Negative Epistasis and Evolvability in TEM-1 β-Lactamase—The Thin Line between an Enzyme's Conformational Freedom and Disorder

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http://dx.doi.org/10.1016/j.jmb.2015.05.011

Edited by A. G. Palmer III

Abstract

Epistasis is a key factor in evolution since it determines which combinations of mutations provide adaptive solutions and which mutational pathways toward these solutions are accessible by natural selection. There is growing evidence for the pervasiveness of sign epistasis—a complete reversion of mutational effects, particularly in protein evolution—but its molecular basis remains poorly understood. We describe the structural basis of sign epistasis between G238S and R164S, two adaptive mutations in TEM-1 β-lactamase—an enzyme that endows antibiotics resistance. Separated by 10 Å, these mutations initiate two separate trajectories toward increased hydrolysis rates and resistance toward second and third-generation cephalosporins antibiotics. Both mutations allow the enzyme’s active site to adopt alternative conformations and accommodate the new antibiotics. By solving the corresponding set of crystal structures, we found that R164S causes local disorder whereas G238S induces discrete conformations. When combined, the mutations in 238 and 164 induce local disorder whereby nonproductive conformations that perturb the enzyme's catalytic preorganization dominate. Specifically, Asn170 that coordinates the deacylating water molecule is misaligned, in both the free form and the inhibitor-bound double mutant. This local disorder is not restored by stabilizing global suppressor mutations and thus leads to an evolutionary cul-de-sac. Conformational dynamism therefore underlines the reshaping potential of protein's structures and functions but also limits protein evolvability because of the fragility of the interactions networks that maintain protein structures.

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Epistasis, including sign epistasis, is pervasive [2,3]. The phenomenon itself is measured and relates to organismal fitness. It is clear, however, that its origins lie in antagonistic interactions between mutations, either in separate genes (or proteins) or within the same gene/protein. However, the molecular basis of epistasis remains poorly understood [10]. We have better understanding of positive epistasis at the single gene level, such as in deer mouse hemoglobin adapted to high altitude [15]. Typically, stabilizing mutations, which have no effect on protein fitness on their own, are revealed as beneficial when they compensate for new function mutations that are typically destabilizing [16]. Such stabilizing mutations can have local, specific compensatory effects [17,18], or global effects, when they can compensate for a whole range of destabilizing mutations [16] in non-contacting residues [19–21]. One case study of positive epistasis in the vertebrate glucocorticoid receptor revealed a mutation that was initially neutral, yet by reorienting an α-helix, it enabled the acceptance of an adaptive mutation that would be deleterious on its own [22]. Overall, it appears that stabilization of the protein’s configuration promotes its adaptive potential and underlines positive epistasis [21,22]. However, a detailed, structural understanding of negative sign epistasis is lacking: Why do certain combinations of mutations, each of which is beneficial individually, become deleterious when combined [23,24]?

Here, we explored a case of negative reciprocal sign epistasis between two adaptive mutations in TEM-1 β-lactamase, an enzyme present in numerous antibiotic-resistant bacteria. These studied mutations, G238S and R164S, are among the most common mutations in clinically isolated variants and therefore represent a highly relevant case study [25,26]. Wild-type TEM-1 confers resistance to natural penicillin antibiotics. However, in the clinic, strains with new resistance mutations are frequently isolated. This increased resistance is the outcome of adaptive evolution of TEM-1 for new, so-called second and third-generation cephaplosporins antibiotics. Adaptive evolution of TEM-1 has occurred in numerous parallel events and can be readily reproduced in the laboratory [25]. The increased catalytic activity of TEM-1 to second and third-generation cephaplosporins antibiotics therefore comprises a broadly accepted model for enzyme evolution [3,25,26]. Owing to the large sampling of TEM-1 sequences by natural and directed evolution, the absence of certain combinations of mutations likely indicates their negative epistatic interaction. This is the case with G238S and R164S, which were individually found in 91 and 51 different variants, respectively, both in laboratory and in clinical contexts [25]. However, their combination has only been observed in one clinical sample [27]. Indeed, the combination of G238S and R164S results in in vivo resistance levels to cefotaxime that are lower than each of the single mutants (4-fold lower than R164S, and 8-fold lower than G238S) and are similar to wild-type TEM-1 (Refs. [7] and [27]). Laboratory experiments attempting to evolve variants carrying the R164S mutation toward high cefotaxime activity indicated that R164S blocks the path to maximal cefotaxime degradation activity [7]. One mutation, E104K, improves R164S’s activity, but the activity of the R164S/E104K double mutant was still inferior to G238S/E104K [24]. The only traceable trajectory to higher cefotaxime resistance in these laboratory evolution experiments was through reversion of R164S and take over by G238S (Ref. [7]). Thus, R164S seems to comprise an evolutionary cul-de-sac [7]. However, the molecular basis underlying the limited adaptive potential of R164S and of the reciprocal sign epistasis with G238S remains unclear.

The strong, nonadditive interaction between R164S and G238S is surprising for two reasons: firstly, the mutation sites are not in direct contact (>10 Å apart), and secondly, combining these mutations does not further compromise TEM-1’s global configurational stability [7,16]. To determine the structural basis for this classical case of reciprocal sign epistasis, and to understand why R164S might lead to an evolutionary cul-de-sac, we solved the crystal structures of the R164S, G238S and double mutant R164S/G238S, in their free forms and bound to an inhibitor. Although these mutations are key first-step mutations in the adaptation of TEM-1 to second and third-generation cephaplosporins antibiotics, the structure of enzymes harboring these mutations with no additional active-site mutations has not been available thus far. The combination of free versus inhibitor bound and of variable temperature X-ray data allowed for increased sampling of the conformational ensemble of TEM-1, thus unraveling the structural basis of negative epistasis and evolvability in TEM-1.

