

Human serine racemase structure/activity relationship studies provide mechanistic insight and point to position 84 as a hot spot for β -elimination function

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There is currently great interest in human serine racemase, the enzyme responsible for producing the NMDA co-agonist D-serine. Reported correlation of D-serine levels with disorders including Alzheimer's disease, ALS, and ischemic brain damage (elevated D-serine) and schizophrenia (reduced D-serine) has further piqued this interest. Reported here is a structure/activity relationship study of position Ser⁸⁴, the putative *re*-face base. In the most extreme case of functional reprogramming, the S84D mutant displays a dramatic reversal of β -elimination substrate specificity in favor of L-serine over the normally preferred L-serine-O-sulfate (~1200-fold change in k_{cat}/K_m ratios) and L (L-THA; ~5000-fold change in k_{cat}/K_m ratios) alternative substrates. On the other hand, the S84T (which performs L-Ser racemization activity), S84A (good k_{cat} but high K_m for L-THA elimination), and S84N mutants (nearly WT efficiency for L-Ser elimination) displayed intermediate activity, all showing a preference for the anionic substrates, but generally attenuated compared with the native enzyme. Inhibition studies with L-erythro- β -hydroxyaspartate follow this trend, with both WT serine racemase and the S84N mutant being competitively inhibited, with $K_i = 31 \pm 1.5 \mu\text{M}$ and $1.5 \pm 0.1 \text{ mM}$, respectively, and the S84D being inert to inhibition. Computational modeling pointed to a key role for residue Arg-135 in binding and properly positioning the L-THA and L-serine-O-sulfate substrates and the L-erythro- β -hydroxyaspartate inhibitor. Examination of available sequence data suggests that Arg-135 may have originated for L-THA-like β -elimination function in earlier evolutionary variants, and examination of available structural data suggests that a Ser⁸⁴-H₂O-Lys¹¹⁴ hydrogen-bonding network in human serine racemase lowers the pK_a of the Ser⁸⁴ *re*-face base.

The discovery of D-serine in the brain and its importance in modulating NMDA receptor activity provided the first *bona fide* example of a D-amino acid in human biology. The quest to uncover the source of this D-serine led to the identification of

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mammalian serine racemase (1). Wolosker *et al.* (1) successfully cloned human serine racemase (hSR)³ at the turn of the millennium. The observation that all D-serine apparently originates in L-serine added another significant branch to the complex metabolic network associated with L-serine and an important new signaling function for the amino acid (Fig. 1).

To be sure, L-serine already was known to possess an array of physiological functions, including serving as both the source of one-carbon equivalents in *N*⁵,*N*¹⁰-methylene tetrahydrofolate (utilized for DNA synthesis; *i.e.* installation of the 5-methyl group in the uracil ring to give the thymine base) and of the neurotransmitter and NMDAR co-agonist, glycine, through the action of a single pyridoxal phosphate (PLP) enzyme, serine hydroxymethyltransferase. L-Serine also serves a central role in maintaining redox homeostasis, because all glutathione equivalents originate in the L-serine backbone, with the sulfur atom from dietary methionine being installed at the β -carbon through the sequential action of two additional PLP-dependent enzymes, cystathionine β -synthase (CBS) and cystathionine eliminase (also known as cystathionine γ -lyase). L-Serine also serves as an important constituent of the phospholipidome and is one of three constituent amino acids of proteins (along with L-threonine and L-tyrosine) that underpin the phosphoproteome, as controlled by the action of protein phosphoserine kinase phosphatases.

From the point of view of neuronal signaling, both D-serine and glycine serve as co-agonists of the NMDA receptor (Fig. 2), binding at the "glycine site" but with the observation that D-serine is a more potent agonist than glycine itself, showing efficacy at several orders of magnitude lower concentration in a seminal study by Ascher and co-workers (2) in a rat hypoglossal motoneuron system (2–4). Whereas it had previously been thought that D-serine is biosynthesized by SR in astroglial cells, more recent evidence indicates that L-serine produced in the astroglia from 3-phosphoglycerate is actively shuttled to the neurons where SR is present and converts the L-serine to D-serine (5, 6).

As is illustrated in Fig. 2A, D-serine generated in the presynaptic neuron serves as stimulatory co-agonist of the postsynap-

³ The abbreviations used are: hSR, human serine racemase; SR, serine racemase; TEA, triethanolamine; Wat, water molecule; PLP, pyridoxal phosphate; CBS, cystathionine β -synthase; L-SOS, L-serine-O-sulfate; L-THA, L-threo- β -hydroxyaspartate; MBP, maltose-binding protein; L-ABH, L-aspartate β -hydroxamate; L-EHA, L-erythro- β -hydroxyaspartate; AOAA, aminoacetate; MD, molecular dynamics.

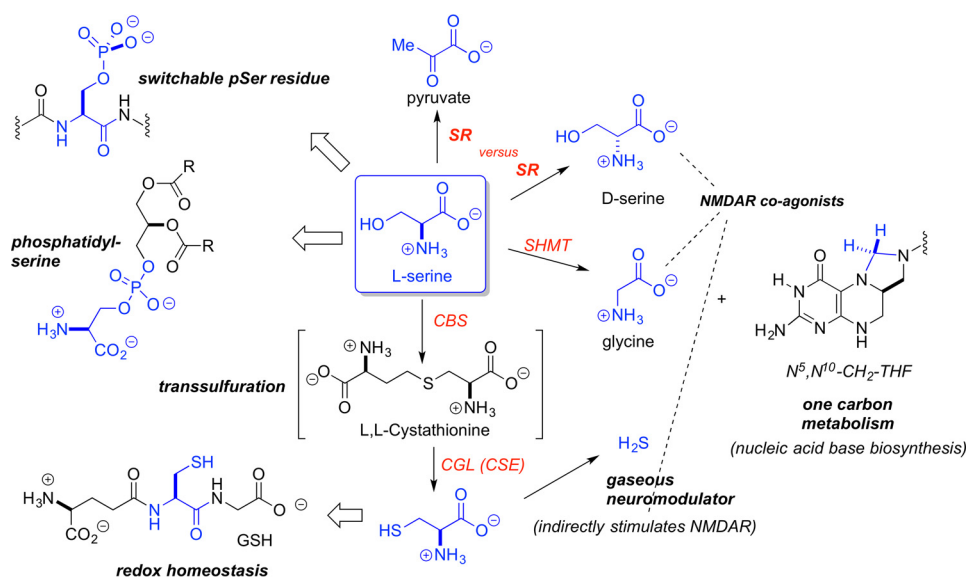


Figure 1. L-Serine as a central metabolite. L-Serine plays a central role in biology, from one-carbon metabolism to transsulfuration, phospholipid/phospho-protein function, and D-serine biosynthesis.

tic NMDA receptor, acting in concert with the primary agonist, L-glutamate. Of particular interest to our laboratory (7–13), the gaseous neurotransmitter, H_2S , produced by another PLP enzyme (CBS) in the brain also elicits an NMDA excitatory response (14), potentially via an adenylate cyclase-cAMP-dependent protein kinase-mediated mechanism (15). Indeed, both SR and CBS have emerged as potential targets for ischemic stroke, because there is evidence that both D-serine (16, 17) and H_2S (18) promote neuronal infarction following such a stroke event. Elevated D-serine levels have also been associated with Alzheimer's disease (19) and ALS (20), suggesting that SR may emerge as a potential target for neurodegenerative disease. On the other hand, low D-serine levels (*i.e.* SR hypofunction) have been correlated with schizophrenia (21–23). In contrast to these examples of SR dysregulation, in the healthy brain, basal D-serine signaling is essential for synaptic efficiency and long-term potentiation associated with learning and memory (24).

The hSR enzyme is known to be activated allosterically by ATP and requires a divalent cation (Mg^{2+} , Mn^{2+}) for activity (Fig. 2B) (25–27). The enzyme is reported to be post-translationally modified by phosphorylation (28), palmitoylation (29), and nitrosylation (30). SR levels can be modulated by ubiquitin tagging for proteasomal degradation (31). The C-terminal PDZ domain is important in protein–protein interactions, with PICK-1 (32, 33), GRIP-1 (34), and PSD-95 (35), for example. X-ray crystallographic structures of the *Schizosaccharomyces pombe* (36), maize (37), mouse (38), and rat and human (39) SR enzymes are available.

Mammalian SR has a type II β -eliminase fold reminiscent of the classical PLP-dependent enzyme, tryptophan synthase. Accordingly, it is perhaps not surprising that the enzyme catalyzes both the β -elimination of L-serine and its racemization to D-serine (Fig. 3). Mechanistically, a dual-base mechanism has been proposed, whereby Lys⁵⁶ serves as the *si*-face base (40), α -deprotonating an appropriately oriented external aldimine of L-serine, giving rise to a common, cofactor-stabilized carban-

ionic intermediate (41). Subsequent *re*-face protonation by the putative *re*-face base, Ser⁸⁴ (42), leads to the D-serine racemization product, whereas expulsion of the (presumably protonated) β -OH leaving group leads to pyruvate, the β -elimination product. L-Serine-O-sulfate (L-SOS) and L-threo- β -hydroxyaspartate (L-THA) are known to serve as very efficient alternative substrates for this latter β -elimination manifold (43). This overall mechanism is consistent with recent QM/MM calculations (44).

Results

We set out to examine the mechanism of hSR, with a particular focus on the influence of the putative *re*-face base upon reactivity. The assignment of Ser⁸⁴ as the *re*-face base itself raises a key mechanistic question, namely how can this residue have an appropriately low pK_a to perform this general acid/base function? Careful examination of available structural information suggests that Lys¹¹⁴ may serve to (de)protonate Ser⁸⁴ through a hydrogen bond network involving an essential water molecule (Fig. 4A). This putative Ser⁸⁴-Wat³⁷²-Lys¹¹⁴ hydrogen bond network appears to resemble the Ser-*cis*-Ser-Lys catalytic triad (Fig. 4B) (45–48) that is typically seen in the amidase signature enzyme family that includes peptide amidases (49–51) and fatty acid amide hydrolases (52–54) as well as some β -lactamases (55) and a recently described hydrazidase enzyme (56). To our knowledge, this model for serine acidification has not previously been proposed for a PLP enzyme.

