

Effect of Single-Site Charge-Reversal Mutations on the Catalytic Properties of Yeast Cytochrome *c* Peroxidase: Mutations near the High-Affinity Cytochrome *c* Binding Site[†]

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ABSTRACT: Fifteen single-site charge-reversal mutations of yeast cytochrome *c* peroxidase (CcP) have been constructed to determine the effect of localized charge on the catalytic properties of the enzyme. The mutations are located on the front face of CcP, near the cytochrome *c* binding site identified in the crystallographic structure of the yeast cytochrome *c*–CcP complex [Pelletier, H., and Kraut, J. (1992) *Science* 258, 1748–1755]. The mutants are characterized by absorption spectroscopy and hydrogen peroxide reactivity at both pH 6.0 and 7.5 and by steady-state kinetic studies using recombinant yeast iso-1-ferrocycytochrome *c* (C102T) as a substrate at pH 7.5. Some of the charge-reversal mutations cause detectable changes in the absorption spectrum, especially at pH 7.5, reflecting changes in the equilibrium between penta- and hexacoordinate heme species in the enzyme. An increase in the amount of hexacoordinate heme in the mutant enzymes correlates with an increase in the fraction of enzyme that does not react with hydrogen peroxide. Steady-state velocity measurements indicate that five of the 15 mutations cause large increases in the Michaelis constant (R31E, D34K, D37K, E118K, and E290K). These data support the hypothesis that the cytochrome *c*–CcP complex observed in the crystal is the dominant catalytically active complex in solution.

Cytochrome *c* peroxidase (CcP)¹ is a detoxification enzyme localized between the inner and outer membranes of yeast mitochondria (1). CcP decreases toxic levels of hydrogen peroxide by catalyzing its reduction to water using ferrocycytochrome *c* (2). The catalytic mechanism involves oxidation of the native enzyme by hydrogen peroxide to an enzyme intermediate called CcP Compound I (CcP-I). CcP-I contains two oxidized sites, an oxyferryl Fe(IV) heme group and a tryptophan π -cation radical located within van der Waals distance of the heme. Interaction with, and electron transfer from, ferrocycytochrome *c* reduces CcP-I back to the

native state via a second enzyme intermediate, CcP Compound II (CcP-II), completing the catalytic cycle. Since 1980, when the three-dimensional structure of CcP was first reported (3, 4), CcP has played an important role in elucidating the structural basis for heme protein reactivity, especially in the activation of hydrogen peroxide (5, 6), long-range electron transfer between heme proteins (7, 8), and protein–protein interactions (9, 10).

The nature of the cytochrome *c*–CcP interaction during catalysis is still under active investigation with a number of questions still unresolved, including the number of cytochrome *c* binding sites on CcP, the cytochrome *c* affinity at each binding site, the location of the binding sites, the dynamic nature of cytochrome *c* bound at each site, and the electron transfer activity of cytochrome *c* bound at the various sites (7, 11).

The first suggestion of multiple cytochrome *c* binding sites on the surface of CcP came from the steady-state velocity studies of Margoliash and co-workers in 1977 (12). Since that time, a large number of studies have been published concerning the nature of the cytochrome *c*–CcP interaction and the potential locations of the cytochrome *c* binding sites on CcP (13–31). Northrup et al. (14) published an influential study modeling the electrostatic interaction between the positively charged cytochrome *c* molecule and the negatively charged CcP using Brownian dynamics simulations. Northrup and colleagues identified three areas on the surface of CcP that had a high probability for formation of productive electron transfer complexes, and these were located near Asp-34, Asp-148, and Asp-217 on the surface of CcP. An

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¹ Mutations in the amino acid sequences of either CcP or cytochrome *c* are indicated by using the one-letter code for the amino acid residue in the wild-type protein, followed by the residue number and the one-letter code for the amino acid residue in the mutant protein; i.e., C102T represents a mutant in which a threonine residue replaces the cysteine residue at position 102 of the wild-type protein. Abbreviations: CcP, generic abbreviation for cytochrome *c* peroxidase whatever the source; yCcP, authentic yeast cytochrome *c* peroxidase isolated from baker's yeast, *Saccharomyces cerevisiae*; rCcP, recombinant CcP expressed in *Escherichia coli* with an amino acid sequence identical to that of yCcP; CcP(MI), recombinant CcP expressed in *E. coli* with four amino acid variations compared to yCcP, a Met-Ile N-terminal extension and mutations T53I and D152G; ZnCcP, CcP in which zinc porphyrin replaces the heme; CcP-I, CcP Compound I, the first intermediate observed in the catalytic cycle produced upon oxidation of CcP with hydrogen peroxide; CcP-II, CcP Compound II, the second intermediate observed in the catalytic cycle produced by one-electron reduction of CcP-I; RZ, reinheitszahl or purity number, the ratio of the absorbance at the Soret maximum to the absorbance at the protein band maximum near 280 nm.

important milestone in the study of cytochrome *c*–CcP interactions was the determination of the crystallographic structures of two cytochrome *c*–CcP complexes by Pelletier and Kraut in 1992 (9). Pelletier and Kraut determined structures of 1:1 complexes of horse heart cytochrome *c* and CcP and of yeast iso-1-cytochrome *c* and CcP. They identified Asp-34 and Asp-290 as critical charged residues on CcP for formation of both complexes (9). A number of other studies using computer modeling (15–17) and chemical modification (18) have identified additional residues on CcP that may be important in cytochrome *c* binding. These include Asp-33, Asp-37, Asp-79, and residues between positions 221–224 and 290–294 as potential interaction sites.

Site-directed mutagenesis of CcP has been used to test some of the suggested locations for the cytochrome *c* binding sites (21–27). These studies consistently find that charge-neutralization or charge-reversal mutants of Asp-34, Asp-37, and Glu-290 inhibit cytochrome *c* binding, while mutations at Glu-32, Glu-35, Asp-79, Asp-148, Asp-217, and Glu-291 have little effect on cytochrome *c* binding. The only experimental study that provides information about the location of the secondary binding site is that of Leesch et al. (24), who found that the K149E mutation increases the level of formation of the 2:1 complex while having little effect on formation of the 1:1 complex.