Results

Mutations that confer TEM-1 with the ability to hydrolyze new antibiotics are located mostly on the loops surrounding the active site: the “Ω-loop” (residues 164–179, Ref. [28]), the “238-loop” (residues 238–242, Ref. [29]) and the “101-loop” (residues 101–111, Ref. [30]). The mutations studied here, G238S and R164S, enhance cefotaxime hydrolysis (third-generation cephaplosporins antibiotics) and reside on the 238-loop and the Ω-loop, respectively [7,29]. Mutation G238S increases cefotaxime resistance levels by ~16-fold: the MIC or the minimal inhibitory concentration increases from 0.06 μg/ml to 1 μg/ml (Table 1). Mutation R164S
Table 1. Kinetic and stability parameters and *E. coli* resistance levels of the TEM-1 variants

<table>
<thead>
<tr>
<th>Variant</th>
<th>Antibiotic</th>
<th>$K_M$ (μM)</th>
<th>$k_{cat}$ (s⁻¹)</th>
<th>$k_{cat}/K_M$ (M⁻¹·s⁻¹)</th>
<th>MIC (μg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>Amp</td>
<td>71 ± 9</td>
<td>1857 ± 27</td>
<td>26 ± 1.3 × 10⁶</td>
<td>4096</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>n.d.</td>
<td>n.d.</td>
<td>1 × 10³ ± 6</td>
<td>0.06</td>
</tr>
<tr>
<td>Stabilized v13</td>
<td>Amp</td>
<td>63 ± 13</td>
<td>1040 ± 21</td>
<td>16.5 ± 1.3 × 10⁶</td>
<td>n.m.</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>n.d.</td>
<td>n.d.</td>
<td>0.07 ± 10²</td>
<td>1</td>
</tr>
<tr>
<td>Wild type + G238S</td>
<td>Amp</td>
<td>26 ± 4</td>
<td>42 ± 0.4</td>
<td>1.4 ± 0.08 × 10⁶</td>
<td>1024</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>403 ± 146</td>
<td>50 ± 3</td>
<td>13 ± 4 × 10⁴</td>
<td>1</td>
</tr>
<tr>
<td>Stabilized v13 + G238S</td>
<td>Amp</td>
<td>18 ± 5</td>
<td>72 ± 0.5</td>
<td>4 ± 0.36 × 10⁶</td>
<td>n.m.</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>502 ± 65</td>
<td>75 ± 1.4</td>
<td>14.5 ± 1 × 10⁶</td>
<td>1</td>
</tr>
<tr>
<td>Wild type + R164S</td>
<td>Amp</td>
<td>37 ± 7</td>
<td>75 ± 25</td>
<td>2.0 ± 0.8 × 10⁵</td>
<td>4096</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>536 ± 51</td>
<td>2.5 ± 0.1</td>
<td>4.7 ± 0.4 × 10⁴</td>
<td>0.5</td>
</tr>
<tr>
<td>Stabilized v13 + R164S</td>
<td>Amp</td>
<td>140 ± 36</td>
<td>164 ± 24</td>
<td>1.1 ± 0.7 × 10⁸</td>
<td>n.m.</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>887 ± 173</td>
<td>5 ± 0.5</td>
<td>5.3 ± 2.5 × 10³</td>
<td>1</td>
</tr>
<tr>
<td>Wild type + G238S + R164S</td>
<td>Amp</td>
<td>11 ± 11</td>
<td>7 ± 3</td>
<td>0.9 ± 0.5 × 10⁶</td>
<td>512</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>578 ± 26</td>
<td>2.6 ± 0.1</td>
<td>4.5 ± 0.5 × 10⁴</td>
<td>0.125</td>
</tr>
<tr>
<td>Stabilized v13 + G238S + R164S</td>
<td>Amp</td>
<td>6 ± 0.85</td>
<td>11 ± 0.2</td>
<td>1.8 ± 0.4 × 10⁵</td>
<td>n.m.</td>
</tr>
<tr>
<td></td>
<td>Ctx</td>
<td>1300 ± 137</td>
<td>12.5 ± 0.85</td>
<td>9.6 ± 2 × 10⁵</td>
<td>1</td>
</tr>
</tbody>
</table>

n.d. is not detectable. A linear dependency of initial rates on substrate concentrations was observed (while applying the highest substrate concentrations possible within maximal absorbance of OD₂₆₄ nm ≤ 1.5), and thus, only $k_{cat}/K_M$ could be determined.

n.m. is not measured. The increased stability may affect the export rates and yields of TEM-1 to the periplasm, and thus, the MIC values of stabilized variants were not measured.

Amp, ampicillin; Ctx, cefotaxime.

* v13 is a stabilized variant of TEM-1 described in Ref. [16] that exhibits essentially the same enzymatic parameters as the wild-type TEM-1.

increases cefotaxime resistance by only ~8-fold [7,31] (MIC =0.5 μg/ml) but exhibits increased resistance levels with other, primarily second-generation antibiotics [31,32], as well as a weaker tradeoff than G238S with ampicillin resistance (Table 1; see Ref. [7]). In the absence of epistasis, the combined effect of the R164S and G238S mutations is expected to be multiplicative at the level of MIC (or additive at the level of log MIC [33]), thus resulting in up to ~120-fold higher cefotaxime resistance levels relative to wild type. However, the double mutant’s MIC is only increased ~2-fold relative to wild type (Table 1). Accordingly, the catalytic efficiency ($k_{cat}/K_M$) of the R164S/G238S mutant with cefotaxime is essentially identical with that of the single mutant R164S and about 20-fold lower than that of G238S (Table 1; see Ref. [7]). These losses in catalytic efficiency, combined with a more minor effect on TEM-1’s conformational stability [7], result in the in vivo MIC levels of the R164S/G238S mutant being lower than each of the single mutants (Table 1; see also Ref. [7]). In addition, the R164S/G238S mutant exhibits a very low $k_{cat}$ value with ampicillin, the original substrate, and accordingly much reduced MIC levels. The $K_M$ of the double mutant with ampicillin was improved. However, given the covalent intermediate mechanism of TEM-1, this change may in fact relate to a drastic slowdown in the rate of deacylation and not improved substrate affinity [34]. Further, in both the wild type and the mutants, the $K_M$ is much lower than the applied antibiotic concentrations, and therefore, the improvement in $K_M$ has little effect on resistance levels. Thus, R164S and G238S exhibit negative reciprocal epistasis: the expected positive effect of both individual mutations is inverted to negative when they are combined.

