

Correction to Analysis of the Structure and Function of YfcG from *Escherichia coli* Reveals an Efficient and Unique Disulfide Bond Reductase [(2009) *Biochemistry 48*, 6559. DOI: 10.1021/bi9008825]. Megan C. Wadington, Jane E. Ladner, Nina V. Stourman, Joel M. Harp, and Richard N. Armstrong*

Table 2: Steady-State Kinetic Constants for the Reduction of 2-Hydroxyethyl Disulfide by YfcG and Its C166A Mutant at 25 $^{\circ}\mathrm{C}$

enzyme	$k_{\rm cat} ({\rm s}^{-1})$	$k_{\text{cat}}/K_{\text{M}}^{\text{GSH}} (\text{M}^{-1} \text{ s}^{-1})$	$K_{\rm M}^{\rm GSH} ({\rm mM})$
YfcG	29 ± 2	$(1.8 \pm 0.3) \times 10^4$	1.6 ± 0.3
YfcG (C166A)	30 ± 2	$(3.0 \pm 0.7) \times 10^4$	1.0 ± 0.2

Because of an error in the analysis of the initial velocity data, the steady-state kinetic constants, $k_{\rm cat}$ and $k_{\rm cat}/K_{\rm M}^{\rm GSH}$, reported in Table 2 are incorrect and smaller than reported by a factor of 6.2. A corrected Table 2 appears above. This correction does not alter the conclusions made in the original report.

DOI: 10.1021/bi101851x Published on Web 11/23/2010