The studies published to date are incomplete in that all potential cytochrome *c* binding sites may not have been identified by the theoretical and computer modeling studies and all of the potential binding sites have not been tested experimentally. We have initiated a systematic study to identify all of the negatively charged groups on CcP that may be involved in formation of both 1:1 and 2:1 yeast iso-1-cytochrome *c*–CcP complexes. We plan to mutate, individually, every aspartate and glutamate residue in CcP to a lysine residue. We will characterize the 45 charge-reversal mutants in terms of their catalytic properties using yeast iso-1-ferrocycytochrome *c* as a substrate (32) and determine equilibrium constants for formation of both the 1:1 and 2:1 complexes using isothermal titration calorimetry (26, 27). If there is a single high-affinity binding site, the charge-reversal mutants that affect formation of the 1:1 complex will be clustered near the crystallographic binding site (9). On the other hand, if there are multiple binding sites with similar affinities for cytochrome *c*, then the locations of the charge-reversal mutations that affect formation of the 1:1 complex will be spread over the surface of CcP. Likewise, we will perform steady-state kinetic and calorimetric studies under conditions that facilitate formation of the 2:1 complex to determine which of the 45 mutations affect binding of the second cytochrome and determine whether these mutations cluster in a single area or are distributed over the surface of CcP.

In this report, we present initial characterization of 15 charge-reversal mutations on the front face of CcP, the surface defined by the cytochrome *c* binding site in the crystal structure of the 1:1 complex (9). The mutants have been characterized by spectroscopic measurements, determination of their hydrogen peroxide reactivity, and steady-state kinetic studies at high ionic strength where formation of only a 1:1 complex of cytochrome *c* and CcP is observed. Five of these mutants show substantial increases in the

Michaelis constant for yeast iso-1-ferrocycytochrome *c*, suggesting that these sites are involved in formation of the 1:1 complex, while 10 of the front-face mutants have essentially no effect on the Michaelis constant, indicating that these sites do not participate in binding of cytochrome *c* at high ionic strengths.

MATERIALS AND METHODS

Starting Clones and Mutagenesis. The expression system for the recombinant CcP used in this study was provided by J. Satterlee (Washington State University, Pullman, WA) (33). The rCcP gene is inserted into the multiple cloning site of Novagene vector pET24a(+) under control of the T7 promoter. The cloned gene has a sequence identical to that of mature baker's yeast CcP with the exception of the methionine codon required for bacterial expression (34). The N-terminal methionine is removed from the recombinant CcP in this expression system so that there are no N-terminal modifications in the recombinant CcP compared to baker's yeast CcP (35).

G. Pielak (University of North Carolina, Chapel Hill, NC) kindly provided the pBTR(C102T) plasmid containing the gene for yeast iso-1-cytochrome *c* (36). The yeast iso-1-cytochrome *c* gene has been altered so cysteine 102 can be replaced with a threonine residue (C102T) to prevent dimerization of the native protein via disulfide bond formation. The plasmid also contains the gene for yeast cytochrome *c* heme lyase for promotion of covalent attachment of the heme (37).

Mutations in CcP were created using Stratagene QuikChange mutagenesis kits and sequenced from 5' to 3' and from 3' to 5' to ensure that, except for the intended mutation, the protein was identical to the published sequence.

Protein Expression and Purification. Recombinant rCcP, the charge-reversal mutants, and recombinant yeast iso-1 cytochrome *c*(C102T) were expressed in *Escherichia coli* strain BL21(DE3) and isolated using published procedures (35–39).

Determination of Protein Concentration. Protein concentrations were determined by absorbance measurements using either a Hewlett-Packard model 8452A diode array spectrophotometer or a Varian/Cary model 3E spectrophotometer. Extinction coefficients and positions of the Soret maxima for the various proteins at pH 6.0 are as follows: $98 \pm 3 \text{ mM}^{-1} \text{ cm}^{-1}$ at 408 nm for yCcP, $101 \pm 3 \text{ mM}^{-1} \text{ cm}^{-1}$ at 408 nm for rCcP, $150 \pm 5 \text{ mM}^{-1} \text{ cm}^{-1}$ at 414 nm for reduced cytochrome *c*(C102T), and $118 \pm 4 \text{ mM}^{-1} \text{ cm}^{-1}$ at 408 nm for oxidized yeast cytochrome *c*(C102T) (29, 39).

Hydrogen Peroxide Concentration. Hydrogen peroxide was a reagent grade 30% (v/v) solution purchased from Aldrich Chemical Co., Inc. The concentrations of hydrogen peroxide stock solutions were determined by titration with cerium(IV) sulfate (40).

Hydrogen Peroxide Reactivity. A model SX.17MV stopped-flow spectrofluorimeter (APL Ltd., Leatherhead, England) was used to investigate the rate of reaction between the mutants and hydrogen peroxide in 0.100 M ionic strength potassium phosphate buffer (pH 6.0 and 7.5) at 25 °C. The reaction was investigated under pseudo-first-order conditions with hydrogen peroxide in excess. The enzyme was generally 1 μM , and the reaction was monitored at 424 nm. The

fraction of active enzyme was estimated by comparing the spectrum of the hydrogen peroxide-oxidized mutant to that of authentic yCcP Compound I.

Steady-State Kinetic Studies. Steady-state kinetic studies and activity measurements were performed at pH 7.5 in potassium phosphate buffers at an ionic strength of 0.100 M. Initial velocities were determined as a function of yeast iso-1-ferrocytochrome *c*(C102T) concentration (generally from 1 to 100 μ M) at a constant hydrogen peroxide concentration (200 μ M). Initial velocities were determined by measuring the change in absorbance upon oxidation of recombinant yeast iso-1-ferrocytochrome *c*(C102T) at multiple wavelengths using a Hewlett-Packard model 8452A diode array spectrophotometer. Buffer, cytochrome *c*, and enzyme were thermally equilibrated at 25 °C in the spectrophotometer; initial absorbance readings were taken, and then the reaction was initiated by addition of hydrogen peroxide. Five different wavelengths, generally chosen from the set of 314, 362, 418, 448, 468, 478, 548, 564, and 574 nm depending upon the substrate concentration, were used to calculate the initial velocity under each set of experimental conditions using eq 1.

$$\frac{v_0}{e_0} = \frac{1}{2(1 - f_{\text{ox}})\Delta\epsilon} \frac{\Delta A}{\Delta t} \quad (1)$$

The symbols in eq 1 include the initial velocity, v_0 , the total enzyme concentration, e_0 , the change in absorbance with time, $\Delta A/\Delta t$, and the difference in extinction coefficient, $\Delta\epsilon$, between oxidized and reduced cytochrome *c*. Samples of the substrate may contain small amounts of oxidized cytochrome *c* that can inhibit the reaction; f_{ox} is the fraction of oxidized cytochrome *c* in the substrate and is used to make small corrections to the initial velocity. The factor of 2 in the denominator converts cytochrome *c* turnover to enzyme turnover.