Crystal structures

Due to their compromised configurational stability, the single mutants, let alone the double mutant, were poorly expressed in *Escherichia coli* [31]. Their tendency to crystallize was also poor, especially of the R164S/G238S double mutant. Indeed, although a whole range of TEM-1 mutant structures is available, a structure of this double mutant has not been thus far attainable, not even at the background of a stabilizing mutation, M182T, that is routinely observed in clinical variants. Therefore, we made use of a stabilized variant of TEM-1 dubbed v13 that carries seven stabilizing mutations located on the surface and away from the active site, including mutations that are commonly observed in clinical variants such as M182T (for a list of mutations and their structural locations, see Supplementary Fig. 1). This variant is essentially identical in structure with wild-type TEM-1 (Ref. [16]), and the kinetic parameters of the studied 164 and 238 mutants in the background of v13 are essentially identical with the same mutations in the wild-type background (Table 1). Further, structures of v13 show the same active-site loop configurations and the same levels of loop mobility as wild-type TEM-1 (Ref. [16]). The individual effects of G238S and R164S, and the nonadditive effects of combining them, on the $k_{cat}/K_M$ values for both cefotaxime and ampicillin are essentially identical with the effects of these mutations at the wild type’s background (Table 1). The in vivo effects
could not be measured because the hyperstability of v13 has major effects on periplasmic export. However, the inconsistencies in the MIC level are the outcome of differences in expression levels and export rates and are therefore irrelevant to issues under study here. The comparison of the in vitro measured kinetic parameters and structures of the wild-type background (Table 1; Ref.[16]) indicate that v13 comprises a valid model for studying the structural and mechanistic origins of the loss of TEM-1’s catalytic function when G238S and R164S are combined as manifested in $k_{cat}/K_{M}$ values and especially in $k_{cat}$ (discussed below).

Overall, we solved 10 different structures of stabilized variant carrying the studied active-site mutations, individually and in combination (R164S/G238S) at both cryogenic and room temperatures. We obtained structures of the enzyme form and structures bound to EC25 (Ref. [35]), a covalent borate-based inhibitor that mimics the deacylation tetrahedral intermediate II [36] (Supplementary Fig. 2). EC25’s structure resembles ampicillin and not cefotaxime. However, a suitable boronate mimic of cefotaxime is not available (the one available, LP08, only mimics the thiazolyl part but not the lactam ring and its bulky substituents [35]). However, in the double mutant, cefotaxime and ampicillin show the very same trend of loss of activity relative to the single G238S mutant, both in relation to MIC values in vivo and to the in vitro kinetic parameters (Table 1). Additionally, as shown below, the origins of negative epistasis are in the disturbance of TEM-1’s catalytic machinery, and the mechanism of catalysis is essentially identical for both cefotaxime and ampicillin [37].

Structures were solved by molecular replacement and refined to 1.05–2.4 Å resolutions. The anisotropic $B$-factors were refined in the high-resolution data as detailed in Supplementary Table 1. All of the newly solved structures align well to one another and to the published wild-type TEM-1 structure (PDB code: 1ZG4; Ref. [38]). The only significant conformational differences between the structures are in the loops surrounding the active site, foremost in the $\Omega$-loop where R164S resides and in the 238-loop that includes G238S. Analyzing the main loop conformations in each of these structures reveals that the G238S mutation samples between two well-defined discrete 238-loop conformations but that R164S results in a diverse $\Omega$-loop conformational ensemble (Fig. 1). It should be noted that the crystal packing of the TEM-1 structures obtained here involves the $\Omega$-loop where R164S resides. Nonetheless, all structures presented in this study exhibit the same space group $C_{2}$, although the published wild-type TEM-1 structure relates to a different one ($P_{2}1_{2}1_{2}$; Ref. [38]). The comparison of the single to the double mutant is therefore based on all structures belonging to the same space group. Additionally, packing interactions tend to be stabilizing and thus to favor a single conformation. Hence, if anything, our data underestimate the conformational ensembles of the mutants and specifically of the double mutant.

The effects of G238S

In v13-G238S mutant, the dominant conformation of the 238-loop is perturbed, while the $\Omega$-loop conformation more closely resembles the wild-type

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**Fig. 1.** Superposition of active-site loops in all structures. (a) Superposition of the main $\Omega$-loop conformation in all wild-type structures (gray) and in the different v13-R164S mutant structures (blue). Room temperature structures are in green; note that, in the double mutant (R164S/G238S) room temperature structure, the $\Omega$-loop could not be modeled and is therefore missing. Shown in sticks are the side chains of N170 that align the deacylating water (W1; gray for wild type, red for mutants) and the catalytic S70 for reference. (b) Superposition of the main 238-loops conformation in all v13 structures containing the G238S mutation (blue), in wild type (gray). The mutant structures determined at room temperature are in green. Shown is A237 whose backbone amide positions the oxyanion hole water (W2). The loop conformations shown relate to the main clearly observed in the electron density of individual structures. Structures included are 4OPY, 4OPQ, 4OP5, 4OQI, 4OQ0, 4OQG, 1ZG4, 4OPQ, 4OP8, 4OP4, 4OQI, 4OPZ, 4OQ0, 4OQG and 1ZG4 (Ref. [38]) (Supplementary Table 1).
conformation (Fig. 2). Inspection of the 238-loop Fourier difference density maps of both the free and EC25-bound structures revealed alternative loop conformations that resemble a wild-type-like conformation (Figs. 2 and 3, G238S, middle top and bottom: density for the loop is fit by the dominant “major” conformation shown in thick sticks and a secondary “minor” conformation shown as thin sticks). The alternative, open conformation, presumably allows access to the much larger cefotaxime substrate and thereby increases the $k_{cat}/K_m$ of G238S by $\geq$130-fold relative to wild type in which the closed conformation prevails. However, the perturbed, open conformation also results in the loss of a key interaction between the $\Omega$-loop and 238-loop (E240’s backbone amide with N170’s backbone carbonyl oxygen; Figs. 2 and 3 shown as red broken circle). Apart from connecting the two loops, the interaction with E240 seems to be critical for stabilizing N170, a key catalytic residue that coordinates the deacylating water molecule [39]. Concordantly, there is electron density consistent with a second N170 side-chain conformation in a noncatalytic orientation. However, unlike v13-R164S, in the v13-G238S structure, the main-chain conformation of N170 closely resembles the catalytically competent wild-type conformation (Fig. 2). Upon ligand binding, the minor conformation of the 238-loop shifts to restore the interaction between the loops (Fig. 3, G238S, middle bottom: the red broken circle encompasses the wild-type and minor 238-loop conformations). Supplementary Movie 2 illustrates the relatively small changes in the $\Omega$-loop’s configuration upon inhibitor binding in comparison to the R164 mutant.