To facilitate experimental studies, it was found that improved hSR solubility could be achieved by expressing the protein as an N-terminal maltose-binding protein (MBP) fusion construct (MBP-hSR). Literature reports of heterologous SR expression indicate that these efforts have been plagued with difficulty, leading to low yields of active enzyme, ranging from 1 to 2 mg/liter of culture, as summarized in Fig. 5 (40, 43, 57) and confirmed in our hands with both the N-terminal His- and GST-tagged constructs. However, the MBP-hSR

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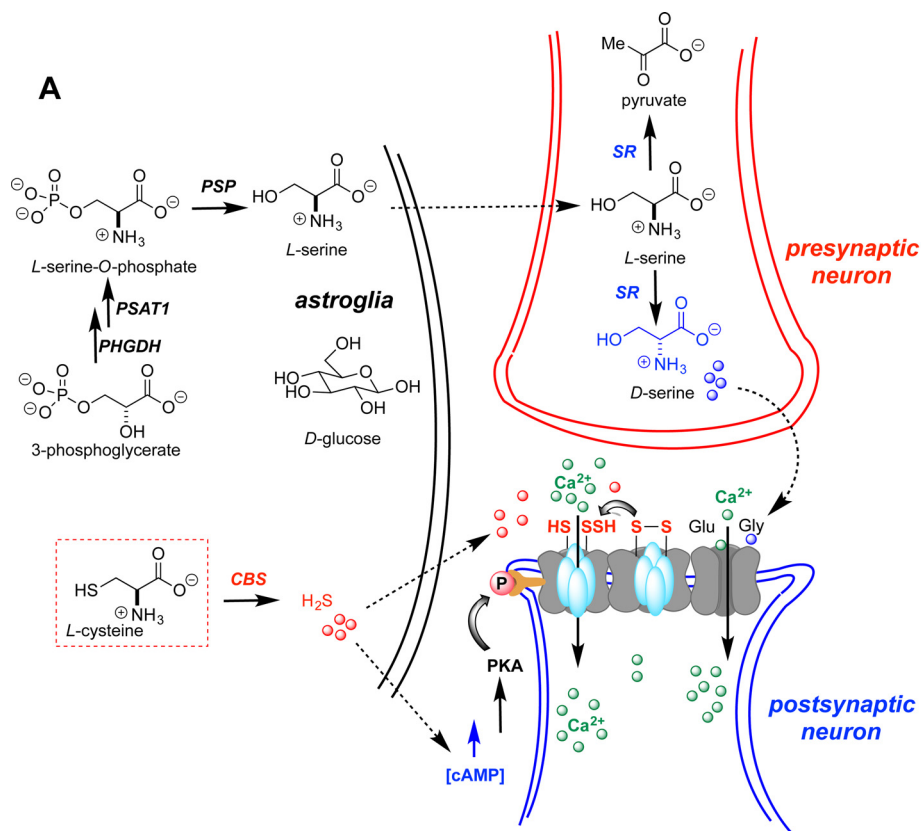


Figure 2. hSR: physiological role and 3D structure. A, biosynthesis of the neuromodulators H_2S and D-serine and their postulated roles in stimulation of the NMDA receptor (schematic); B, homology model for hSR based upon Protein Data Bank entries 3L6B (recombinant modified human; internal aldimine) and 2ZPU (*S. pombe*; substrate-modified internal aldimine). Blue, C-terminal PDZ domain (interacts with GRIP-1 and PICK-1); lavender, structural dication (Mg^{2+} , Ca^{2+} or Mn^{2+}); salmon, palmitoylation site (Thr²²⁷); red helical region, putative nitrosylation site (Cys¹¹³); green, ATP-binding site; red loop region, putative phosphorylation site (Thr⁷¹).

construct reproducibly gave ~ 15 mg of purified fusion protein. This is largely attributed to improved solubility. Pixel densitometric gel analysis demonstrates that whereas His–hSR showed only 7% soluble hSR protein, the MBP–hSR fusion is estimated to give 48% of the protein in the supernatant, a nearly 7-fold increase in solubility. The 15 mg of MBP–hSR translates to ~ 6.4 mg of hSR *versus* the 1–2 mg obtained for His–hSR, a significant improvement. Removal of the tag via factor Xa digestion followed by ATP column purification resulted in a doubling of specific activity, as expected.

Native PAGE experiments are indicative of a dimeric structure for the hSR–MBP fusion protein, consistent with previous reports and crystal structures for the SR (39). Interestingly, gel filtration (Sephacryl S-200) shows an apparent molecular mass of 247 Da for the new construct, suggestive of an oligomeric composition of 3.1. This may be reflective of an equilibrium between a dimeric and tetrameric form of MBP–hSR under the conditions of the gel filtration experiment. This notion would be consistent with a recent observation by Mozzarelli and co-workers (58) that hSR is capable of forming an active tetramer

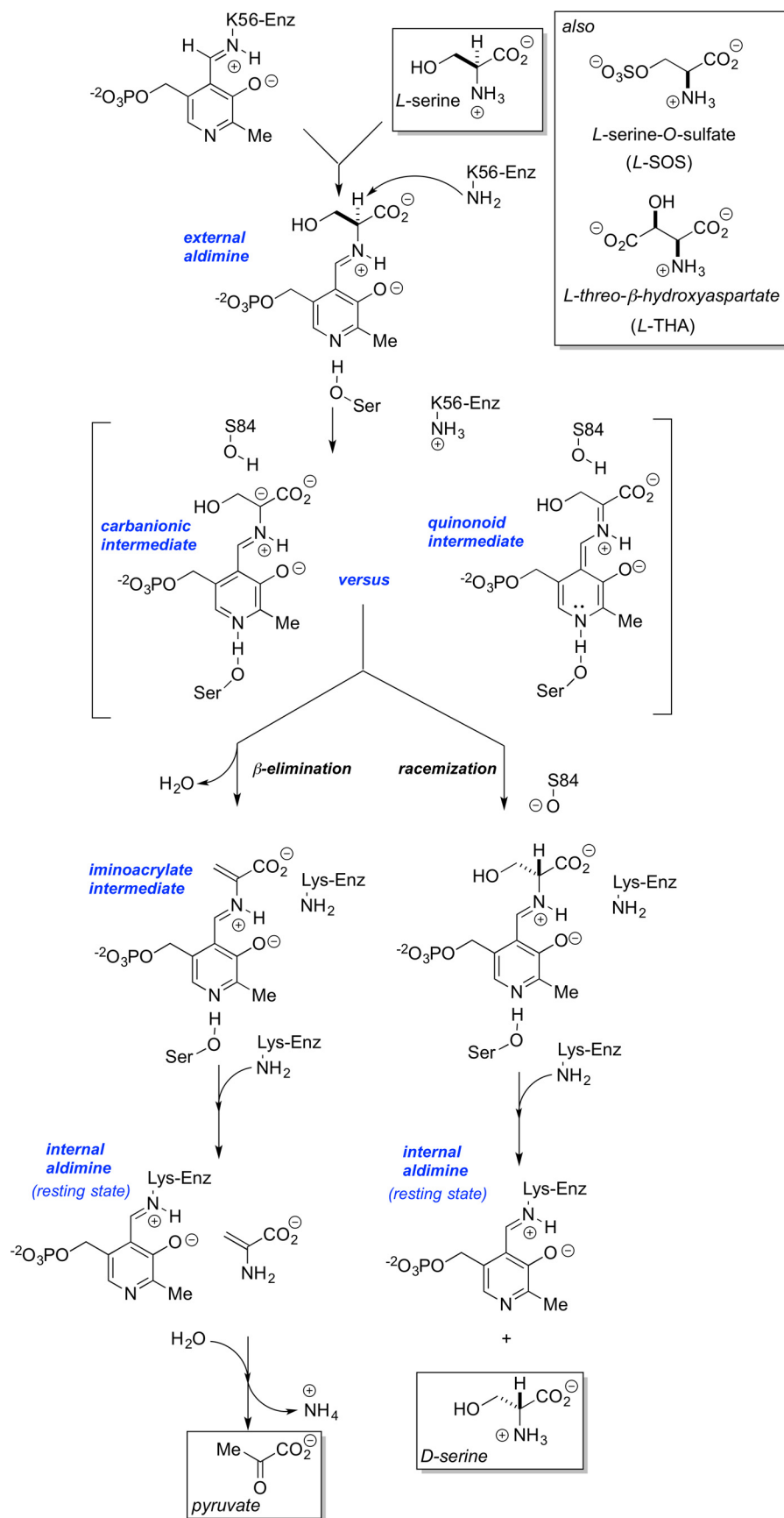


Figure 3. Mechanism of serine racemase. Shown is the racemization manifold versus the β -elimination manifold. Note that both carbanionic and fully delocalized quinonoid intermediate pathways can be considered.

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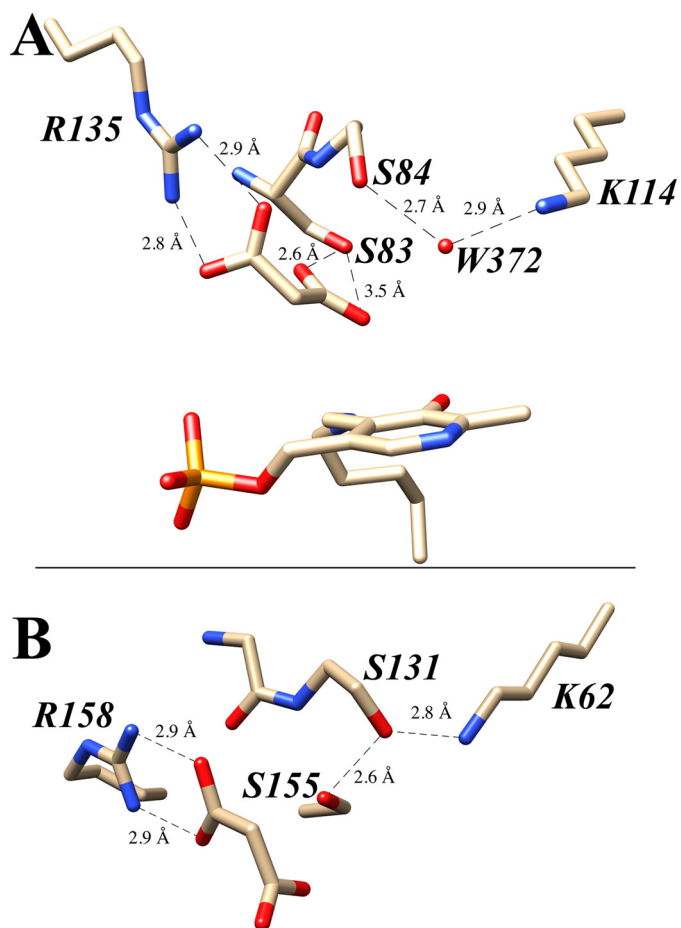


Figure 4. Acidity of Ser⁸⁴: Ser-(OH₂)-Lys versus Ser-(cis-Ser)-Lys triad. *A*, proposed hydrogen-bonding network involving the *re*-face base, Ser⁸⁴, and the neighboring Wat³⁷² molecule and Lys¹¹⁴ residue. *B*, comparison with the Ser¹⁵⁵-*cis*-Ser¹³¹-Lys⁶² catalytic triad in the amidase signature family of enzymes. Crystal structures 3L6B (hSR; *A*) and 1OCL (malonamidase; *B*) are used in this image; both structures contain a bound malonate as inhibitor (*A*) and product (*B*), respectively.

in the presence of ATP and the divalent cation (Ca²⁺ or Mg²⁺) normally associated with the protein.