RESULTS

Locations of the Charge-Reversal Mutation Sites. The surface of CcP shown in Figure 1 contains the cytochrome *c* binding site identified using X-ray crystallography (9). We call this surface the “front face” of CcP. The cytochrome *c* binding site in Figure 1 is represented by the three residues shown in black, Asp-34, Ala-193, and Glu-290. Asp-34 and Glu-290 are within 4.2 and 4.4 Å of the cytochrome *c* surface residues, Lys-87 and Lys-73, respectively, and can potentially form salt bridges through small adjustments of the side chains (9). Ala-193 of CcP is in contact with a methyl group on the cytochrome *c* heme and is thought to be the primary electron transfer point between ferrocytochrome *c* and CcP during catalysis (9).

The locations of the mutation sites relative to the cytochrome *c* binding site can be somewhat misleading in Figure 1 due to the curvature of the CcP surface. Additional views of the mutation sites are shown in Figures S1–S3 of the Supporting Information. The figures in the Supporting Information include the bound cytochrome *c* and provide a perspective on the relationship between each of the mutation sites on CcP and the bound cytochrome *c*.

Spectroscopic Properties of the Charge-Reversal Mutants. UV–visible absorption spectroscopy was used for the initial

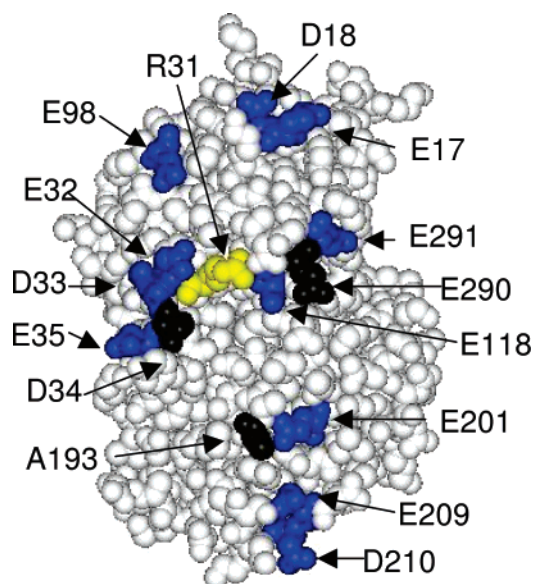


FIGURE 1: Space-filling model of the front face of CcP showing the location of residues that define the cytochrome *c* binding site (black) and the location of residues mutated in this study (blue and yellow). Asp-34, Ala-193, and Glu-290 (black) define the yeast iso-1-cytochrome *c* binding site. The aspartate and glutamate residues on the front face of CcP, excluding Asp-34 and Glu-290, are colored blue. The 13 aspartate and glutamate residues colored blue were individually mutated to lysine residues in this study. Arg-31 is colored yellow and was mutated to a glutamate residue. The one-letter abbreviations and sequence numbers are used to identify the amino acid residues. Data from ref 9; Protein Data Bank (PDB) entry 2PCC.

characterization of the mutants. Spectra were recorded at pH 6.0, the center of the pH stability region for CcP, and at pH 7.5, the pH at which the steady-state and transient-state kinetic studies were performed. The spectra for all rCcP mutants at both pH 6.0 and 7.5 are shown in Figures S4–S19 of the Supporting Information. Selected spectroscopic parameters, including the RZ value (purity number, the ratio of absorbance at the Soret maximum to the absorbance at the maximum of the protein band near 280 nm) at pH 6.0, the wavelength of the Soret maximum at both pH 6.0 and 7.5, and the ratio of the absorbance at the Soret maximum to that at 380 nm at both pH 6.0 and 7.5, are included in Table 1. Data for yCcP, CcP(MI), and rCcP are included in Table 1 for reference.

At pH 6.0, most of the charge-reversal mutants have spectra similar to that of pentacoordinate CcP, but three mutants, D37K, E209K, and D210K, have spectra characteristic of mixtures of penta- and hexacoordinate heme groups. Representative spectra for the two groups of mutants are shown in Figure 2 with the D34K mutant having a spectrum nearly identical to that of pentacoordinate CcP and the spectrum of the D37K mutant showing a significant contribution from hexacoordinate heme.

The most characteristic spectral feature of the pentacoordinate heme group in CcP is the high absorbance in the δ band near 380 nm (Figure 2). We have used the ratio of absorbancies at the Soret maximum to that at 380 nm (A_{Soret}/A_{380}) to monitor penta- and hexacoordination (Table 1). Pentacoordinate yCcP, CcP(MI), and rCcP have A_{Soret}/A_{380} values between 1.52 and 1.54. Twelve of the 15 mutants have A_{Soret}/A_{380} values ranging between 1.53 and 1.62. Three mutants exhibit significant fractions of hexacoordinate heme

Table 1: Spectroscopic Properties for rCcP and Its Charge-Reversal Mutants^a

	pH 6.0				pH 7.5	
	λ_{protein} (nm)	λ_{Soret} (nm)	RZ ^b	A_{Soret}/A_{380}	λ_{Soret} (nm)	A_{Soret}/A_{380}
yCcP ^c	282	408	1.28 ± 0.03	1.52 ± 0.04	409	1.60 ± 0.04
CcP(MI) ^c	282	408	1.28 ± 0.04	1.53 ± 0.02	410	1.67 ± 0.03
rCcP	282	408	1.31	1.54	409	1.62
Negative Cluster Mutants						
R31E	281	407	1.40	1.53	410	1.78
E32K	282	409	1.29	1.58	409	1.62
D33K	282	409	1.29	1.60	410	1.69
D34K	282	408	1.29	1.57	408	1.63
E35K	282	408	1.30	1.58	408	1.64
D37K	283	411	1.34	2.23	414	2.59
Mutants near Ala-193						
E201K	280	409	0.73	1.60	410	1.75
E209K	280	409	1.22	1.87	411	2.12
D210K	281	409	1.27	1.71	411	1.93
Mutants near Glu-290						
E118K	279	409	1.12	1.62	410	1.76
E290K	281	407	1.26	1.57	410	1.84
	281	408	1.23	1.53	411	1.87
Top Front-Face Mutants						
E17K	280	408	1.22	1.61	411	2.00
D18K	281	408	1.28	1.60	410	1.74
E98K	281	408	1.31	1.55	409	1.62

^a Absorbance values were obtained at 1 nm intervals between 240 and 700 nm in 0.10 M ionic strength potassium phosphate buffer. ^b RZ is the purity number, the ratio of the absorbance at the Soret maximum relative to the absorbance at the maximum of the protein band near 280 nm. ^c Data from studies reported in refs 39 and 44.

