature for 0, 5, 20, 40, 80, 120, and 210 min prior to spectral recording.

Figure 2.

Time-dependent inhibition of LuxS by [3-Br]SRH. At time 0, Co-VhLuxS (100 μ M) and [3-Br]SRH (0–3.0 mM) were mixed at pH 7.4 and 0 $^{\circ}$ C. At the indicated time points (0–5 h), 10- μ L aliquots were withdrawn and added to a solution containing 20 μ M SRH and 150 μ M DTNB (total volume of 1 mL) and the remaining LuxS activities were determined and plotted as a function of the preincubation time.

1 2 3 4 5

Figure 3. Paper chromatography showing LuxS-catalyzed Br^- release from $[3\text{-Br}]\text{SRH}$. Lane 1, wild-type LuxS; Lane 2, C84S LuxS; Lane 3, E57D LuxS; Lane 4, 3-Br-SRH only (no LuxS); Lane 5, NaBr (positive control).

Figure 4. ^{19}F NMR spectra (376.5 MHz) of LuxS-catalyzed F^- elimination. (A) [3- F]SRH (3.0 mM) and wild-type BsLuxS (75 μM or 1.5 mM) were mixed and incubated, and spectra were collected at the designated time points.

Figure 5.

Electron density observed in the active site of Co-BsLuxS co-crystallized with [3-Br]SRH. LuxS is shown in stick form with one subunit of the dimer in cyan and the other subunit in yellow. The Co^{2+} atom is shown as a pink sphere. The inhibitor molecule (**4**) is shown in green stick form. The blue cage is the $2F_o - F_c$ electron density map, calculated at 1.9 Å resolution and contoured at 1 σ . The red cage is an $F_o - F_c$ electron density map calculated after refinement with compound **4** omitted. The figure was prepared using Pymol (Delano Scientific)

Scheme 1.
Proposed mechanism of LuxS-catalyzed reaction

Scheme 2.

Synthesis of 3-Keto Intermediate 7^a

^aReagents and Conditions: (a) NaH, PMBCl, DMF, 0 °C-RT, overnight, 79%; (b) ^tBuOK, DMSO, 80 °C, 2 h; (c) I₂, pyridine/THF/H₂O, 1 h, 89% over two steps; (d) NaBH₄, MeOH, 0 °C-RT, overnight, 93%; (e) NaH, PMBCl, DMF, 0 °C-RT, 2 days, 85%; (f) 60% AcOH, dioxane, overnight, 82%; (g) SOCl₂, Et₃N, CH₂Cl₂, 78-0 °C, 1 h; (h) RuCl₃, NaIO₄, CCl₄, CH₃CN, H₂O, 1 h, 78% over two steps; (i) BuLi, DMF, BocNH-CH(CH₂CH₂SH)-CO₂tBu, 0 °C-RT, 4 h; (j) H₂SO₄, THF/H₂O, 0 °C, 1 h, 80% over two steps; (k) Dess-Martin periodinane, CH₂Cl₂, 1 h, 86%; and (l) 10% anisole in TFA, 7 h, 75%.

Scheme 3. Synthesis of [3-Br]SRH^a

^aReagents and Conditions: (a) TBDPS-Cl, imidazole, CH₂Cl₂, 0 °C-RT, 2 h, 75%; (b) TPP, NBS, imidazole, chlorobenzene, reflux, 4 h, 48%; (c) TBAF, THF, 0 °C-RT, 4 h, 75%; (d) TsCl, Et₃N, CH₂Cl₂, 0 °C-RT, overnight, 70%; (e) BocNH-CH(CH₂CH₂SH)-CO₂tBu, K₂CO₃, acetone, reflux, 3 h, 49%; and (f) TFA/anisole/H₂O, 0 °C-RT, 3.5 h, 71%.

Scheme 4. Synthesis of [3-F]SRH^a

^aReagents and Conditions: (a) AcCl, MeOH, CH₂Cl₂, H₂O; (b) Deoxo-Fluor, pyridine/CH₂Cl₂, 72 h, 33%; (c) Dry HCl/acetone, 42 h, 52%; (d) H₂, Pd/C, EtOH, 24 h, 92%; (e) TsCl, Et₃N, CH₂Cl₂, 0 °C-RT, overnight, 83%; (f) BocNH-CH(CH₂CH₂SH)-CO₂tBu, K₂CO₃, acetone, reflux, 4 h, 84%; (g) TFA/H₂O/anisole, 0 °C-RT, 6 h, 71%.

Scheme 5.

Possible Mechanisms of LuxS Inhibition by [3-Br]SRH and [3-F]SRH.

Scheme 6.
Chemical Trapping of Intermediates **33** and **37**

Table 1
Catalytic Properties of Co-LuxS Variants toward SRH, 2-Ketone **4**, and 3-Ketone **7**

enzyme	3-ketone 7			2-ketone 4			SRH	
	$k_{cat}(s^{-1})$	$K_M(\mu M)$	$k_{cat}/K_M(M^{-1} s^{-1})$	$k_{cat}(s^{-1})$	$K_M(\mu M)$	$k_{cat}/K_M(M^{-1} s^{-1})$	$k_{cat}(s^{-1})$	$K_M(\mu M)$
WT (Bs) ^a	0.038 ± 0.001	3.6 ± 0.1	11000	0.031 ± 0.004	1.5 ± 0.3	21000	0.030 ± 0.005	2.3 ± 0.4
WT (Ec)	0.40 ± 0.06	170 ± 30	2300	ND ^c	ND	ND	0.38 ± 0.01	17 ± 1
WT (Vh) ^b	0.64 ± 0.02	27 ± 2	24000	0.40 ± 0.02	38 ± 1	10000	0.40 ± 0.01	37 ± 4
C83A (Vh)	0.022 ± 0.002	700 ± 120	31	ND	ND	ND	ND	ND
E57A (Vh)	0.012 ± 0.001	98 ± 13	120	ND	ND	ND	ND	ND
E57D (Vh) ^a	0.026 ± 0.003	730 ± 120	36	0.008 ± 0.001	29 ± 2	280	ND	ND
R39M (Vh) ^b	0.016 ± 0.001	150 ± 20	110	0.008 ± 0.001	22 ± 2	350	ND	ND
H11Q (Vh) ^b	0.024 ± 0.002	520 ± 60	47	0.008 ± 0.001	39 ± 2	200	ND	ND
S6A (Vh) ^b	0.14 ± 0.01	40 ± 3	3400	0.102 ± 0.002	96 ± 7	1100	0.027 ± 0.005	41 ± 5
								660

^aData for 2-ketone **4** and SRH are from ref⁸.

^bData for 2-ketone **4** and SRH are from ref⁹.

^cNot determined.

Table 2
Inhibition Constants of SRH Analogues against Co-LuxS Variants

Enzyme	[3-F]SRH		[3-Br]SRH	
	K_I (μM)	K_I^* (μM)	K_I (μM)	K_I^* (μM)
BsLuxS	10.6 ± 2.4	0.70 ± 0.26	7.9 ± 1.6	1.4 ± 0.4
EcLuxS	ND ^a	ND	47 ± 4	5.9 ± 0.5
VhLuxS	ND	ND	47 ± 4	1.6 ± 0.1

^a Not determined