Active site mutants/kinetic evaluation

We next chose to examine the influence of *re*-face base on enzyme function. In previous work by Wolosker and co-workers (40), mutagenesis of residues 151–154 led to a reversal of functional preference with L-serine as the substrate. As noted (Fig. 6), normally mammalian serine racemase favors L-serine elimination activity over L-serine racemization activity, in a ratio of ~4:1. Their most notable mutant, Q154D, displayed a dramatic reversal of these activities to a ratio of 1:3.

Focusing on the active site, previous reports had identified Ser⁸⁴ as the *re*-face base in homologous enzymes belonging to *S. pombe* (42) and slime mold (*Dictyostelium discoideum*) (59). In both cases, the *re*-face Ser to Ala mutant was constructed and resulted in loss of function for racemization. Utilizing the MBP-hSR platform, we generated the corresponding mammalian SR S84A mutant, in addition to the S84D, S84N, and S84T variants. There had been one earlier report of the former construct, but the activity of this mutant was only studied with the natural substrate, L-serine (60). The rationale was to examine

changes in hydrogen bond donor ability, charge, and sterics for putative *re*-face base by studying the behavior of these mutants across four assays: racemization and β -elimination reaction across a battery of three substrates.

The results are summarized in Table 1. Compared with the wild-type hSR, the S84D mutant shows the most dramatic difference in substrate preference as measured by the catalytic efficiency (*i.e.* measured k_{cat}/K_m value). Whereas the charged substrates L-SOS and L-THA are highly favored in wild-type SR, S84D shows a dramatic reversal of this preference. In the most pronounced case, for L-THA compared with L-serine, this preference changes from 100:1 in the wild-type to 1:50 (see Table 1). This represents a 5000-fold change in substrate preference. A similar ~1200-fold change is observed for L-SOS. It is important to note that this was not a “mutate-and-kill” effect, because the catalytic efficiency of L-serine only decreased ~6-fold in the S84D mutant. That PLP enzymes can exhibit significant catalytic promiscuity even with subtle changes has recently been highlighted by Patrick and co-workers (61) in detailing the promiscuous alanine racemase activity seen with mutant cystathionine- β -lyase, both in *Escherichia coli*.

The S84N mutant shows intermediate behavior, exhibiting neither the strong preference for charged substrates seen in the wild type nor the inverted substrate preference seen in the S84D mutant. Specifically, the S84N mutant shows a modest 2.5:1 preference in L-SOS over L-serine and a 7:1 preference of L-THA over L-serine for β -elimination (Table 2). Although unable to catalyze L-serine racemization, this mutant is also the most fit mutant in catalyzing L-serine elimination, displaying ~75% of the catalytic efficiency of the native enzyme.

The S84A mutant also loses L-Ser racemization completely and has attenuated L-Ser elimination activity (>6-fold drop in catalytic efficiency, almost equally due to k_{cat} and K_m effects), and although this mutant displays quite respectable k_{cat} values for the charged substrates, it pays a significant penalty in K_m for L-THA (34-fold).

Whereas the alterations in substrate profile seen with the S84D mutant were dramatic, the S84T mutant displayed a more nuanced change in substrate preference that is also useful in considering the hSR mechanism. Thus, S84T-hSR retains the native hSR preference for charged substrates, but that preference is now selective for L-SOS over L-THA. In other words, the β -elimination of L-THA is now favored only 35-fold *versus* the β -elimination of L-serine compared with the original 100-fold, whereas L-SOS is now eliminated some 350-fold more efficiently than L-serine. Expressed differently, but perhaps more succinctly, whereas the k_{cat} for L-SOS elimination is unchanged for the S84T mutant, the corresponding k_{cat} values for L-serine and L-THA elimination suffer a 4–5-fold penalty with this subtle mutation in *re*-face base structure. Thus, the S84T-hSR is able to discriminate between these two charged substrates, showing an order of magnitude higher catalytic efficiency for processing L-SOS *versus* L-THA, whereas no such preference is seen in wild-type enzyme.

Molecular dynamics/docking studies

These kinetic results inspired us to employ molecular dynamics simulation and substrate docking experiments with

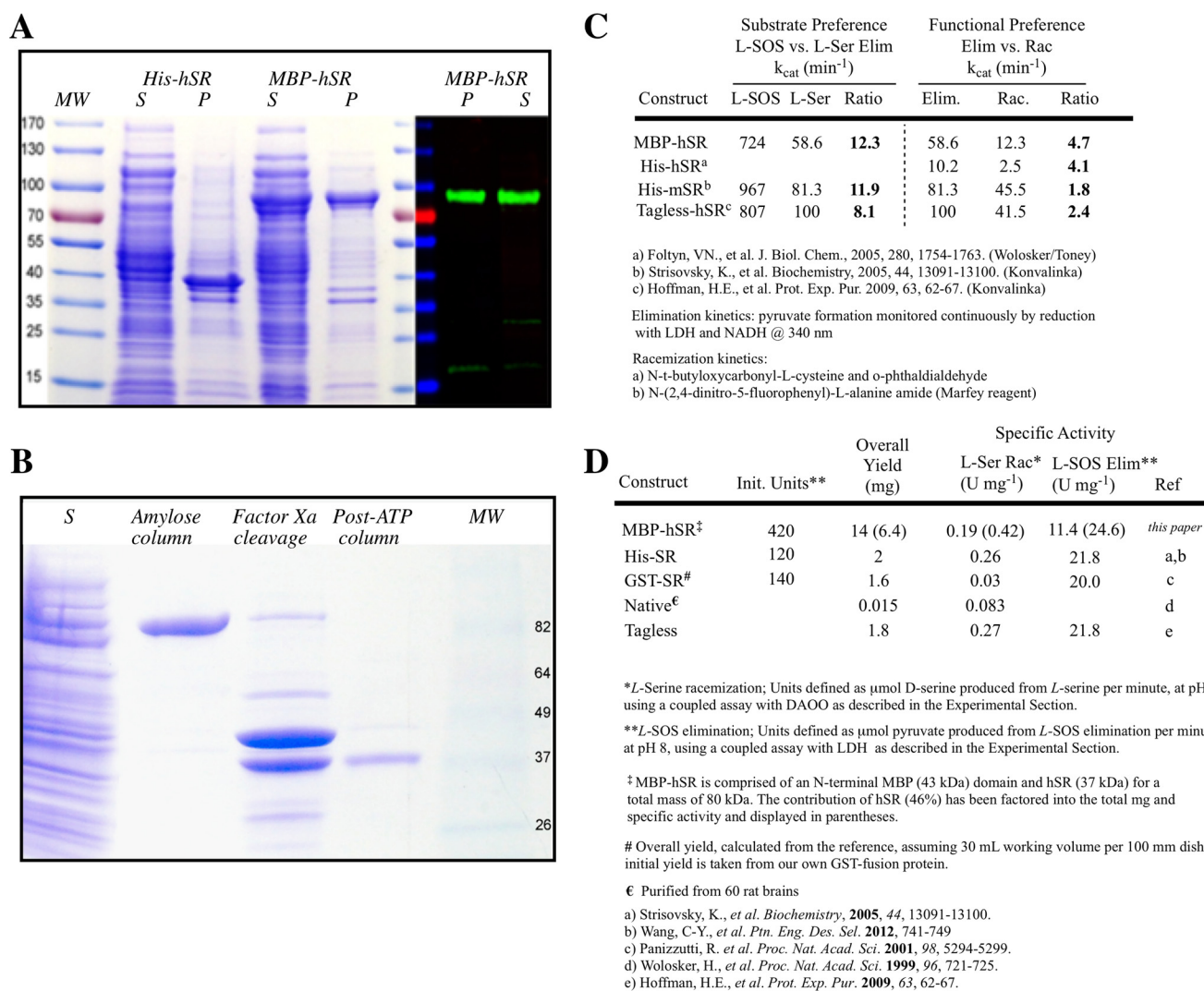


Figure 5. Use of the MBP-hSR construct for expression and purification. A, purification gel comparison of the MBP-hSR and His₆-hSR constructs by Coomassie Blue staining and Western blotting. B, removal of the MBP tag. C, comparison of kinetic properties of the MBP-hSR with hSR values reported in the literature (40, 43, 57). D, comparison of MBP-hSR specific activity with that of other reported SR constructs (1, 43, 57, 60, 75).

the goal of potentially shedding light on the origins of these observations on substrate preference. See “Experimental procedures” for details on how the homology model was constructed and how the molecular dockings were carried out and analyzed. All three elimination substrates were docked to both wild-type hSR and the S84D mutant. Displayed in Fig. 6 is the structure of L-THA-derived external aldimine docked into the wild-type hSR active site. The β -carboxylate of the substrate is engaged in a salt bridge interaction with Arg¹³⁵. This is somewhat reminiscent of the published crystal structures of SR with bound malonate inhibitor, wherein one carboxylate of the malonate appears to be similarly engaged (39). However, unlike L-THA, malonate is, of course, not covalently engaged with the PLP cofactor at all.