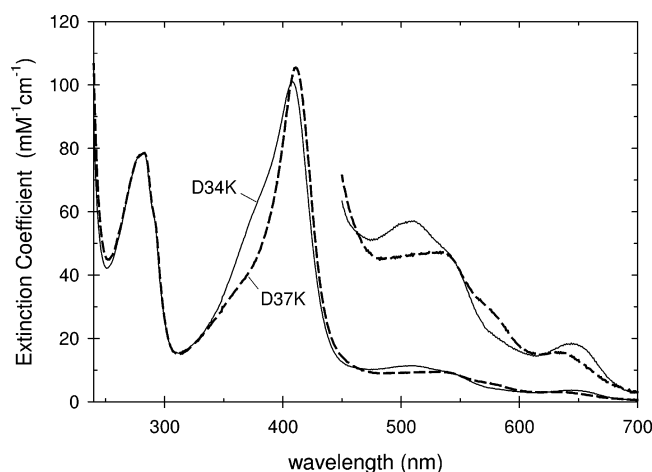


FIGURE 2: Spectra of the D34K and D37K mutants of rCcP at pH 6.0. The heme group in the D34K mutant is predominantly pentacoordinate and high-spin, while the heme in the D37K mutant is a mixture of penta- and hexacoordinate forms.

and have A_{Soret}/A_{380} values between 1.71 and 2.23. At pH 7.5, the spectra of CcP and all of the mutants show an increased contribution from hexacoordinate heme relative to that at pH 6.0 (Table 1).

Steady-State Velocity Measurements. The dependence of the steady-state velocity on the yeast iso-1 ferrocycytochrome *c* concentration was determined in 0.100 M ionic strength phosphate buffer (pH 7.5). Representative steady-state velocity plots are shown in Figure 3. Under these conditions, rCcP and 13 of the 15 mutants exhibit simple Michaelis–Menten behavior, characterized by a Michaelis constant, K_M , and a

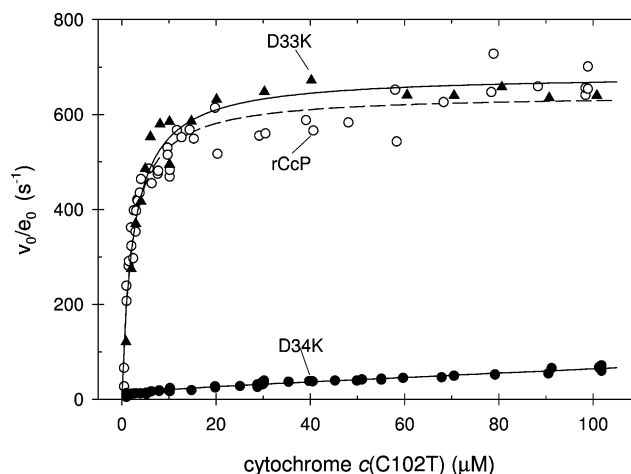


FIGURE 3: Steady-state velocities as a function of the recombinant yeast iso-1-ferrocycytochrome *c*(C102T) concentration for rCcP (○) and two of its mutants, D33K (▲) and D34K (●). Experimental conditions: 0.100 M ionic strength potassium phosphate buffer, pH 7.5, 25 °C, and 200 μM H₂O₂.

Table 2: Steady-State Parameters for rCcP and 15 Charge-Reversal Mutants^a

	major phase		minor phase	
	K_M (μM)	V_{max}/e_0 (s ⁻¹)	K_M (μM)	V_{max}/e_0 (s ⁻¹)
rCcP ^b	2.1 ± 0.2	640 ± 20		
Negative Cluster Mutants				
R31E	>100	>870	1.2 ± 0.4	30 ± 3
E32K	4.5 ± 0.5	750 ± 20		
D33K	2.4 ± 0.3	680 ± 20		
D34K	>100	>450	2.7 ± 0.8	20 ± 2
E35K	4.4 ± 0.4	760 ± 20		
D37K	82 ± 12	41 ± 3		
Mutants near Ala-193				
E201K	2.9 ± 0.9	2.8 ± 0.2		
E209K	2.8 ± 0.5	200 ± 10		
D210K	1.9 ± 0.3	470 ± 10		
Mutants near Glu-290				
E118K	51 ± 8	230 ± 20		
E290K	60 ± 11	140 ± 10		
E291K	3.8 ± 0.4	520 ± 10		
Top Front-Face Mutants				
E17K	2.3 ± 0.4	420 ± 20		
D18K	2.0 ± 0.3	380 ± 10		
E98K	3.1 ± 0.7	690 ± 40		

^a Experimental conditions: 0.100 M ionic strength potassium phosphate buffer, pH 7.5, 25 °C, and 200 μM H₂O₂. ^b Data from ref 30.

maximum velocity, V_{max}/e_0 . The steady-state parameters are collected in Table 2.

Two of the mutants, R31E and D34K, exhibit a biphasic dependence on the cytochrome *c* concentration with a minor phase characterized by V_{max}/e_0 and K_M values and a major phase in which the velocity increases linearly up to the highest substrate concentrations used in the study. The major kinetic phase for the R31E and D34K mutants can only be characterized by giving lower limits for V_{max}/e_0 and K_M values, and these are included in Table 2. On the basis of the lower limits for the V_{max}/e_0 values for the major phases of the reaction, the minor phases contribute less than 5% to the maximum activity of R31E and D34K. Steady-state velocity plots for the R31E and D34K mutants are shown in Figure S20 of the Supporting Information.