The incomplete shift of the minor conformation to the catalytically competent conformation suggests that formation of the transition states leading to and/or from the acyl-enzyme intermediate involves a higher entropic penalty for G238S than for wild type. As shown below, this cost seems far more pronounced in the R164S mutant that samples between many perturbed $\Omega$-loop conformations in the free enzyme.

The effects of R164S

As previously reported [28,40,41], the R164S mutant increased the conformational freedom of the $\Omega$-loop, as indicated by high B-factors and a weak electron density (Table 2 and Supplementary Figs. 4 and 5) relative to other parts of the protein. Indeed, the increased conformational freedom of the $\Omega$-loop was suggested to increase TEM-1’s active-site accessibility for larger substrates such as cefotaxime [25,31]. The other active-site loops, including the 238-loop, are well ordered in the R164S free and inhibitor-bound v13 structures (Fig. 2). The dominant conformation of the $\Omega$-loop is distinct from the wild-type conformation ($C^\alpha$ RMSD is $\sim$2 Å; Fig. 2). Accordingly, the interaction between E240-NH1 and N170-O is lost (Fig. 3, shown as the red broken lines). Indeed, in the inhibitor-free structure, N170 exhibits an outward rotation relative to the wild-type conformation, away from the deacylating water (Fig. 2). The deacylating water is accordingly shifted, although its other ligating residue, E166, exhibits wild-type conformation. Upon EC25 binding, the mobility of the $\Omega$-loop is reduced, the critical interaction with the 238-loop is restored and N170, as well as E166, adopts wild-type conformation (Table 2 and Fig. 2). Supplementary Movie 1 illustrates the changes in the $\Omega$-loop’s configuration upon inhibitor binding, the concomitant gain of interaction between the $\Omega$-loop and 238-loop and the relocation of N170.

The binding of EC25 to TEM-1 is covalent (Supplementary Fig. 2), thereby promoting its ability to shift the equilibrium toward the well-organized conformation of the $\Omega$-loop and N170. The cost of N170’s alternative, noncatalytic conformation in the case of the actual substrate is therefore likely to be much higher. These results suggest that substrate binding to the R164S mutant involves an increased entropic penalty due to the poor preorganization of the $\Omega$-loop, as indicated by N170’s perturbed conformation, and the elevated B-factors for the entire $\Omega$-loop (a detailed analysis is provided below). However, the perturbed $\Omega$-loop conformational dynamics of the free enzyme are likely required to accommodate larger second and third-generation cephalosporin antibiotics. The observed differences in loop configurations, and particularly of the $\Omega$-loop, may relate to crystal packing and may thus fail to report the enzyme’s actual configurations in solution. Note, however, that all v13 structures presented here belong to the same space group (Supplementary Table 2). Additionally, the $\Omega$-loop’s configuration does not seem to vary differently between structures belonging to different space groups, suggesting that crystal packing does not significantly alter this loop’s configuration (Supplementary Fig. 3).

The combined effect of R164S and G238S

A systematic analysis of the v13 double mutant’s structure did not reveal potential effects of the mutations on substrate binding. Indeed, neither R164S nor G238S is in direct contact with the substrate (as far as can be judged from the EC25 complex) and neither takes part in catalysis. Therefore, we examined the structures for longer-range effects. Both single mutants exhibit conformational changes within the active-site loops that are necessary for accommodating the larger cefotaxime substrate [29,41]. However, these two mutations have different side effects on the
Fig. 2. The $\Omega$-loop and 238-loop interactions network. A snapshot of the $\Omega$-loop, the 238-loop and the key catalytic residues in the free structures (left column) and in structures with the inhibitor EC25 bound (right column). Shown are wild-type TEM-1 (gray), the single v13 mutants R164S (yellow) and G238S (blue) and the double v13 mutant R164S/G238S (green). Catalytic residues are in smoked pink and EC25 is in light pink balls and sticks. Minor conformations of the loops are shown in light blue. Broken lines represent putative interactions involving R164 and G238, and interactions lost in the R164S/G238S double mutant are marked in blue in the wild type. The nonnative interaction in the double mutant between E171 and S238 is in red. A putative stacking interaction between R164 and R178 (Ref. [42]) is also marked.
active-site conformational ensemble, suggesting that the mechanism underlining TEM-1’s adaptation toward cefotaxime is more complex than previously proposed [43,44]. In the v13 R164S/G238S double mutant, a wider ensemble of conformations that includes nonproductive conformations in both the free and inhibitor-bound states is seen (Fig. 2). Specifically, like in the v13-R164S mutant, the Ω-loop of the double mutant is conformationally heterogeneous. Additionally, like in the v13-G238S mutant, the perturbed conformation of the 238-loop dominates, and the near-wild-type conformation is scarcely populated (Figs. 2 and 3, R164S/G238, right top: the major and minor conformations of the 238-loop are shown as thin sticks with the position of E240-N indicated by spheres in both minor and major conformations). However, in the double mutant, the 238-loop and Ω-loop fail to establish interactions that stabilize the catalytically competent conformation of N170. Further, N170’s side-chain carbonyl oxygen takes the place of the deacylating water that is not present in the double mutant’s free structure, although E166 exhibits a wild-type-like conformation. Even covalent inhibitor binding is insufficient to drive the rearrangement. Supplementary Movies 3 and 4 address the minor and major 238-loop conformations, respectively. The morph movies illustrate the profound changes in the Ω-loop’s configuration upon inhibitor binding, the loss of interaction between the Ω-loop and 238-loop in both the free and inhibitor-bound structures and the displacement of N170.