When L-THA is docked into the active site of the S84D-hSR mutant, the results are much different. What we observe is Arg¹³⁵ now interacting with the newly mutated S84D. The new *re-face* base, Asp⁸⁴, has “hijacked” Arg¹³⁵, forming an intramolecular salt bridge, which effectively renders Arg¹³⁵ unavailable to assist with the binding and proper positioning of the charged

substrates, L-SOS and L-THA. As can be seen in Fig. 6, this leads to a significant distortion of the substrate in the active site. This is illustrated with an overlaid image of the two structures (WT in lavender and S84D mutant in green).

The wild-type hSR enzyme is represented by purple residues, whereas the S84D mutant is shown in green. The external aldimine formed between L-THA and PLP is shown in silver. The interaction between the substrate and wild-type Arg¹³⁵ is again displayed. However, when Ser⁸⁴ is converted to Asp⁸⁴ (green), it becomes quite apparent how far Arg¹³⁵ has now migrated in the active site. This key active site residue no longer engages the substrate; rather, it interacts solely with Asp⁸⁴.

How might this simple mutation lead to the kinetic differences that we observe for the three hSR β -elimination substrates? During the course of the docking studies, we noticed a large change in substrate orientation within the active site. These orientations were examined closely with an eye toward how PLP enzymes operate normally, according to the Dunathan hypothesis (62). Looking more closely at the docking results for the external aldimine complexes of all three sub-

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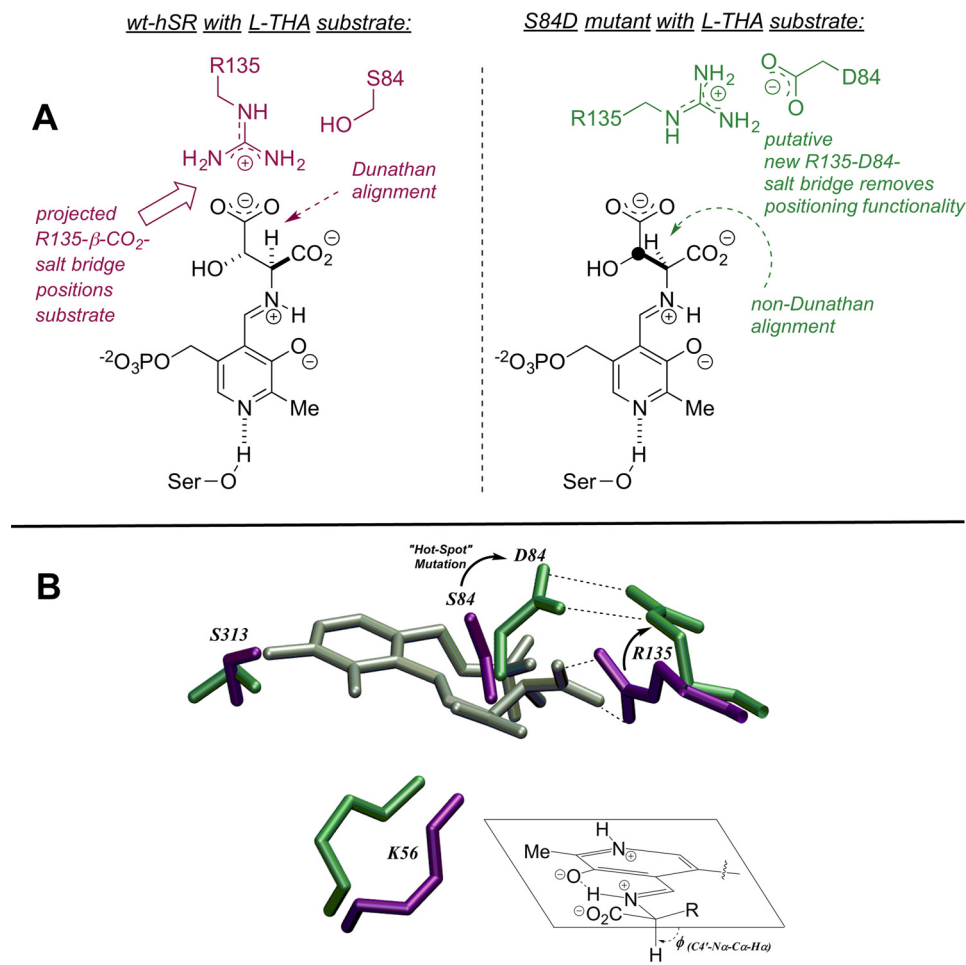


Figure 6. Molecular modeling results: WT versus S84D. Relaxation of the hSR homology model from Protein Data Bank entries 3L6B and 2ZPU (Fig. 2) with GROMACS version 4, followed by molecular docking of the L-THA-PLP-external aldimine (Autodock version 4) leads to the active site structures shown, both schematically (A) and in 3D (B). Arg¹³⁵ is projected to have an important role in L-THA substrate positioning in wild-type hSR (*lavender*). On the other hand, Arg¹³⁵ is engaged in a new salt bridge with Asp₈₄ (*green*) in the S84D mutant, making it unavailable for L-THA or L-SOS substrate positioning.

strates, a large difference was observed in the behavior of L-serine compared with that of its charged counterparts.

In Fig. 7 (*left column*), a typical example of the preferred docking orientation of each of the three substrates in the wild-type SR active site is shown. For all three cases, the substrate is aligned with a proper Dunathan orientation of the α -C–H bond to be broken. In other words, for all three cases, the α -C–H is nearly parallel with the π -system of the cofactor imine. In this case, Arg¹³⁵ is not only important for substrate binding/recognition; it appears to also be vital to substrate positioning for catalysis.

For the mutant S84D, we no longer see proper orientation. Whereas L-serine remains relatively unchanged in position, L-SOS and L-THA exhibit a substantial rotation about the key C_{4'}-N-C α -H dihedral angle. This non-Dunathan orientation is consistent with the dramatic decrease in β -elimination activity for these substrates with the S84D mutant, in good agreement with the experimental kinetic results. This apparent importance of Arg¹³⁵ raises interesting questions about the evolutionary history of serine racemase, as will be discussed below.

In an effort to understand the intermediate substrate preferences of the S84N mutant, molecular docking was also under-

taken here. In nearly all docked structures, the new asparagine 84 residue was seen to interact with aspartate 238, probably through hydrogen bonding. Perhaps because residue 84 is no longer able to contribute to the α -carboxylate binding site, one sees a major cluster of conformers in which Arg¹³⁵ is now engaged with the α -carboxylate.

Indeed with the charged substrates, L-SOS and L-THA, whereas in the WT enzyme these substrates appear to be “locked” into position for catalysis through electrostatic pairing with Arg¹³⁵ (Figs. 6 and 7), in the S84N mutant, molecular docking identifies two nearly equally populated clusters of conformers, the aforementioned Arg¹³⁵- α -carboxylate cluster (Fig. 8, *left*) and a second cluster of conformers in which the Arg¹³⁵-side chain interaction is retained (Fig. 8, *right*). Inspection of the C_{4'}-N-C α -H dihedral angle for the individual members of each of these major clusters gives a dihedral angle range of 140–190° for the Arg¹³⁵- α -carboxylate cluster and of 80–105° for the Arg¹³⁵-side chain cluster. The latter dihedral angle window appears to be in the stereoelectronically allowed range for Dunathan-compliant α -deprotonation. Representative examples of members of each cluster for both the bound L-SOS-external aldimine (*top half*) and the bound L-THA-external aldimine are depicted in Fig. 8. Overall, this Arg¹³⁵-toggle model for

Table 1
Kinetic inventory of hSR variants across the major enzymatic reaction manifolds

L-Ser \longrightarrow Pyruvate				
Variant	k_{cat} (min^{-1})	K_{m} (mM)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{mM}^{-1}\text{min}^{-1}$)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{M}^{-1}\text{s}^{-1}$)
WT	58.6 \pm 1.7	10.0 \pm 0.6	5.9	98.3
S84A	25.7 \pm 0.5	27.6 \pm 0.6	0.93	15.5
S84D	12.7 \pm 0.3	12.6 \pm 1.1	1.0	16.7
S84N	49.1 \pm 1.0	11.2 \pm 1.0	4.38	73.1
S84T	13.7 \pm 0.2	20.9 \pm 0.8	0.65	10.8

L-Ser \longrightarrow D-Ser				
Variant	k_{cat} (min^{-1})	K_{m} (mM)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{mM}^{-1}\text{min}^{-1}$)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{M}^{-1}\text{s}^{-1}$)
WT	12.3 \pm 0.42	7.8 \pm 0.7	1.57	26.1
S84A	-	-	-	-
S84D	-	-	-	-
S84N	-	-	-	-
S84T	3.8 \pm 0.24	29.2 \pm 1.1	0.13	2.16

L-SOS \longrightarrow Pyruvate				
Variant	k_{cat} (min^{-1})	K_{m} (mM)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{mM}^{-1}\text{min}^{-1}$)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{M}^{-1}\text{s}^{-1}$)
WT	724 \pm 9.8	1.2 \pm 0.1	603	1.0x10 ⁴
S84A	520 \pm 12	7.7 \pm 0.7	67.4	1.1x10 ³
S84D	9.8 \pm 1.2	122 \pm 20	0.08	1.3
S84N	72.1 \pm 3.0	6.5 \pm 1.2	11.1	184
S84T	700 \pm 19	2.9 \pm 0.3	241	4.0x10 ³

L-THA \longrightarrow Oxaloacetate				
Variant	k_{cat} (min^{-1})	K_{m} (mM)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{mM}^{-1}\text{min}^{-1}$)	$k_{\text{cat}}/K_{\text{m}}$ ($\text{M}^{-1}\text{s}^{-1}$)
WT	1400 \pm 16	2.5 \pm 0.3	560	9.3x10 ³
S84A	843 \pm 15	85 \pm 2.5	9.9	165
S84D	0.85 \pm 0.2	43 \pm 2.3	0.02	0.33
S84N	181 \pm 6	5.8 \pm 1.8	31.2	521
S84T	422 \pm 9.7	13 \pm 0.8	33	550

Table 2
Relative catalytic efficiencies of substrates as a function of hSR variant

Variant	$\frac{k_{\text{cat}}/K_{\text{m}}(\text{Sub.})}{k_{\text{cat}}/K_{\text{m}}(\text{L-Ser})}$		
	L-SOS	L-THA	L-Ser vs L-SOS vs L-THA
WT	100:1	93:1	1:100:93
S84A	1:12	1:50	1:71:34
S84D	1:12	1:50	50:4:1
S84N	2.5:1	7:1	1:2.5:7
S84T	370:1	50:1	1:370:50

substrate binding for the S84N mutant is consistent with the modest preference for these charged substrates displayed.