Table 3: Kinetic Parameters for the Reaction of rCcP and Its Charge-Reversal Mutants with H_2O_2^a

	fast phase		slow phase			inactive ^b
	% enzyme	k_1 ($\mu\text{M}^{-1} \text{s}^{-1}$)	% enzyme	k_2 ($\mu\text{M}^{-1} \text{s}^{-1}$)	k_3 (s^{-1})	
yCcP ^c	96	45 ± 3	0	—	—	4
CcP(MI) ^c	76	47 ± 4	12	—	11 ± 7	12
rCcP	82	48 ± 2	3	1.3 ± 0.1	—	15
Negative Cluster Mutants						
R31E	59	44 ± 3	0	—	—	41
E32K	98	43 ± 6	2	2.1 ± 0.3	—	0
D33K	94	41 ± 7	0	—	—	6
D34K	94	49 ± 4	0	—	—	6
E35K	89	44 ± 6	3	2.2 ± 0.3	—	8
D37K	10	37 ± 6	2	—	6 ± 3	88
Mutants near Ala-193						
E201K	42	43 ± 7	0	—	—	58
E209K	27	40 ± 12	0	—	—	73
D210K	58	39 ± 11	0	—	—	42
Mutants near Glu-290						
E118K	57	44 ± 8	0	—	—	43
E290K	61	51 ± 6	8	2.5 ± 0.4	—	31
E291K	40	36 ± 5	7	1.8 ± 0.3	—	53
Top Front-Face Mutants						
E17K	31	38 ± 8	10	2.0 ± 0.3	—	59
D18K	68	43 ± 14	7	2.8 ± 1.5	—	25
E98K	93	42 ± 6	5	—	22 ± 7	2

^a Experimental conditions: 0.100 M ionic strength potassium phosphate buffer, pH 7.5, and 25 °C. ^b Percent inactive enzyme estimated from the increase in the absorbance at 424 nm in the presence of a stoichiometric excess of hydrogen peroxide relative to that for yeast CcP. ^c Data from studies reported in refs 39 and 44.

Rate of Reaction between H_2O_2 and the Charge-Reversal Mutants. The mutant enzymes were further characterized by determining their rate of reaction with H_2O_2 using stopped-flow techniques at both pH 6.0 and 7.5. At pH 7.5, seven of the mutants gave monophasic kinetics with H_2O_2 , while eight of the mutants gave biphasic kinetics, with the slow phase of the reaction being at least 10 times slower than the fast phase. The rates of monophasic reaction and the fastest phase of the biphasic reactions are linearly dependent upon the H_2O_2 concentration and are due to the bimolecular reaction between the enzyme and H_2O_2 to form Compound I. The bimolecular rate constant is defined as k_1 . Values of k_1 vary between 36 ± 5 and $51 \pm 6 \mu\text{M}^{-1} \text{s}^{-1}$ for E291K and E290K, averaging $45 \pm 4 \mu\text{M}^{-1} \text{s}^{-1}$ for the 15 mutants. The average value of k_1 for the mutant enzymes is identical to the rate constant of $45 \pm 3 \mu\text{M}^{-1} \text{s}^{-1}$ determined for yCcP at pH 7.5 (41, 44). The results of the stopped-flow studies at pH 7.5 are summarized in Table 3. Kinetic data for the H_2O_2 reaction at pH 6.0 are given in Table S1 of the Supporting Information, along with a discussion of the slow phase of the H_2O_2 reaction.

Estimation of the Fraction of the H_2O_2 -Reactive Enzyme for the Charge-Reversal Mutants. During the stopped-flow studies of the reaction between the charge-reversal mutants and H_2O_2 , it became apparent that the absorbance changes at 424 nm for some of the mutants were much smaller than those for wild-type CcP. This led us to determine the absolute spectrum of the H_2O_2 -oxidized form of the mutant enzymes. Some of the mutants, such as E32K, react with H_2O_2 producing Compound I spectra essentially identical to that of yCcP Compound I (Figure 4), while other mutants, such as D37K, exhibited relatively small changes in the absorp-

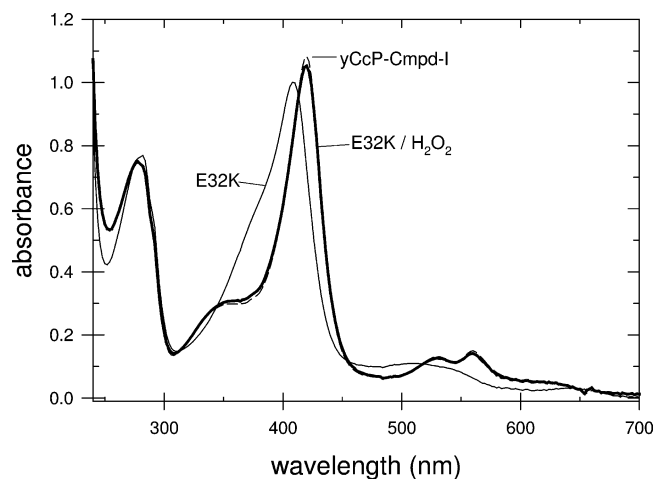


FIGURE 4: Spectra of the E32K mutant of rCcP in the absence (thin line) and presence (thick line) of a slight stoichiometric excess of hydrogen peroxide at pH 7.5. The spectrum of yCcP Compound I is also given (dashed line).

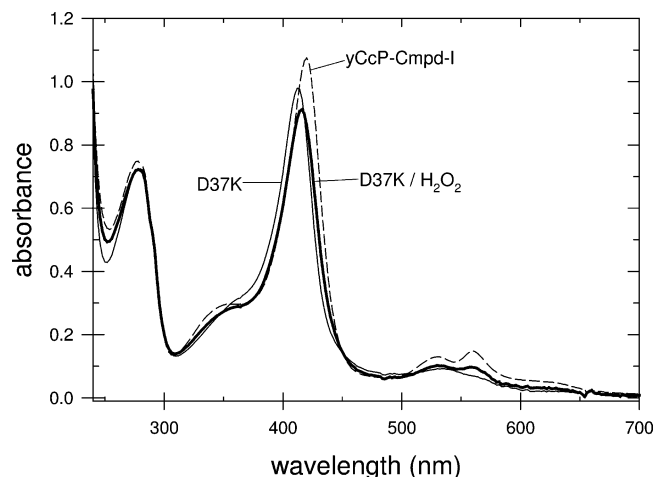


FIGURE 5: Spectra of the D37K mutant of rCcP in the absence (thin line) and presence (thick line) of a slight stoichiometric excess of hydrogen peroxide at pH 7.5. The spectrum of yCcP Compound I is also given (dashed line).

tion spectrum upon addition of a slight excess of H_2O_2 (Figure 5).