Table 2. The Ω-loop and 238-loop B-factors.

<table>
<thead>
<tr>
<th>Structure</th>
<th>Relative loop B-factors (Å²)</th>
<th>Free enzyme</th>
<th>EC25 complex</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild-type 238-loop</td>
<td>0.98</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Wild-type Ω-loop</td>
<td>1.24</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>G238S 238-loop</td>
<td>1.18</td>
<td>1.06</td>
<td>—</td>
</tr>
<tr>
<td>G238S Ω-loop</td>
<td>2.56</td>
<td>1.75</td>
<td>—</td>
</tr>
<tr>
<td>R164S 238-loop</td>
<td>1.02</td>
<td>1.12</td>
<td>—</td>
</tr>
<tr>
<td>R164S Ω-loop</td>
<td>2.54</td>
<td>2.22</td>
<td>—</td>
</tr>
<tr>
<td>R164S/G238S 238-loop</td>
<td>1.46</td>
<td>1.24</td>
<td>—</td>
</tr>
<tr>
<td>R164S/G238S Ω-loop</td>
<td>4.75</td>
<td>3.34</td>
<td>—</td>
</tr>
</tbody>
</table>

B-factors indicate the level of thermally induced mobility of the loops’ backbone atoms.

*Calculated for the set of the structures obtained at cryotemperature (Supplementary Table 1). Noted are the average B-factors for backbone atoms of all loop residues, normalized to the average B-factor for all backbone atoms in a given structure.

Supplementary Movies 3 and 4 address the minor and major 238-loop conformations, respectively. The morph movies illustrate the profound changes in the Ω-loop’s configuration upon inhibitor binding, the loss of interaction between the Ω-loop and 238-loop in both the free and inhibitor-bound structures and the displacement of N170.

The loss of the catalytically competent N170 conformation relates to a new, nonproductive interaction seen exclusively in the double mutant: in the bound structure, E171’s side chain interacts with the backbone amide of E240 within the perturbed conformation of the 238-loop (Figs. 2 and 3, R164S/G238, right top: The interaction between E171 and E240 is indicated by red broken lines). The nonproductive 171-240 interaction originates to the...
loss of E171’s side-chain interaction with R164 upon mutating this residue to Ser. However, the 171-240 interaction does not occur in the v13-R164S single mutant simply because the perturbed 238-loop conformation is only populated when G238 is mutated to Ser. Therefore, negative epistasis between these two mutations has two primary structural causes: (1) excessive conformational flexibility when the two mutations are combined decreases the preorganization of the enzyme active site in the free state, and (2) a nonnative 171-240 interaction between the Ω-loop and 238-loop seems to shift the equilibrium against the productive active-site conformation. As discussed in detail below, the shift toward nonproductive conformations is also apparent in the double mutant’s free structure, where the deacylating water molecule is missing and, further, N170’s side chain and the catalytic S70’s hydroxyl are within contacting distance. This interaction is likely to severely hamper the active site (Fig. 2).

The disturbance of active-site preorganization

Enzyme active sites, including TEM-1’s active site, are preorganized in a catalytically optimal conformation [45,46]. However, the high mobility of the Ω-loop, in both the v13 single R164S mutant and especially in the v13-R164S/G238S double mutant, induced alternative, suboptimal configurations (Fig. 4a). To determine the magnitude of conformational change the apo-enzyme exhibits in order to get to the catalytically optimal conformation, we examined the differences between the free and the inhibitor-bound enzyme structures. Specifically, we observed the displacement of N170, a residue that coordinates the deacylating water molecule [39]. In wild type and G238S, the overall deviations in the active site’s structure upon binding are minimal (RMSD ≤ 0.39 Å; Table 3) and thus N170 is largely prealigned for catalysis. In v13-R164S and R164S/G238S, however, N170 is misaligned in the free structures, and the Ω-loop exhibits a large conformational change upon inhibitor binding (RMSD ≈ 1.85 Å; Table 3 and Fig. 4). This displacement dominates the conformational ensembles of the v13-R164S mutant and the v13 double mutant R164S/G238S but is only a minor

![Image](image_url)

**Table 3. Structural deviations between the free and inhibitor-bound structures**

<table>
<thead>
<tr>
<th>Structure</th>
<th>All residues (H25-W286)</th>
<th>Ω-loop (E164-R179)</th>
<th>238-loop (S238-S242)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>0.37</td>
<td>0.18</td>
<td>0.22</td>
</tr>
<tr>
<td>G238S</td>
<td>0.34</td>
<td>0.39</td>
<td>0.20</td>
</tr>
<tr>
<td>R164S</td>
<td>0.39</td>
<td>1.83</td>
<td>0.07</td>
</tr>
<tr>
<td>R164S/G238S</td>
<td>0.33</td>
<td>1.86</td>
<td>0.12</td>
</tr>
</tbody>
</table>

Noted are average RMSD values for main-chain atoms (in angstroms).
conformation in the v13-G238S mutant (Fig. 4a and Table 4). Consistent with this structural analysis, the R164S mutation was shown to impair deacylation in class A β-lactamases [40,41]. Thus, the prevailing conformational states of v13-R164S and v13-R164S/G238S probably exhibit no or poor catalytic activity. The entropic cost of selecting from multiple conformations, combined with the impaired configuration of N170 even in the inhibitor-bound structure, is manifested in the very low $k_{\text{cat}}$ value of the double mutant. As expected, inhibitor binding shifts the equilibrium toward the catalytic conformation in all mutants (Fig. 4b and Table 4). However, even in the inhibitor-bound v13-R164S/G238S structure, a wild-type-like configuration of N170 is only observed at 0.5 occupancy alongside an alternative, highly perturbed conformation that is unlikely to be catalytically active (Fig. 4b, in green; Table 4).