Overviewing this set of hSR mutants, then, in light of these docking results, one sees that WT enzyme appears to permit a sort of three-point binding interaction with Ser⁸³, Ser⁸⁴ (α -carboxylate), and Arg¹³⁵ (charged side chains). Mutation to S84N appears to remove residue Asn⁸⁴ from the binding pocket as it becomes engaged with Asp²³⁸. This appears to drive an Arg¹³⁵-toggle in binding both anionic groups in the charged substrates. Mutation to S84D removes both residue Asp⁸⁴ and Arg¹³⁵ because they are predicted to combine to form a salt bridge. Whereas these *in silico* models are consistent with the relaxed preference for L-SOS and L-THA seen in the S84N mutant and with the dramatic reversal of substrate preference seen in the S84D mutant, they remain to be tested by structural biology studies in the future.

Probing the active site with inhibitors

That said, we next set out to undertake complementary experiments to provide additional data with which to evaluate this interesting hSR binding model. Namely, given the importance ascribed to Arg¹³⁵, particularly in binding and positioning the charged substrates, L-SOS and L-THA, in this model, it seemed prudent to challenge this array of hSR active sites with a battery of inhibitors. A set of four inhibitor candidates was chosen: (i) malonate, (ii) L-aspartate β -hydroxamate (L-ABH), (iii) L-erythro- β -hydroxyaspartate (L-EHA), and (iv) aminooxyacetate (AOAA). Three of these inhibitors bear anionic side chains that might be expected to engage Arg¹³⁵, particularly in light of the substrate binding model being put forward here. The β -hydroxamate in L-ABH might also be expected to interact with Arg¹³⁵, although probably not as strongly. Only for malonate is crystal structure information available (39), and indeed such an interaction with Arg¹³⁵ is seen (Fig. 4).

All experiments were conducted in competition with L-serine, the native substrate. The first two hSR inhibitors showed dramatically different behavior with WT-hSR *versus* the S84N and S84D mutants (Fig. 9 and Table 3). Malonate displayed competitive inhibition with $K_i = 65 \pm 3.2 \mu\text{M}$ (reported $K_i = 27\text{--}71 \mu\text{M}$ (38, 43, 57)). L-ABH also inhibited WT-hSR competitively, with $K_i = 155 \pm 11.2 \mu\text{M}$ (reported K_i of 93 μM (63)) but showed no inhibition of either the S84N or the S84D mutant. As

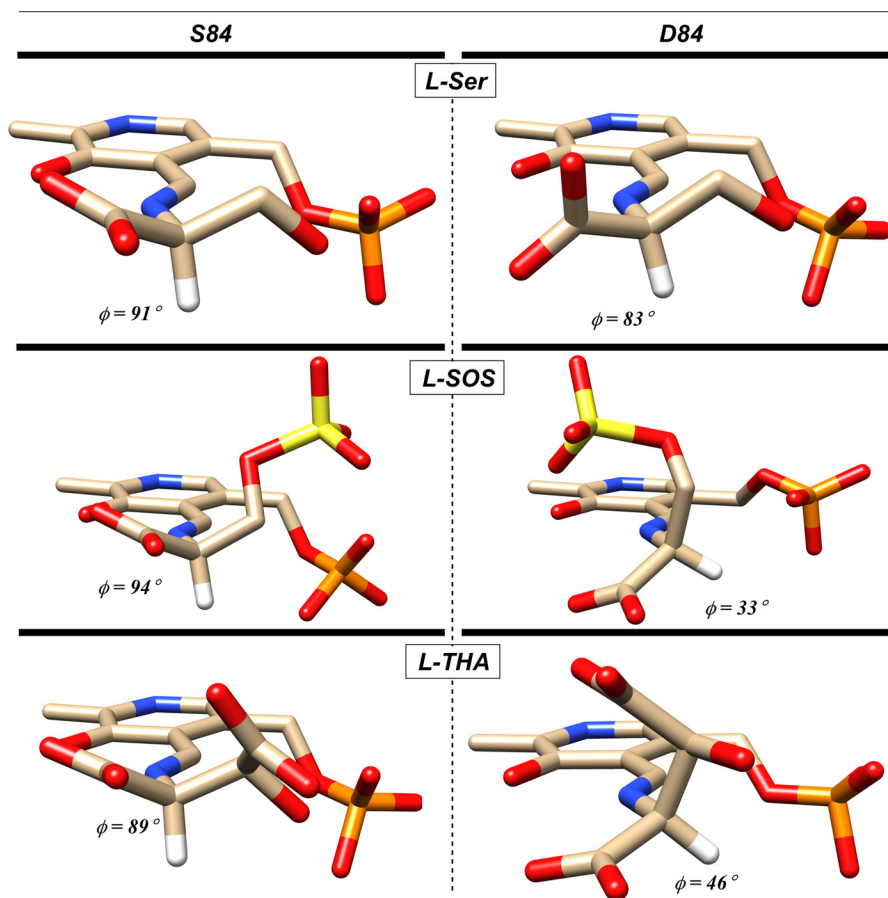


Figure 7. Molecular modeling to examine stereoelectronics in external aldimines for WT- and S84D-hSR. Molecular docking (Autodock version 4) results imply that whereas the L-Ser, L-SOS, and L-THA β -elimination substrates show proper Dunathan alignment in their respective external aldimines for WT-hSR, this alignment is significantly altered in the S84D mutant.

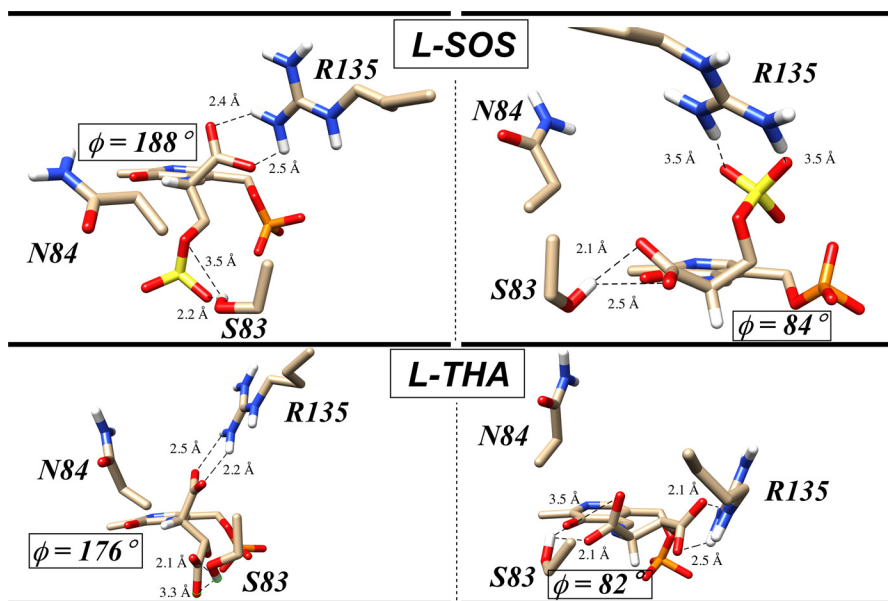


Figure 8. Molecular modeling suggests two types of bound conformations for L-SOS and L-THA in S84N-hSR. Molecular docking (Autodock version 4) results show two clusters of bound external aldimine conformers for the L-SOS and L-THA substrates with the S84N mutant. The conformers on the *left* exhibit an Arg¹³⁵- α -carboxylate salt bridge and are non-Dunathan-aligned, whereas in conformers on the *right*, Arg¹³⁵ is engaged with the charged side chain, leading to proper alignment for α -deprotonation.

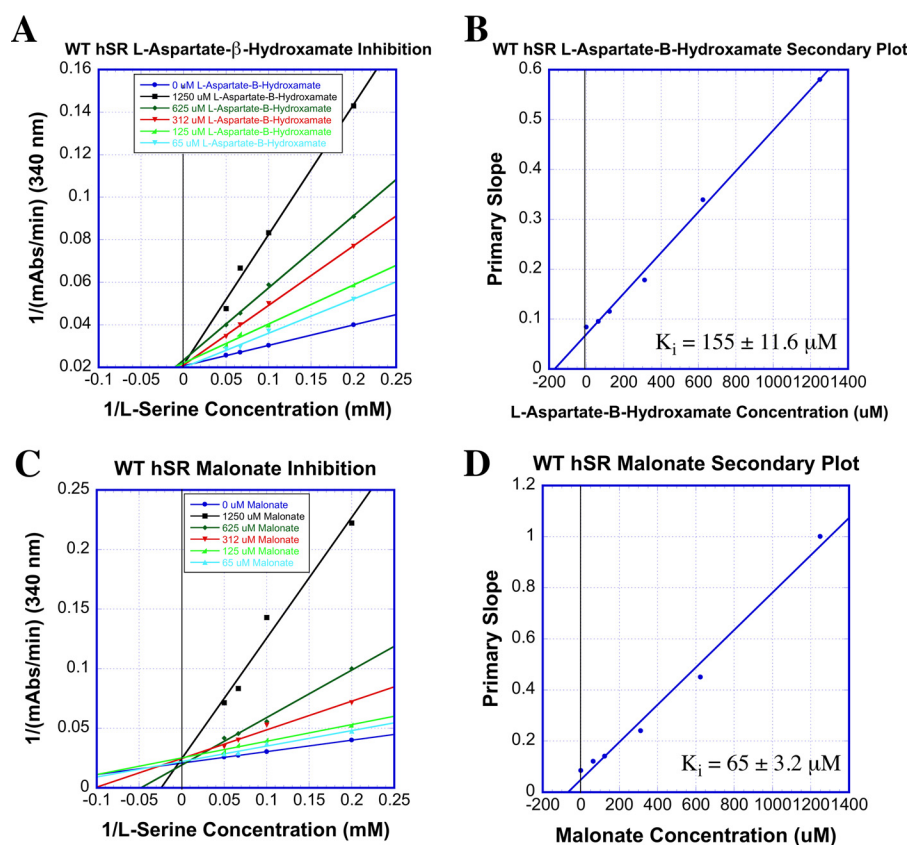


Figure 9. Competitive inhibition of WT-hSR with malonate and L-ABH. A, Lineweaver–Burk plot for WT-hSR inhibition with malonate; B, secondary plot for these data. C, Lineweaver–Burk plot for WT-hSR inhibition with L-ABH; D, secondary plot.