The most likely reason for the small absorbance changes for the H_2O_2 reaction in some of the mutants is that these mutants do not react stoichiometrically with H_2O_2 . An estimate of the fraction of H_2O_2 -reactive enzyme for all of the mutants was made on the basis of the assumption that the spectrum of Compound I for the mutant enzymes is identical to that of yCcP Compound I. This is a reasonable assumption since the heme ligation for Compound I of the mutants should be identical to that of yCcP Compound I. The fraction of H_2O_2 -reactive mutant was estimated on the basis of the observed absorbance change at 424 nm in the presence of a slight excess of H_2O_2 relative to the absorbance change at 424 nm calculated for the complete conversion of the mutant enzyme to the equivalent of yCcP Compound I (Figures 4 and 5). On the basis of these estimates, the fraction of H_2O_2 -reactive enzyme varies from 100% for E32K to 12% for D37K at pH 7.5. These estimates, converted to the percent of inactive enzyme, are included in the last column in Table 3. The results of equivalent estimates at pH 6 are given in Table S1 of the Supporting Information. A discussion of

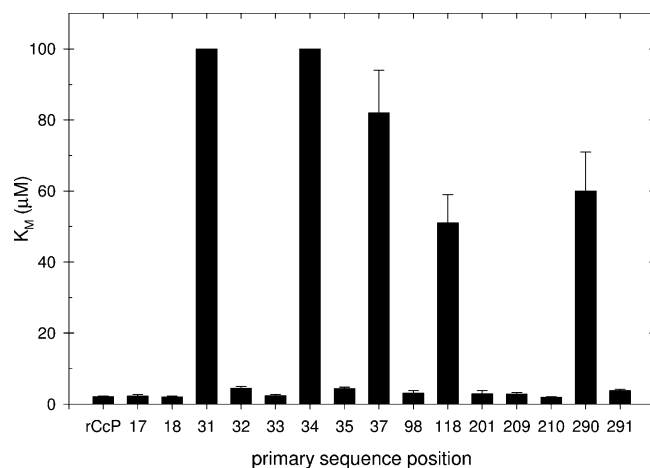


FIGURE 6: Michaelis constants, K_M , for the interaction of yeast iso-1-ferrocytochrome *c* with rCcP and with 15 charge-reversal mutants of CcP. The K_M value for rCcP is given in the left-most position followed by the K_M values of the mutants listed in order of the primary sequence position of the charge-reversal mutations. The K_M values for R31E and D34K are lower limits, while error bars are included for all other K_M values. Experimental conditions: 0.100 M ionic strength potassium phosphate buffer, pH 7.5, 25 °C, and 200 μ M H_2O_2 .

potential errors in estimating the fraction of inactive enzyme by this procedure is included in the Supporting Information.

DISCUSSION

Variation in K_M for the Charge-Reversal Mutants. We are most interested in those mutants exhibiting an altered Michaelis constant since a review of the literature shows that the K_M values for the cytochrome *c* dependence of the steady-state kinetic velocities are within experimental error of the equilibrium dissociation constants for the cytochrome *c*–CcP complex when the reactions are carried out under identical ionic strength conditions (11, 30). In this study, K_M has been used to assess the effects of charge-reversal mutations on the front face of CcP on the interaction of rCcP with yeast iso-1-cytochrome *c* (C102T). Figure 6 shows a bar graph of the K_M values for the charge-reversal mutants as a function of the primary sequence position within CcP. It is obvious that five of the mutations, R31E, D34K, D37K, E118K, and E290K, have very large effects on the binding of cytochrome *c* with K_M values that are more than 24 times larger than that of wild-type rCcP (Table 2). The properties of these five mutants will be discussed below. The K_M values for the 10 remaining charge-reversal mutants are within a factor of ~ 2 of that for wild-type rCcP, varying between 1.5 and 4.5 μ M, and these mutations are thought to have a minimal effect on the interaction between CcP and yeast cytochrome *c*.

Effect of D34K and E290K on Binding. The large increase in K_M for both D34K and E290K (Table 2) is consistent with the crystal structure of the 1:1 yeast iso-1-cytochrome *c*–CcP complex (9), which shows potential interactions between Asp-34 and Glu-290 and lysines 87 and 73 on cytochrome *c*, respectively. Previous studies have also shown that charge-neutralization or charge-reversal mutations at positions 34 and 290 decrease the affinity for cytochrome *c* (24–27). Steady-state kinetic studies with D34N and E290N mutants show 4–18-fold increases in the K_M values for yeast iso-1-ferrocytochrome *c*, depending upon the ionic strength (25), while calorimetry studies with these same mutants produce

a 4-fold decrease in binding affinity for horse cytochrome *c* in 50 mM ionic strength buffers at pH 6.0 (26). Photoinitiated electron transfer kinetic studies with E290K and zinc-substituted horse cytochrome *c* indicate that the affinity for horse cytochrome *c* is reduced 20-fold in 18 mM ionic strength buffer at pH 7.0 (24). All of these data are consistent with the involvement of both Asp-34 and Glu-290 in binding cytochrome *c* to form a 1:1 complex.

Effect of E118K on Binding. The E118K mutation increased the K_M for yeast cytochrome *c* by a factor of 24 (Table 2). Although Glu-118 is not in direct contact with the bound cytochrome (see Figure S3 of the Supporting Information), Glu-118 is near Glu-290 and located on the surface of CcP between the two protein molecules. The carboxylate side chain of Glu-118 is partially buried with OE2 hydrogen-bonded to the peptide nitrogen of Leu-289 in the interior of CcP. Mutation of Glu-118 to lysine will disrupt the hydrogen bond to Leu-289 and most likely cause the positively charged lysine side chain to rotate toward the surface of CcP. This would strengthen the electrostatic repulsion between CcP and cytochrome *c*, weakening the binding as observed. Another possible contribution to the weakening of binding could be the disruption of the Glu-118...Leu-289 hydrogen bond, altering the conformation of the polypeptide backbone from Leu-289 through the C-terminal residue, Leu-294, including Glu-290. This could affect the Glu-290–Lys-73 interaction between CcP and cytochrome *c*.