### B-Factors analysis

To further test the idea that increased conformational flexibility underlies the sign negative epistasis between R164S and G238S, we examined the refined crystallographic temperature factors ($B$-factors) along TEM-1’s polypeptide chain. $B$-Factors model how much an atom deviates from its average position in the crystal [47,48]. However, this model includes both dynamic and static disorders and is sensitive to model errors. $B$-Factors are especially useful reporters of molecular flexibility when discrete conformations cannot be distinguished in the electron density. Since alternative conformations for the 238-loop are apparent at low electron density levels (Supplementary Fig. 4 and 5), we carefully modeled these fragments and then normalized the $B$-factors within each structure to indicate the relative mobility of atoms [48]. The normalized $B$-factors indicate that, in

<table>
<thead>
<tr>
<th>Interaction distances (Å)</th>
<th>Interaction number</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>7</th>
<th>8</th>
<th>9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Free enzyme</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wild type</td>
<td></td>
<td>2.5</td>
<td>2.8</td>
<td>2.8</td>
<td>2.4</td>
<td>3.2</td>
<td>2.9</td>
<td>2.8</td>
<td>3.0</td>
<td>2.7</td>
</tr>
<tr>
<td>R164S</td>
<td></td>
<td>2.7</td>
<td>3.5</td>
<td>3.2</td>
<td>2.9</td>
<td>2.9</td>
<td>3.1</td>
<td>2.7</td>
<td>2.9</td>
<td>2.7</td>
</tr>
<tr>
<td>G238S</td>
<td></td>
<td>2.5</td>
<td>2.3</td>
<td>2.9</td>
<td>2.8</td>
<td>2.4</td>
<td>2.4</td>
<td>3.0</td>
<td>2.7</td>
<td>2.9</td>
</tr>
<tr>
<td>6R164S/G238S</td>
<td></td>
<td>n.d.</td>
<td>n.d.</td>
<td>n.d.</td>
<td>2.9</td>
<td>2.9</td>
<td>3.1</td>
<td>2.7</td>
<td>2.8</td>
<td>2.7</td>
</tr>
<tr>
<td>EC25 bound</td>
<td></td>
<td>2.8</td>
<td>2.9</td>
<td>2.2</td>
<td>2.6</td>
<td>3.8</td>
<td>3.1</td>
<td>2.9</td>
<td>2.5</td>
<td>3.1</td>
</tr>
<tr>
<td>Wild type</td>
<td></td>
<td>2.5</td>
<td>2.6</td>
<td>2.6</td>
<td>2.5</td>
<td>2.5</td>
<td>2.8</td>
<td>2.8</td>
<td>2.8</td>
<td>2.6</td>
</tr>
<tr>
<td>R164S</td>
<td></td>
<td>2.5</td>
<td>2.7</td>
<td>2.5</td>
<td>2.5</td>
<td>2.8</td>
<td>2.8</td>
<td>2.7</td>
<td>2.8</td>
<td>2.8</td>
</tr>
<tr>
<td>G238S</td>
<td></td>
<td>2.5</td>
<td>2.7</td>
<td>2.4</td>
<td>2.4</td>
<td>3.0</td>
<td>2.8</td>
<td>2.7</td>
<td>2.8</td>
<td>2.7</td>
</tr>
<tr>
<td>R164S/G238S</td>
<td></td>
<td>2.5</td>
<td>6.1</td>
<td>6.1</td>
<td>6.1</td>
<td>n.a.</td>
<td>n.a.</td>
<td>n.a.</td>
<td>n.a.</td>
<td>n.a.</td>
</tr>
</tbody>
</table>

The table depicts a superposition of the key active-site residues in free (gray, PDB code: 1ZG4; Ref. [38]) and bound (green, PDB code: 4OQG) wild-type structures. The distances for the indicated bonds are provided in the table. n.d. is not detected; the interaction is not observed because at least one of the interacting TEM-1 side-chain atoms is disordered. n.a. is not applicable. These contacts are not hydrogen bonds. The table’s figure is TEM-1’s active-site configuration.

\( ^{a} \) N170 appears in two alternative conformations in this structure.

\( ^{b} \) The oxyanion hole water or the borate oxygen atom mimicking it.

\( ^{c} \) The deacylating water or the borate oxygen atom mimicking it.
the free structures of the mutants, both the Ω-loop and G238-loop exhibit higher flexibility in the mutants than in wild-type TEM-1. Interestingly, while R164S mutation induces only local changes in the flexibility of the Ω-loop, the G238S mutation induces alternative conformations of the 238-loop and increased B-factors in both the 238 and the Ω-loop. In the double mutant, we observed that both of the loops exhibit increased B-factors. However, the Ω-loop exhibits normalized B-factors that are almost 2-fold higher than in each of the single mutants (Table 2). To determine how flexibility changes upon inhibitor binding, we analyzed the patterns of B-factors in the inhibitor-bound structures. In both of the single mutants, inhibitor binding reduced the B-factor values. However, in the double mutant, the reduction in the loop’s B-factors upon binding is much lower than in any of the single mutants (Table 2). The B-factor analysis therefore supports the hypothesis that the double mutant exhibits a higher degree of conformational freedom compared to the single mutants. Hence, we suggest that both the negative epistasis between R164S and G238S mutant and the decreased evolvability of alleles carrying R164S result from the mutations widening the conformational ensembles to a level that penetrates into the active site and hampers catalysis.

