

## Corrections

Identifying the Physiological Electron Transfer Site of Cytochrome *c* Peroxidase by Structure-Based Engineering, by Mark A. Miller,\* Lois Geren, Gye Won Han, Aleister Saunders, James Beasley, Gary J. Pielak, Bill Durham, Francis Millett, and Joseph Kraut, Volume 35, Number 3, January 23, 1996, pages 667–673.

Page 670. In the legend to Figure 2, the sentence beginning on line 10 should read as follows: (Panel B, bottom) The positions of the yCc (white) and hCc (black) hemes are shown relative to CcP(MI) when bound in their respective modes.

Page 671. In Table 1,  $k_{et}$  for CcP:yCc at pH 7 should read 170. Also, errors are present in the footnote assignments and footnote legend. The table should appear as follows:

Table 1: Kinetic Parameters for ET from yCc to CcP and MPB-CcP(MI,A128,C193)<sup>a</sup>

	pH	$k_{et}^b$ ( $\times 10^{-4} \text{ s}^{-1}$ )	$k_{cat}^c$ ( $\times 10^{-2} \text{ s}^{-1}$ )	$k_a^d$ ( $\times 10^{-7} \text{ M}^{-1} \text{ s}^{-1}$ )	$k_b^e$ ( $\times 10^{-7} \text{ M}^{-1} \text{ s}^{-1}$ )	$k_{cat}/K_m^c$ ( $\times 10^{-7} \text{ M}^{-1} \text{ s}^{-1}$ )
CcP:yCc	6		22	20 <sup>f</sup>	3.5 <sup>f</sup>	3.8
	7	170 <sup>g</sup>		17 <sup>h</sup>	2.3 <sup>h</sup>	
MPB mutant:yCc	6		0.32 (1.5)	1.0 <sup>f</sup> (5)	0.19 <sup>f</sup> (5)	0.14 (4)
	7	1.3 (1.1)		0.5 <sup>h</sup> (3)	0.10 <sup>h</sup> (4)	
CcP:hCc	6	6.1	2.5	20 <sup>i</sup>	3 <sup>i</sup>	0.6
MPB mutant:hCc	6	3.0 (49)	1.2 (47)	8.0 <sup>i</sup> (40)	1.0 <sup>i</sup> (33)	0.6 (100)

<sup>a</sup> Transient ET rates using Ru-bipyridine derivatives of hCc and yCc (Geren et al., 1991; Hahm et al., 1993) and steady-state kinetic measurements were made as described under Materials and Methods. The values in parentheses are the percentage of the CcP(MI) parent. <sup>b</sup> Measured with Ru-39-yCc, in 3 mM sodium phosphate buffer. <sup>c</sup> Measured in 5 mM sodium phosphate buffer, pH 6.0, adjusted to 110 mM ionic strength with NaCl. <sup>d</sup> Bimolecular reduction of Trp 191 radical of compound I measured with stopped-flow spectrophotometer. <sup>e</sup> Bimolecular reduction of oxyferryl heme of compound II measured with stopped-flow spectrophotometer. <sup>f</sup> Measured in 10 mM MES, 300 mM NaCl. <sup>g</sup> The measurement of this parameter will be described in detail elsewhere (Geren et al., manuscript in preparation). <sup>h</sup> Measured in 5 mM sodium phosphate, 300 mM NaCl. <sup>i</sup> Measured in 10 mM MES, 100 mM NaCl.

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