Effect of D37K on Binding. Previous studies have shown that the D37K mutation decreases the affinity for both horse and yeast cytochrome *c*, with decreases ranging from 5- to 67-fold depending upon the experimental conditions (21–23). Our steady-state data are consistent with previous studies, showing that the D37K mutation has an ~ 40 -fold decrease in affinity for yeast cytochrome *c* at pH 7.5 and an ionic strength of 100 mM. However, Asp-37 is not involved in binding cytochrome *c* in the crystal structures of the horse and yeast cytochrome *c*–CcP complexes (9). We do not believe that the influence of Asp-37 is the result of a through-space electrostatic repulsion of cytochrome binding at the crystallographic site unless there is a major reorientation of the lysine side chain in the mutant with respect to the aspartate side chain in wild-type CcP. This is based on the observation that Glu-32, Asp-33, and Glu-35 are closer to the bound cytochrome in the crystal structure than Asp-37 is, and the E32K, D33K, and E35K mutations have no significant effect on K_M (Table 2). It is also difficult to visualize an orientation of cytochrome *c* that would allow the cytochrome to simultaneously interact with Asp-37, Asp-34, and Glu-290 without also involving Glu-32, Asp-34, and Glu-35.

Another consideration that must be taken into account is the localized nature of the charge-reversal mutation effects. D34K causes a 48-fold decrease in cytochrome *c* affinity, while mutations at both adjacent residues, D33K and E35K, have almost no effect. This is also seen with E290K and E291K. The former mutation causes a 30-fold decrease in binding affinity, while the latter has essentially no effect. Assuming that the D37K mutation causes a localized effect on cytochrome *c* binding leads to the conclusion that cytochrome *c* can bind in an orientation different from that shown in the crystal structure of the 1:1 complex, one that

involves direct interaction between Asp-37 and a positively charged residue on cytochrome *c*. Alternatively, Asp-37 could be part of a second binding site that is adjacent to and/or overlaps the crystallographic site as suggested by Nocek et al. (7). Nocek et al. suggest that the second binding site is essentially that proposed by Poulos and Kraut (15) involving residues Asp-34, Asp-37, Asp-79, Gln-86, Asn-87, and Asp-217. Asp-37 is at the boundary between the crystallographic site and the Poulos–Kraut site and could affect cytochrome *c* binding at both sites. On the other hand, the data of Corin et al. (21, 22) would seem to eliminate Asp-79 and Asp-217 from the second site.

A recent NMR study (31) of the CcP–yeast iso-1-cytochrome *c* complex demonstrates that the bound cytochrome is considerably mobile but generally corroborates the crystallographic structure, indicating that cytochrome *c* resides at the crystallographic site ~70% of the time and in much more dynamic encounter complexes ~30% of the time. The encounter complexes encompass much of the CcP surface as first postulated by Northrup et al. (14). Using covalently attached paramagnetic spin-labels, Volkov et al. (31) showed that encounter complexes exist near residues 38, 200, and 288 on the surface of CcP and but not at residues 137 and 263. The spin-label probe at residue 38 should detect cytochrome *c* binding near Asp-37.

There is also the possibility that the D37K mutation decreases the affinity for cytochrome *c* by altering the surface conformation of CcP. The carboxylate side chain of Asp-37 is rotated toward the interior of CcP with the OD2 atom hydrogen bonding to NE2 of His-181 (4). His-181 is part of a hydrogen bonding network that connects residues in the distal heme pocket with those in the proximal heme pocket (43). Disruption of the hydrogen bonding network by the lysine for aspartate substitution could account for the increased concentration of hexacoordinate heme in this mutant (Table 1 and Figure 2) and, perhaps, for an altered surface conformation of the cytochrome *c* binding site. It is likely that the lysine for aspartate substitution would rotate the lysine side chain toward the surface of CcP, bringing it closer to cytochrome *c* bound at the crystallographic site, strengthening the electrostatic repulsion of bound cytochrome *c*, and decreasing the affinity at the crystallographic site.

Effect of R31E on Binding. The fifth mutation with a large effect on the Michaelis constant is the R31E mutation. Arg-31 is a positively charged residue that precedes the string of negatively charged residues, Glu-32–Asp-37. We decided to mutate Arg-31 to a glutamate in anticipation that it might increase the affinity of CcP for cytochrome *c*. Much to our surprise, the R31E mutation had a profound effect on the steady-state kinetics, generating a biphasic Michaelis–Menten plot (Figure S20 of the Supporting Information), with the major phase having a K_M value so large that only a lower limit of 100 μ M could be established. The steady-state data suggest that the R31E mutation decreases the binding affinity for cytochrome *c* by a factor of more than 50. This is difficult to understand on the basis of simple electrostatic considerations, and something more profound may be happening in the R31E mutant.

Within the yeast cytochrome *c*–CcP complex (9), the side chain of Arg-31 is hydrogen bonded to the C-terminal carboxylate of Leu-294 and is within 4.7 Å of the ϵ -amino

group of Lys-86 in cytochrome *c*. The effect of the R31E mutation appears to be the result of opposite effects, strengthening the interaction with Lys-86 but disrupting the hydrogen bond with Leu-294. Disruption of the R31E–Leu-294 interaction apparently has a more profound effect, perhaps by changing the conformation of the C-terminus and altering the Glu-290–Lys-73 interaction, which was proposed for the effects of the E118K mutation described above. Both Arg-31 and Glu-118 are involved in hydrogen bonding to the last few residues at the C-terminus of CcP and certainly play some role in stabilizing the conformation of the C-terminus. The C-terminus includes Glu-290, which can form a salt bridge with Lys-73 of cytochrome *c* in the 1:1 complex (9).

Relation to Other Studies. The data presented here are consistent with other studies that support the hypothesis that there is a unique catalytically competent cytochrome *c*–CcP complex and this complex has a structure similar to, or identical to, the structures determined by Pelletier and Kraut (9) for the yeast iso-1-cytochrome *c*–CcP and horse heart cytochrome *c*–CcP complexes in the crystalline state. This hypothesis is supported by the cytochrome *c*–CcP cross-linking studies of Poulos and co-workers (28, 45, 46) as well as those of Nakani and colleagues (29, 30). Pappa and Poulos (45) and Papa et al. (46) used site-directed mutagenesis to engineer specific cysteine residues into both yeast cytochrome *c* and CcP to covalently attach cytochrome *c* to the Pelletier–Kraut binding site through a CcP Cys-290–cytochrome *c* Cys-73 disulfide bond. The covalent complex is electron transfer competent with the covalently bound ferrocyanochrome *c* rapidly reducing the Trp-191 radical when CcP Compound I is formed. Equally important is the observation that the covalent complex has a very slow turnover rate for exogenous ferrocyanochrome *c*, indicating that if a second cytochrome *c* binding site exists, its catalytic activity is low. Nakani et al. (29) also synthesized the CcP Cys-290–cytochrome *c* Cys-73 covalent complex and showed that the residual activity of the covalent complex during steady-state turnover of exogenous ferrocyanochrome *c* is due to small amounts of rCcP(E290C) that copurified with the covalent complex and that the covalent complex is completely inactive toward exogenous ferrocyanochrome *c*, eliminating the possibility of a second catalytically active cytochrome *c* binding site on CcP. Guo et al. (28) synthesized a second covalent complex in which cytochrome *c* is bound at the Pelletier–Kraut binding site through a CcP Cys-197–cytochrome *c* Cys-81 disulfide bond. This covalent complex has rapid electron transfer from the covalently bound ferrocyanochrome *c* to the Trp-191 radical in CcP Compound I and also has a very slow rate of electron turnover of exogenous ferrocyanochrome *c*. Guo et al. (28) determined a high-resolution X-ray structure of the covalent complex and demonstrated that it closely resembles the Pelletier–Kraut structure (9). As in the Pelletier–Kraut structure, there are no direct salt bridges between charged residues in CcP and cytochrome *c* at the interface between the two proteins in the covalent complex but there are a number of water-mediated interactions between polar groups on CcP and cytochrome *c* (28).