Discussion

Sign epistasis modulates the fitness landscape such that traversing from one peak (i.e., a given sequence and function) to another may occur via only few trajectories [3,49,50]. Understanding the underlying causes of this particular form of epistasis may make evolutionary processes, and enzyme design and engineering, easier to understand and predict [2]. Here, we revealed the basis of reciprocal sign epistasis between two mutations in TEM-1 β-lactamase that initiate alternative resistance pathways to second and third-generation cephalosporins antibiotics including cephalosporins. As nearly all other new function mutations in enzymes, R164S and G238S occur not in the catalytic residues and not even in residues that directly contact the substrate [51]. Rather, they are located in loops that surround the active site and in residues that are >10 Å apart. Nonetheless, these mutations strongly interact phenotypically at the level of enzyme function. Individually, each of the studied adaptive mutations introduces conformational changes in the active-site loops that enable TEM-1’s active site to better accommodate a new larger substrate. However, the level of conformational diversity introduced by these mutations and the side effects of widening the enzyme’s conformational ensemble differ between these two mutations. While one mutation, G238S, introduces new well-defined conformations, the other, R164S, introduces disorder along large segments of the Ω-loop (Fig. 1). Further, in the G238S mutant, the 238-loop conformation dominates in both the free and inhibitor-bound enzyme forms—hence, this mutation simply reshapes the loop (Table 3). In contrast, R164S induces an ensemble of different conformations, most of which are of nearly equal stability and are hence rarely represented (manifested in the weak electron density of the Ω-loop in the crystal). Binding of the inhibitor and, accordingly, substrate binding and formation of the transition state shift the equilibrium toward the catalytically active conformation (Table 3).

The existence of multiple nonactive conformations is manifested in the $k_{cat}$ value of the R164S/G238S double mutant with ampicillin (TEM-1’s preferred substrate) being ~10-fold lower than that of R164S and 260-fold lower than that of wild type (Table 1). Assuming “conformational selection”, that is, multiple preexisting conformations and substrate binding shifting the equilibrium toward the functional conformation, the measured $k_{cat}$ relates to the $k_{cat}$ of individual conformers multiplied by their relative representation. Thus, a loss of ~100-fold in $k_{cat}$ upon combining R164S and G238S (relative to the expected additive effect) suggests that the fraction of the catalytically optimal conformers dropped to ~1%. The role of multiple preexisting conformers in mediating the multifunctionality of proteins [52,53] and their evolutionary potential [54] is widely recognized. However, the evolutionary cost of excessive conformational freedom, as observed here, is generally overlooked. One case highlighting the cost of increased conformational freedom relates to a laboratory engineered trypsin/chymotrypsin loop swap [55]. This swap resulted in an enzyme that was not only catalytically impaired but also lacked the expected chymotrypsin-like specificity. The low turnover rate and specificity related to the high mobility of active-site loops and their stabilization via a single point mutation led to the recovery of the enzyme’s rate and specificity. In triosephosphate isomerase [56], mutations destabilizing the loops surrounding the active site severely hamper catalytic activity and especially the $k_{cat}$ as observed here. Similarly, in RNase H [57], insertion/deletion of a glycine altered the ensemble of loop conformers and modulated the catalytic rate. Here, we addressed the effects of mutations that occurred along adaptive evolutionary trajectories. We found that both G238S and R164S increase the representation of a new active-site conformation that better accommodates cefotaxime. However, combining these two mutations results in noncatalytic conformers dominating the conformational ensemble and in nonnative interactions between the loops such as the hydrogen bond between E171 and E240 (Fig. 2; R164S/G238S, red broken lines).
Thus, in TEM-1, sign epistasis between mutations that increase enzyme function seems to be the outcome of “conformational anarchy” whereby the catalytically optimal conformation becomes diluted to a degree that severely hampers catalytic efficiency. Previous work described the role of compensatory mutations with global stabilizing effects in the evolution of new enzyme functions and specifically in TEM-1 [31,51]. However, as far as the catalytic parameters are concerned, the negative epistasis effect between R164S and G238S in TEM-1 is not alleviated by global, stabilizing, compensatory mutations—not by M182T (Refs. [7,24] and [31]). TEM-1’s most frequently observed stabilizing mutation and not even in v13 that carries seven stabilizing mutations including another known global suppressor, L201A (Ref. [58]). The inability to evolve a more stable and rigid core and scaffold to compensate for increased mobility of the active-site loops relates to the polarized architecture of TEM-1 (Ref. [16]).

Our results therefore reveal a delicate balance between the adaptive benefit associated with increased structural freedom and its cost. An example of this tradeoff is antibody affinity maturation. Germ-line antibodies exhibit high conformational plasticity yet low affinity, which is compensated by the higher avidity of IgM antibodies. Maturation involves the funneling of conformations toward the one that binds the antigen and hence increase affinity [59,60].

Why do these TEM-1 mutations behave so differently—R164S leading to a local adaptive peak versus G238S leading toward higher fitness? Both R164 and G238 are part of an elaborate network that shapes TEM-1’s active site (Fig. 2). However, whereas R164 has seven contacts with neighboring residues, G238 has only two. Contacts density correlates with how buried a residue is [61], yet R164 and G238 exhibit the same relative solvent accessibility. Thus, R164 exhibits higher than average contact density for its degree of burial (mean = 4.8 contacts per residue) whereas G238 is far below average (Supplementary Fig. 6). This difference goes beyond the expected for the different sizes of these amino acids. The high connectivity of R164 results in the collapse of this interaction network upon mutating to serine (Fig. 2) and the appearance of noncatalytic conformations (Fig. 4). Negative epistasis in TEM-1 may therefore relate to the network properties of protein folds. These so-called “small-world” networks [62,63] are characterized by most residues (nodes) being connected via a few hubs. In such networks, highly connected residues as R164 are relatively rare, and their perturbation severely undermines the network [63]. These networks, however, are highly robust to changes in weakly connected nodes as G238. Accordingly, position 238 shows ~10-fold faster evolutionary rate than 164 (238 evolves 3× slower than TEM-1’s average and R164 30-fold slower as calculated by rate4site [64]). Owing to its limited connectivity within the active site’s interactions network, position 238 exhibits high robustness and innovability—the potential to promote new enzymatic functions [65].