Table 3
Inhibitor profile across the hSR mutant array

Inhibitor Candidate	Structure	Inhibition Behavior - SR Panel		
		wt-hSR	S84N-hSR	S84D-hSR
L-EHA (L-erythro- β -hydroxy-Asp)		$K_i = 31 \pm 1.5 \mu\text{M}$ (competitive)	$K_i = 1.5 \pm 0.1 \text{ mM}$ (competitive)	none (up to 20 mM)
Malonate		$K_i = 65 \pm 3.2 \mu\text{M}$ (competitive)	none (up to 20 mM)	none (up to 20 mM)
L-ABH (L-Asp- β -hydroxamate)		$K_i = 155 \pm 11.6 \mu\text{M}$ (competitive)	none (up to 20 mM)	none (up to 20 mM)
AOAA (aminoxyacetate)		$K_i = 1 \pm 0.1 \mu\text{M}$ (competitive)	$K_i = 7 \pm 0.4 \mu\text{M}$ (competitive)	$K_i = 6 \pm 0.4 \mu\text{M}$ (competitive)

a positive control, the oxime-forming global PLP-dependent enzyme inactivator, AOAA (1), was tested and displayed potent inactivation of WT-hSR as well as the S84N and S84D mutants (Table 3).

L-EHA exhibited competitive inhibition kinetics with $K_i = 31 \pm 1.5 \mu\text{M}$ for WT-hSR in our hands (reported K_i of 11–43 μM (43, 57)). Upon mutation of the active-site Ser⁸⁴ to Asn, however, a pronounced 50-fold decrease in inhibition potency was observed as K_i increased to 1.5 mM. Even more extreme, the S84D mutant showed no inhibition with L-EHA at concentrations up to 20 mM. (Fig. 10 and Table 3).

Discussion

The effectiveness of all four inhibitors with native hSR and the inability of all but the universal PLP inactivator, AOAA, to inhibit the S84D mutant is a striking contrast and is consistent with the Arg¹³⁵-charged side chain binding model that emerged from the earlier substrate scan/molecular modeling. Much as in the β -elimination substrate studies, the S84N mutant again displayed behavior that is intermediate between WT-hSR and S84D-hSR in the inhibition studies. Namely, in contrast to the S84D mutant, the S84N mutant is inhibited by L-EHA ($K_i = 1.5 \pm 0.1 \text{ mM}$) but considerably less well than WT-hSR ($K_i = 31 \pm 1.5 \mu\text{M}$). These results are in line with the details of the molecular modeling that emerged earlier whereby it was predicted that Arg¹³⁵ would be fully available for side chain carboxylate binding in WT-hSR (Fig. 6), partially available in the S84N mutant (Fig. 8) and unavailable in the S84D mutant (Fig. 6). Indeed, molecular modeling of the putative L-EHA-external aldimine with both WT-hSR and S84N hSR indicates that these active sites are also capable of engaging the β -carboxylate of L-EHA in a salt bridge with Arg¹³⁵ (Fig. 11), consistent with the ability of this compound to inhibit both enzymes.

This study also raises interesting questions from the observation of the S84T mutant, particularly its ability to retain all hSR functions and yet to discriminate between the L-SOS (preferred) and L-THA substrates. Recent investigations into the mechanisms of the related PLP-dependent β -eliminases *Drosophila* CBS (64) and tryptophan synthase (65) may provide

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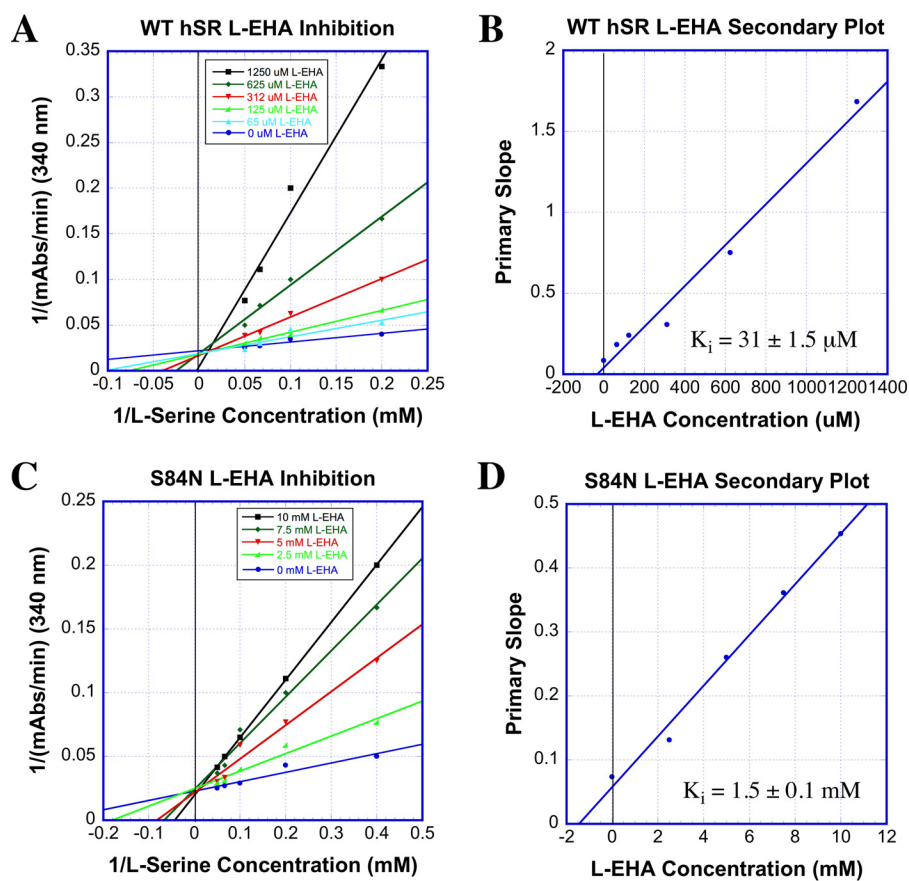


Figure 10. L-EHA inhibition steady-state kinetic results for WT- and S84N-hSR. A, Lineweaver–Burk plot for WT-hSR inhibition with L-EHA; B, secondary plot for these data. C, Lineweaver–Burk plot for S84N-hSR inhibition with L-EHA; D, secondary plot.

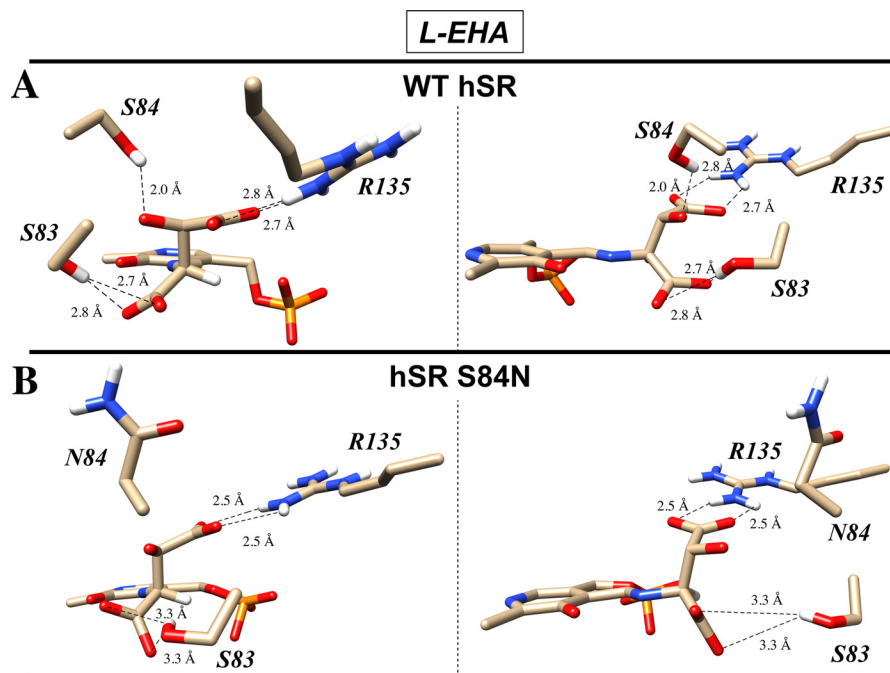


Figure 11. L-EHA inhibition molecular modeling. Shown are docking results (Autodock version 4) for the inhibitor *L*-erythro- β -hydroxyaspartate (A) in the wild-type hSR active site and in the S84N-hSR active site (B).

guidance here. Both of these enzymes also catalyze the β -elimination of PLP-aldimine-bound *L*-serine along the normal reaction coordinate, for LL-cystathionine and *L*-tryptophan bio-

synthesis, respectively. In the *Drosophila* CBS study (64), the authors claim to observe a carbanionic intermediate that is generated upon *L*-serine deprotonation. It is argued that this spe-