Crane and co-workers (47–49) have provided some of the most definitive evidence that the 1:1 CcP–cytochrome *c* complexes observed in the crystalline state are the electron

transfer-active species. Crane and co-workers have crystallized several complexes between zinc porphyrin CcP (ZnC-cP) and cytochrome *c* and have shown that the rate of quenching of the zinc porphyrin triplet by the Fe(III) cytochrome *c* within the crystal is essentially the same as that in solution (47, 48). This provides strong support for the idea that the 1:1 complexes observed in the crystalline state are in fact the complexes involved in electron transfer between cytochrome *c* and CcP.

Another important aspect of the work of Crane and colleagues is the fact that they have provided crystal structures for six additional 1:1 complexes of various modified forms of CcP and cytochrome *c* (47, 48) to go along with the original yeast and horse cytochrome complexes with wild-type CcP provided by Pelletier and Kraut (9). These complexes include the yeast and horse cytochrome complexes with ZnCcP (47) and four Phe-82 mutants of yeast cytochrome *c* binding to ZnCcP (48). In the crystal structures of all eight complexes published to date, cytochrome *c* binds to the same general surface region of CcP but with some variation as originally found for the yeast and horse cytochrome *c*-CcP structures (9, 48). Kang and Crane (48) have also described a low-resolution crystal structure of a ninth complex, that between a K72S/F82Y double mutant of yeast cytochrome *c* and ZnCcP, in which the cytochrome binds in a substantially different region on the surface of ZnCcP and shows no electron transfer activity in the crystal. Interestingly, none of the cytochrome *c* molecules in any of the nine complexes is near Asp-37 in CcP, and the inhibition of binding of cytochrome *c* by the D37K mutant still remains an enigma.

Variation in V_{\max}/e_0 . At the beginning of these studies, we had anticipated that the charge-reversal mutations would modulate the binding affinity of cytochrome *c* for CcP but that cytochrome *c* would bind to the same site and have essentially the same maximum velocity as the wild-type enzyme. The observation of the large variation in the maximum velocity was unexpected. The maximum velocities actually have a larger variation than the Michaelis constants, varying >300-fold, from 2.8 s^{-1} for E201K to $>870 \text{ s}^{-1}$ for R31E (Table 2). A major part of the variation in V_{\max}/e_0 is due to the presence of H_2O_2 -unreactive forms in the mutant enzymes (Table 3). If the V_{\max}/e_0 values are corrected for the fraction of H_2O_2 -reactive enzyme, there is much less variation in the maximum turnover rates.

The fraction of H_2O_2 -inactive enzyme correlates with the amount of hexacoordinate heme as monitored by the A_{Soret}/A_{380} ratio (Tables 1 and 3). The observation that charge-reversal mutations on the surface of CcP caused changes in heme ligation in the interior of the protein was another unanticipated finding, although not central to the major objective of this study. We hypothesize that the charge-reversal mutation affects the apparent pK_a for the alkaline transition in CcP (42, 50) by modulating the binding of hydroxide ion to the heme iron, the predominant hexacoordinate heme species observed in the CcP mutants at pH 6 and 7.5 (Table 1). A discussion of the effects of the charge-reversal mutations on heme ligation and the consequent changes in H_2O_2 reactivity is included in the Supporting Information.

CONCLUSIONS

Five (R31E, D34K, D37K, E118K, and E290K) of the 15 charge-reversal mutants of CcP investigated in this study have significantly decreased affinity for yeast iso-1-ferrocytochrome *c* as monitored by the Michaelis constant in steady-state kinetic studies. Three of the five (D34K, D37K, and E290K) have previously been shown to decrease affinity for cytochrome *c* (21–27), while two (R31E and E118K) have been shown to decrease the binding affinity for the first time. Three of the mutation sites that affect cytochrome *c* binding, D34K, E118K, and E290K, are within the cytochrome *c* binding domain identified in the crystal structure of the yeast iso-1-cytochrome *c*-CcP complex (9), while two are on the periphery, R31E and D37K. The R31E mutation most likely alters the cytochrome *c* affinity by changing the conformation of the C-terminus of CcP, the region that includes Glu-290. The effect of Asp-37 on the binding of cytochrome *c* is not completely understood. It is unlikely that the effect is due to direct electrostatic repulsion of cytochrome *c* bound at the Pelletier–Kraut site (9). The D37K mutation could alter the conformation of CcP near Asp-34 and Asp-35, indirectly weakening the binding of cytochrome *c*, or Asp-37 could be part of a second binding site of the left-hand side of CcP, not yet observed in any of the crystal structures of the various cytochrome *c*-CcP complexes (9, 47, 48).

A second important result of these studies is that 10 of the 15 mutation sites on the front face of CcP (Figure 1) do not affect cytochrome *c* binding. Previously, three of these 10 sites (E32K, E35K, and E291K) have been shown not to affect cytochrome binding (24, 25). This study adds seven more sites that can be eliminated as potential cytochrome *c* binding sites for formation of 1:1 complexes.

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SUPPORTING INFORMATION AVAILABLE

Figures describing the localization of mutation sites relative to the cytochrome *c* binding site on CcP, spectra of all mutants at pH 6.0 and 7.5, hydrogen peroxide reactivity at pH 6.0, and steady-state velocity plots for selected charge-reversal mutants. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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