Overall, our analysis suggests a tight link between protein structural networks, conformational ensembles and evolvability. At least for the case of TEM-1 β-lactamase, we observed a delicate balance between the potential of mutations to introduce new conformations that may initiate new functions and their cost in over-broadening the protein’s conformational ensemble. The latter appears to result in negative epistasis and what seems to be an evolutionary cul-de-sacs in TEM-1’s adaptation via the R164S mutation. Better understanding of how a protein’s architecture, structural networks and conformational dynamics at different timescales affect this balance will promote our understanding of protein evolution and also advance the design and engineering of new proteins [66,67].

Materials and Methods

Crystallization

Wild-type TEM-1 and mutants were expressed and purified as previously described [16]. For crystallization, concentrated solutions of TEM-1 variants (60 mg/ml) in 25 mM Tris (pH 8.4) and 100 mM NaCl were submitted to crystallization trials. Crystallization was performed using the hanging-drop vapor diffusion method. Equal volumes (0.5 μl) of protein and reservoir solutions were mixed, and the resulting drops were equilibrated at 293 K against a 400-μl reservoir solution made of 9% (wt/vol) polyethylene glycol (PEG) 8000, 100 mM 4-morpholineethanesulfonic acid (Mes) (pH 6.2) and 200 mM Ca(OAc)₂ for all mutants and 6% PEG 8000, 100 mM Mes buffer (pH 6.2), 200 mM Ca(OAc)₂ and 50 μM NaF for the wild type. Microseeding was performed to obtain larger crystals. For the structures bound to EC25 (Ref. [35]), the crystals were soaked for 1 h at 293 K in a solution containing 50 mM EC25, 20% (wt/vol) PEG 8000, 25% PEG 600, 100 mM Mes (pH 6.5) and 200 mM Ca(OAc)₂ for all mutants. The wild-type TEM-1 crystals were soaked using the same protocol in a solution containing 50 mM EC25, 25% PEG 600, 100 mM Mes (pH 6.5) and 200 mM Ca(OAc)₂.

Data collection at 100 K

Crystals of ligand-free proteins were mounted on microloops (MiTeGen, USA) and transferred into a cryoprotectant solution containing 20% PEG 8000, 25% PEG 600, 100 mM Mes (pH 6.5) and 200 mM Ca(OAc)₂ for 1 min and were then flash-cooled in liquid nitrogen. For the bound wild-type and v13 structures, the soaking solution was used as cryoprotectant. X-ray diffraction data of mutant G238S free and in complex with EC25, R164S and R164S/G238S in complex with EC25 were collected on a Rigaku R-AXIS IV++ imaging plate area detector mounted on a Rigaku RU-3R.
generator with CuKα radiation focused by Osmic confocal mirrors (in-house source). The X-ray diffraction data of the wild-type crystals in complex with EC25, of v13 mutants R164S and R164S/G238S (free forms), were collected at ID29 beamline (European Synchrotron Facility, Grenoble, France) using a PILATUS 6M detector. The diffraction data of R164S crystals were collected on ID14-1 beamline (European Synchrotron Facility) using an ADSC Q210 detector.

**Data collection at 293 K**

Crystals were mounted on microloops (MiTeGen) and transferred into capillaries (Hampton, UK). Diffraction data were collected on a Rigaku R-AXIS IV++ imaging plate area detector mounted on our in-house source.

**Data processing and refinement**

Diffraction data were integrated and scaled using XDS package [68]. Molecular replacement was performed using MOLREP [69] using the structure of wild-type TEM-1 (PDB code: 1ZG4; Ref. [38]) as the starting model. Manual model improvement was performed using Coot [70] and the refinement was performed using REFMAC5 (Ref. [69]) and PHENIX [71]. Data collection and refinement statistics are provided in Supplementary Table 1. Figures depicting structures were prepared with PyMOL.

**Relative loops B-factors**

The occupancies of all residues in all analyzed structures were set at a value of 1 (for residues with alternative conformations, the sums of occupancies were set at 1). The structures were re-refined using REFMAC5 (Ref. [69]). The relative loops B-factor values were obtained by normalizing the B-factor values of the loops main-chain atoms by the average B-factor of main-chain atoms of the whole structure as described in Ref. [16].

**RMSD between free and bound structures**

RMSD calculations were performed using the Swiss-PDB viewer, using main-chain atoms of the loops or the entire structures.

**Evolutionary rates**

Evolutionary rates were calculated using rate4site [64] as described in Ref. [18]. This method computes the relative evolutionary rate for individual protein positions across the entire phylogeny, via Bayesian Estimation. The rates are normalized with respect to the phylogeny such that the expected rate over all sites is 1. Hence, the rate of each site indicates how rapidly the site evolves relative to the mean. The sequence alignment was taken from ConSurf and contained 150 sequences with 95–35% identity to wild-type TEM-1.

Number of contacts and accessible surface area values were calculated as in Ref. [16].

**Accession codes**

Protein Data Bank: Atomic coordinates and structure factors have been deposited with the accession codes PDB codes: 4OPY, 4OPQ, 4OP5, 4OQI, 4OQH, 4OQO, 4OOG, 4OP8, 4OPR and 4OPZ.

Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.jmb.2015.05.011.

**Acknowledgments**

Financial supports by the Israel Science Foundation (D.S.T), and EMBO Visiting Fellowship and National Institutes of Health (OD009180) to J.S.F., are gratefully acknowledged. D.S.T. is the Nella and Leon Benoziyo Professor. We thank Linda Shimon for invaluable assistance with the home source and room temperature X-ray data collections.

Received 9 January 2015; Received in revised form 8 May 2015; Accepted 12 May 2015
Available online xxxx

**Keywords:**
protein evolution; protein disorder; protein folds; conformational diversity; interactions network

**Abbreviations used:**
PEG, polyethylene glycol; Mes, 4-morpholineethanesulfonic acid.

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Please cite this article as: Dellus-Gur Eynat, et al, Negative Epistasis and Evolvability in TEM-1 β-Lactamase—The Thin Line between an Enzyme’s Conformational Freedom and Disorder, J Mol Biol (2015), http://dx.doi.org/10.1016/j.jmb.2015.05.011
Negative Epistasis and Evolvability in TEM-1