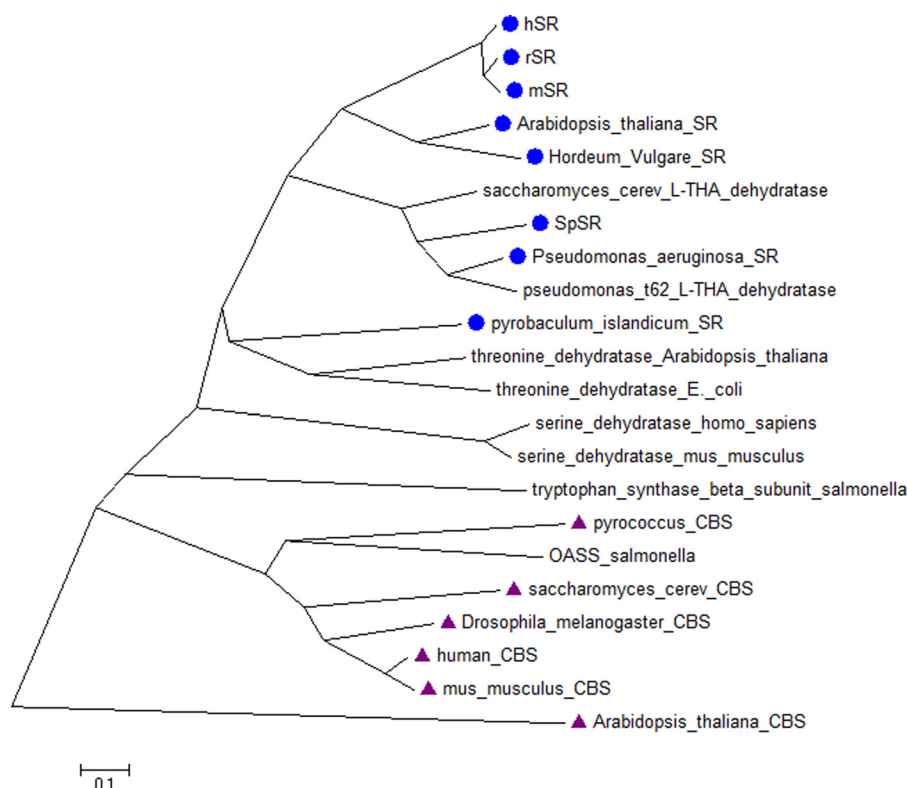


Figure 12. Phylogenetic tree of various fold type II eliminases. Enzymes annotated as SR are marked with blue dots; enzymes annotated as having CBS activity are given lavender triangles.

cies is an incompletely delocalized, PLP-stabilized α -carbanion, as opposed to a fully delocalized quinonoid intermediate. The authors argue that both the *si*-face lysine ammonium ion and a *re*-face serine residue are of central importance in stabilizing this key mechanistic intermediate.

Mueller and Dunn (65) have recently described a similar carbanionic intermediate for the archetypical β -replacement enzyme, tryptophan synthase, utilizing a new biophysical method that combines solid-state NMR, X-ray crystallography, and computational chemistry. Here too, the active site lysine ammonium ion is thought to stabilize the negative charge in this intermediate. It is, of course, possible that a similar carbanionic intermediate forms along the reaction coordinate for hSR. It may be that replacement of a β -H with a β -methyl group (S84T) results in a subtle repositioning of this residue, rendering it less efficient at stabilizing the developing negative charge in the enzyme-bound substrate aldimine upon α -deprotonation.

Indeed, it might well be the case that a stepwise mechanism involving initial rate-limiting substrate deprotonation takes place for the L-serine and L-THA hSR substrates, but not for L-SOS. The latter substrate, by virtue of having a good leaving group in sulfate, could presumably undergo a concerted β -elimination without the need to stabilize a discreet α -carbanionic PLP-substrate intermediate. This explanation is consistent with the observation that the hSR S84T mutant more effectively catalyzes the β -elimination of L-SOS as compared with that of L-THA, in contrast to WT-hSR. However, this is still quite speculative at this juncture. The nature of the PLP-centered intermediate in hSR catalysis has remained elusive

heretofore, so future experiments will be needed to evaluate this notion. It may be possible to shed light on this hypothesis, at least indirectly, through the sort of Mueller/Dunn solid-state NMR/X-ray studies described above or through successful co-crystallization of hSR with known inhibitor L-EHA.

Evolutionary considerations

To examine the potential evolutionary implications of the favorable L-THA elimination kinetics observed here, we decided to construct a phylogenetic tree of various type II β -eliminases acting on L-serine (Fig. 12) (66). The blue dots represent enzymes known to racemize serine. What we observe is that only recently do we see serine racemase activity. Furthermore, an apparent ancestor is shared with enzymes that are annotated as L-THA dehydratases from *Saccharomyces cerevisiae* (67) and *Pseudomonas* sp. T62 (68). Very recently, such activity has been observed in a *Caenorhabditis elegans* enzyme as well (69). Looking more closely at the sequences of a range of proteins in this broad β -eliminase family, we observe the following trends. First, Ser⁸⁴ is conserved in serine racemases and L-THA dehydratases. Perhaps most striking, Arg¹³⁵ is also conserved among these examples while being absent in all other type II dehydratases (Fig. 13). These results are consistent with the substrate preferences (*i.e.* L-SOS and L-THA as preferred β -elimination substrates over L-Ser (Tables 1 and 2)) observed for the wild-type serine racemase. This also suggests that perhaps serine racemase is still early on in its evolution of function (see Fig. 12).

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	84	135
hSR	HSSGNHG	DESRENV
mSR	HSSGNHG	DESREKV
rSR	HSSGNHG	DESRENV
atSR	HSSGNHA	MSSREEI
spSR	FSSGNHA	KDDRREKM
hvSR	HSSGNHA	MESRESI
paSR	FSSGNHA	TEDREQI
piSR	ASSGNHA	GESYYEA
sTHADH	FSSGNHA	TEDREQI
pTHADH	FSSGNHA	TEDREQI
hSDH	-SAGNAG	LDEAFEL
mSDH	-SAGNAG	LDEAIQV
atTDH	SSAGNHA	YDQAQAH
ecTDH	ASAGNHA	FDEAKAK
sOASS	-TNGNTG	KGMK121
sTrpS	-GAGQHG	SATLKDA
hCBS	-TSGNTG	ARFDSPE
mCBS	-TSGNTG	ARFDSPE

Figure 13. Alignment of fold type II enzymes. Arg¹³⁵ is conserved among enzymes showing L-THA eliminase activity consistent with the functional role of this residue posited herein. *h*, human; *m*, mouse; *r*, rat; *at*, *Arabidopsis thaliana*; *sp*, *S. pombe*; *hv*, *Hordeum vulgare*; *pa*, *Pseudomonas aeruginosa*; *pi*, *Pyrobaculum islandicum*; *sc*, *S. cerevisiae*; *p*, *Pseudomonas T62*; *ec*, *E. coli*; *s*, *Salmonella*.

Conclusions

Since the discovery of D-serine as a potent co-agonist of the NMDA receptor, there has been growing interest in the enzyme responsible for its biosynthesis, serine racemase. Its implication in various disease states has further raised interest in developing selective inhibitors (ischemic stroke and potentially Alzheimer's disease and ALS) or stimulators (schizophrenia) of this enzyme. A new expression construct for hSR is reported here, namely the MBP-hSR fusion protein, that yields higher titers of soluble enzyme and serves as an excellent platform to study active-site mutants. Characterization of these mutants has led to a new understanding of important residues in the mechanism of serine racemase. Specifically, these complementary "enzyme/substrate mutation" studies focusing on position 84 in the enzyme and the three principal hSR β -elimination substrates have raised mechanistic hypotheses for hSR function. Studies with the hSR S84D and S84N mutants suggest an important mechanistic role for Arg¹³⁵ in substrate positioning for charged β -elimination substrates that is consistent with the Dunathan hypothesis. Indeed, these observations, along with sequence alignment studies point to Arg¹³⁵ as a key residue that probably was selected for in this β -eliminase family to optimize the β -elimination of L-THA (conserved in all annotated L-THA dehydratases).

Phylogenetic analysis supports the notion that Arg¹³⁵ is historically important for L-THA eliminase activity. But where would organisms experience evolutionary selection pressure to evolve L-THA eliminase activity? Recently, Katane *et al.* (69) have speculated that *C. elegans* may have evolved its T01H8.2 protein, an apparent L-THA dehydratase, as an important

defense mechanism. Namely, microorganisms such as *Athrinium phaeospermum* and *Streptomyces* are known to produce L-THA, and this amino acid is known to inhibit L-glutamate transport. The thinking is that *C. elegans* might well feed on such microorganisms and take advantage of this evolved L-THA dehydratase activity to detoxify ingested L-THA, and whereas this may be the evolutionary origin of residue Arg¹³⁵ in hSR, it may well be possible to leverage this vestigial ligand-binding ability in future hSR inhibitor design studies.

Experimental procedures

Recombinant His-hSR

Human serine racemase cDNA was purchased from Origene and amplified using the following primers (IDT): 5'-CCT TCT TGC TAG CTG TGC TCA GTA TTG CAT C-3' (forward) and 5'-CAC GCG CTC GAG AAT TCC CAC CAT TTC-3' (reverse) for the NheI and XhoI restriction sites, respectively. Gene Runner software was used to design the primers, and the 10198-0118 PCR Reagent System (Invitrogen) was utilized. The cDNA was inserted into the pGEM-T cloning vector (Promega) and used to transform DH5 α -derived *E. coli* made competent by the calcium chloride method. Plasmids from overnight cultures were extracted using a plasmid miniprep spin kit (Qiagen) and verified by 1% agarose gel as well as sequencing with T7 and T7term primers. After double digestion, the fragment was inserted into the pET-28c plasmid. BL21(DE3)-pLysS *E. coli* was used for expression.

Starter cultures of 3-ml volume, grown overnight, were used to inoculate 1 liter of Luria-Bertani broth (Difco), supplemented with an additional 5 g/liter yeast extract. Cells were grown at 37 °C and 250 rpm. When the cells reached an A_{600} of 0.8, the temperature was reduced to 25 °C, and the cells were induced with 0.1–0.5 mM isopropyl 1-thio- β -D-galactopyranoside for 18 h at 300 rpm. Cells were pelleted by centrifugation for 15 min at 10,000 \times g and stored at –80 °C.

Purification of His-hSR

Cells were resuspended in assay Buffer A (200 mM TEA, 150 mM KCl, 10 mM DTT, 5 mM MgCl₂, 2.5 mM ATP, and 50 μ M PLP, pH 8.0) at a volume of 1 ml/g of wet cell mass. Cells were disrupted by sonication on ice for 5 cycles (1 min on/1 min off) and centrifuged for 15 min at 15,000 \times g. The supernatant contained modest levels of hSR and no visible SDS-PAGE band, whereas the pellet, when solubilized in the same volume of Buffer B (10 mM Tris, 100 mM sodium phosphate, 8 M urea, pH 7) as the crude supernatant, exhibited a very intense band at ~37 kDa. To verify that this band was insoluble His-tagged hSR protein, the solubilized pellet fraction was applied to a nickel-nitrilotriacetic acid column. The protein did bind and was successfully eluted, with decreasing pH, yielding a single band on SDS-PAGE that matched the expected molecular weight for hSR.

Recombinant GST-hSR

Human serine racemase cDNA was purchased from Origene (catalog no. TC11289) and amplified using the primers (IDT) 5'-CGT TGC GGA TCC ATG TGT GCT CAG TAT TGC-3'

(forward) and 5'-CAC GCG CTC GAG AAT TCC CAC CAT TTC-3' (reverse) for restriction sites BamHI and XhoI, respectively. The cDNA was double-digested and inserted into the pGEX-4T1 expression vector (GE Healthcare) and used to transform DH5 α -derived *E. coli* made competent by the calcium chloride method. Plasmids from overnight cultures were extracted using a plasmid miniprep spin kit (Qiagen) and used to transform BL21(DE3)pLysS *E. coli* for expression.

Purification of GST-hSR

Cells were resuspended in assay Buffer A at a volume of 1 ml/g of wet cell mass. Cells were disrupted by sonication on ice for five cycles (1 min on/1 min off) and centrifuged for 15 min at 15,000 \times *g*. The supernatant displayed more units of activity per liter of culture compared with the His-hSR construct but did not yield a visible band by SDS-PAGE. When the pellet was solubilized in Buffer B, there did exist a strong band at the expected molecular weight of the GST-hSR construct. SDS-PAGE indicated that most of the expressed protein was relegated to the insoluble pellet, so further purification was not attempted.

Recombinant MBP-hSR

The protein was designed to bear the N-terminal MBP tag present in the pMAL-c2X (New England Biolabs) vector. The following primers were employed in this system (IDT): 5'-CGT TGC GGA TCC ATG TGT GCT CAG TAT TGC-3' (forward) and 5'-CAC CTA GTC GAC AAT TCC CAC CAT TTC C-3' (reverse), for restriction sites BamHI and Sall, respectively. The respective hSR PCR products were ligated into the respective vectors and used to transform our DH5 α -competent cells. The amplified plasmids were then analyzed and used to transform BL21(DE3)pLysS *E. coli* strain for expression.

Purification of hSR

Cells were resuspended in assay Buffer A at a volume of 1 ml/g of wet cell mass. Cells were disrupted by sonication and collected by centrifugation (15,000 \times *g*). The supernatant was mixed with prebalanced amylose resin (New England Biolabs) and shaken at 4 $^{\circ}$ C for 1 h. The slurry was centrifuged 15 min at 5000 \times *g* and decanted. The resin was applied to a column of the same resin, washed with 3 column volumes of Buffer C (50 mM Tris, 150 mM KCl, 10 mM β -mercaptoethanol, 15 μ M PLP, pH 8.0), and eluted with a linear gradient (0–10 mM) of maltose. MBP-hSR was observed to elute at 3–5 mM maltose. Fractions were pooled by activity and UV₂₈₀ absorbance. Pooled fractions were then concentrated (Amicon cell, 30-kDa membrane) to a final concentration of 20 mg/ml further clarified by centrifugation (15 min, 15,000 \times *g*, 4 $^{\circ}$ C). Protein was flash-frozen in 50% (v/v) and stored at –80 $^{\circ}$ C.

To obtain tag-free hSR, the MBP tag of pMAL-c2X was cleaved by factor Xa (New England Biolabs), leaving a 6-mer (ISEFGS) N-terminal overhang immediately preceding the initial methionine of hSR. Digestion took place in Buffer D (20 mM Tris, 50 mM NaCl, 0.2 mM β -mercaptoethanol, and 10 μ M PLP). After overnight digestion, the solution was applied to a 3-ml column of C-8-linked ATP-agarose (Sigma), washed with 3

column volumes of Buffer D, and eluted with 2.5 mM ATP in Buffer D.

β -Elimination activity assay

β -Elimination of L-SOS to form pyruvate was conducted under the following assay conditions: 200 mM TEA, 150 mM KCl, 5 mM MgCl₂, 2.5 mM ATP, 50 μ M PLP, 0.15 units of lactate dehydrogenase, 0.24 mM NADH, and 10 mM L-serine-O-sulfate. Activity was monitored by observing the decrease in absorbance at 340 nm.

L-Serine-O-sulfate β -elimination

A Shimadzu UV-2101PC spectrophotometer equipped with a 6-cell changer was used to monitor the decrease in NADH concentration as pyruvate produced from L-serine-O-sulfate elimination was reduced by lactate dehydrogenase. Assay conditions were as follows: 37 $^{\circ}$ C, pH 8, 200 mM TEA, 150 mM KCl, 5 mM MgCl₂, 2.5 mM ATP, 50 μ M PLP, 0.24 mM NADH, and 0.15 units of lactate dehydrogenase. Each point represents the average of three experiments. The kinetic parameters for MBP-hSR (79.5 kDa) were estimated by least squares hyperbolic fitting to the Michaelis–Menten equation.

L-Serine elimination

L-Serine elimination activity was evaluated by the same assay described for L-SOS elimination, because both reactions give pyruvate as the hSR product.

L-threo- β -Hydroxyaspartate β -elimination

L-THA elimination was monitored by the formation of oxaloacetate. Oxaloacetate formation was measured by reduction with malate dehydrogenase and NADH under the same conditions described for L-SOS.

L-Serine racemization

To obtain racemization data, D-amino acid oxidase from porcine kidney (Sigma) was used to oxidize D-serine to 3-hydroxypyruvate and H₂O₂. The resulting H₂O₂ was reduced in the presence of horseradish peroxidase and Amplex Red to generate resorufin that can be monitored continuously at 570 nm. Racemization was also monitored via a time point assay by derivatizing the products with Marfey's reagent (1-fluoro-2,4-dinitrophenyl-5-L-alanine) and resolved using reversed-phase HPLC with glycine as an internal standard.

Inhibition studies

Wild-type hSR and the S84N and S84D mutants were probed for inhibition by malonate, L-EHA, L-ABH, and AOAA. Utilizing the L-serine elimination assay at various L-serine concentrations (5, 10, 15, and 20 mM) and a battery of inhibitor concentrations, *K_i* values were determined following standard steady-state kinetic analysis. Wild-type hSR was examined with all inhibitors (except AOAA) at inhibitor concentrations of 65, 125, 312, 625, and 1250 μ M. For the S84N mutant (aside from AOAA), inhibition was seen only with L-erythro- β -hydroxyaspartate upon incubation at 2.5, 5, 7.5, and 10 mM concentrations. The S84D mutant showed no inhibition up to 20 mM concentration with malonate, L-EHA, and L-ABH. The WT-

Human serine racemase structure/activity relationship studies

hSR and S84N and S84D mutants were all effectively inhibited by AOAA, which was tested at 1, 2, 3, 4, 5, and 10 μM . Activity was measured by observing the decrease in absorbance at 340 nm associated with the consumption of NADH in the coupled lactate dehydrogenase reaction converting pyruvate to L-lactate.

Oligomeric state determination

Native PAGE and gel filtration analysis were used to determine the oligomeric state of the MBP-hSR construct. For native gels, molecular weight markers from GE Healthcare (High MW) were used. Electrophoresis was run using 4% stacking, 9% resolving discontinuous polyacrylamide minigels at constant current (40 mA) under flow-cooling in a Hoeffer Mighty Small II apparatus. A calibration curve (R_f versus $\log M_r$) was constructed for the standards. The primary MBP-hSR band displayed an R_f value of 0.397, corresponding to an apparent molecular mass of 185 kDa, which corresponds to a dimer (2.3 monomeric units) based on the calculated monomeric molecular mass of 79.6 kDa. A similar analysis of the His₆-hSR construct by native PAGE showed an apparent molecular mass of 81.5 kDa, again corresponding to a dimeric structure (2.1 monomeric units). Gel filtration analysis employing a GE Healthcare S-200 High-Prep Sephacryl HS column with Bio-Logic DuoFlow software was utilized. A calibration curve was constructed using the GE Healthcare high molecular weight gel filtration calibration kit and revealed an apparent molecular mass of 247 Da for the new construct, suggestive of an oligomeric composition of 3.1 (possible dimer-tetramer equilibrium (58)).

Homology modeling

To construct a hSR homology model, the 340-amino acid human serine racemase protein sequence (NCBI accession number NP_068766) was BLASTed against the NCBI structural database (70). A multiple-sequence alignment was performed on hSR, 3L6B, 1WTC, and 2ZPU, using the ClustalW algorithm (71). This alignment forms the basis for construction the hSR model by MODELLER (72).

ATP, PLP, Ca^{2+} , and Mg^{2+} were copied into the homology model from the template structures. PLP is bound to all template structures in a similar orientation. Of these, PLP from 3L6B was chosen for the hSR model. 1WTC contains a bound ATP/ Mg^{2+} analogue that was copied into the hSR model and energy-minimized so as to represent bound ATP.

Molecular dynamics and docking experiments

All MD simulations were performed using the GROMACS software package (73). Before each run, the structure was solvated in a water box using the spc216 water model. Box dimensions were set so that the hSR model was no closer than 1 nm to the edge of the box. Simulations were performed using the GROMOS96 force field, periodic boundary conditions, standard temperature-coupling schemes, and the particle-mesh Ewald method for determining long-range electrostatics. Each MD run was preceded by a 1000-step steepest descent energy minimization and a 20-ps position-restrained MD simulation.

Each full MD simulation was performed for 2 ns at 300 K, using 2-fs time steps.

The first simulation contained Ca^{2+} , $\text{ATP}/\text{Mg}^{2+}$, and non-covalently bound PLP. This setup was chosen to allow the active-site lysine (Lys⁵⁶) to reorient itself to a position that would not obstruct external aldimine docking. Following the initial 2-ns MD simulation, the hSR structure was energy-minimized and prepared for docking. Autodock version 4 was used to dock the external aldimine of L-serine, L-serine-O-sulfate, L-threo- β -hydroxyaspartate, and L-erythro- β -hydroxyaspartate into the hSR active site (74).

Author contributions—D. B. B., D. L. N., G. A. A., M. L. B., and D. L. G. designed the experiments. D. L. N., G. A. A., M. L. B., and D. L. G. performed the experiments. D. B. B., D. L. N., G. A. A., M. L. B., and D. L. G. reviewed and analyzed the results, contributed to writing the manuscript, and approved the final version.

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