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Strategic roles of axial histidines in structure formation and redox regulation of tetraheme cytochrome c3

Abstract

Tetraheme cytochrome c3 (cyt c3) exhibits extremely low reduction potentials and unique properties. Since axial ligands should be the most important factors for this protein, every axial histidine of *Desulfovibrio vulgaris* Miyazaki F cyt c3 was replaced with methionine, one by one. On mutation at the fifth ligand, the relevant heme could not be linked to the polypeptide, revealing the essential role of the fifth histidine in heme linking. The fifth histidine is the key residue in the structure formation and redox regulation of a c-type cytochrome. A crystal structure has been obtained for only H25M cyt c3. The overall structure was not affected by the mutation except for the sixth methionine coordination at heme 3. NMR spectra revealed that each mutated methionine is coordinated to the sixth site of the relevant heme in the reduced state, while ligand conversion takes place at hemes 1 and 4 during oxidation at pH 7. The replacement of the sixth ligand with methionine caused an increase in the reduction potential of the mutated heme of 222-244 mV. The midpoint potential of a triheme H52M cyt c3 is higher than that of the wild type by ~50 mV, suggesting a contribution of the tetraheme architecture to the lowering of the reduction potentials. The hydrogen bonding of Thr24 with an axial ligand induces a decrease in reduction potential of ~50 mV. In conclusion, the bis-histidine coordination is strategically essential for the structure formation and the extremely low reduction potential of cyt c 3. © 2008 American Chemical Society.

Keywords

tetraheme, regulation, redox, c3, formation, axial, cytochrome, strategic, structure, roles, histidines

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Strategic Roles of Axial Histidines in Structure Formation and Redox Regulation of Tetraheme Cytochrome c_3 [†]

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ABSTRACT: Tetraheme cytochrome c_3 (cyt c_3) exhibits extremely low reduction potentials and unique properties. Since axial ligands should be the most important factors for this protein, every axial histidine of *Desulfovibrio vulgaris* Miyazaki F cyt c_3 was replaced with methionine, one by one. On mutation at the fifth ligand, the relevant heme could not be linked to the polypeptide, revealing the essential role of the fifth histidine in heme linking. The fifth histidine is the key residue in the structure formation and redox regulation of a c -type cytochrome. A crystal structure has been obtained for only H25M cyt c_3 . The overall structure was not affected by the mutation except for the sixth methionine coordination at heme 3. NMR spectra revealed that each mutated methionine is coordinated to the sixth site of the relevant heme in the reduced state, while ligand conversion takes place at hemes 1 and 4 during oxidation at pH 7. The replacement of the sixth ligand with methionine caused an increase in the reduction potential of the mutated heme of 222–244 mV. The midpoint potential of a triheme H52M cyt c_3 is higher than that of the wild type by ~50 mV, suggesting a contribution of the tetraheme architecture to the lowering of the reduction potentials. The hydrogen bonding of Thr24 with an axial ligand induces a decrease in reduction potential of ~50 mV. In conclusion, the bis-histidine coordination is strategically essential for the structure formation and the extremely low reduction potential of cyt c_3 .

Cytochrome c_3 (cyt c_3)¹ isolated from sulfate-reducing bacteria is a periplasmic electron transport protein whose molecular mass is typically ~14 kDa, and it possesses four hemes attached to a single polypeptide. It is involved in sulfate respiration, which is the prototype of the oxygen

respiration system. Because of its unusual properties, it is classified as a class III c -type cytochrome (1). The roles of the four hemes in regulation of the electron transport pathway in sulfate respiration are biologically relevant. The three-dimensional structures of ferrous and ferric cyt c_3 from various strains have been determined by means of X-ray crystallography and nuclear magnetic resonance spectroscopy (NMR) (2 and references cited therein). Although the degree of sequence homology among cyt c_3 species is not high (~30%), the heme architecture and the fold of the protein are conserved. The fifth and sixth ligands of each heme are histidines (His or H). Cyt c_3 has a variety of interesting properties (3). The most important of them is its extremely low reduction potentials. This can be attributed to the long history of sulfate respiration during biological evolution. The low potentials mean that cyt c_3 carries highly reductive heme irons. Its tetraheme architecture was suggested to be important for regulating redox processes (3). It is also known that a dried film of it becomes electric conductive on reduction (4). Therefore, this protein is intriguing not only from a biological point of view but also from physical and chemical ones.

Different from the monoheme cyt c involved in oxidative phosphorylation, cyt c_3 has two types of heme-binding motifs, namely, -Cys-(Xaa)₂-Cys-His- and -Cys-(Xaa)₄-Cys-His- (CXXXXCH), where Cys (C) and Xaa (X) are cysteine and

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¹ Abbreviations: cyt c_3 , cytochrome c_3 ; DvMF, *Desulfovibrio vulgaris* Miyazaki F; DvH, *D. vulgaris* Hildenborough; CD, circular dichroism; MPOH, 3-mercapto-1-propanol; NMR, nuclear magnetic resonance; NaP_i, sodium phosphate buffer; L₅ⁿ or L₆ⁿ mutant, mutant with the fifth or sixth ligand, respectively, of heme n changed; Ccm, cytochrome c maturation system; SAM, self-assembled monolayer.

any amino acid, respectively, and His is the fifth ligand. A classification of the cyt c_3 subfamilies based on the combination of heme binding motifs has been reported (5). The reduction potentials of the four hemes were found to be lower for the 2-4-2-4 subfamily than for other subfamilies. The potentials are in the ranges of -200 to -380 mV (6–10) and -90 to -355 mV (11, 12), respectively. The name of each subfamily is based on the number of Xaa residues in the four heme-binding motifs from the N-terminus to the C-terminus. Cyt c_3 from *Desulfovibrio vulgaris* Miyazaki F (*DvMF*), the subject of this work, belongs to the 2-4-2-4 subfamily. Its macroscopic and microscopic reduction potentials have been determined, using differential pulse polarography and NMR (13, 14). Its oxidized (15, 16) and reduced (2) structures have been reported. Since an efficient overexpression system for cyt c_3 has been established (17), we have carried out a series of mutation works (3). This is one of the best-characterized cyt c_3 species. Using this system, we have tried to elucidate the significance of the bis-imidazole coordination and the tetraheme architecture. Although it is well-known that the reduction potential of the His–His coordinations is much lower than that of methionine (Met or M)–His coordinations (18), it is important to determine the actual values to understand the redox properties of cyt c_3 . For this purpose, the axial His residues were replaced with Met one by one in this work. Mutations at the sixth axial ligand in *Desulfovibrio vulgaris* Hildenborough (*DvH*) have been reported previously (7, 8). However, extensive work has not been carried out because of the difficulty with overexpression. In particular, heme-specific reduction potentials were not investigated. This has been done in this work. Furthermore, we could replace His not only at the sixth but also at the fifth coordination site. This work will also provide a complete view of the roles of the aromatic residues in cyt c_3 , because we have mutated all noncoordinated aromatic residues in *DvMF* cyt c_3 (5, 16). The results revealed that the axial histidine coordination is essential not only for realizing the extremely low reduction potentials but also for the structure formation of cyt c_3 .

MATERIALS AND METHODS

Site-Directed Mutagenesis. All mutations were introduced into the pKFC3k plasmid, which contains the *DvMF* cyt c_3 gene (16). Site-directed mutagenesis was performed with a Mutan-Super Express Km kit (Takara Bio Inc.), and synthetic oligonucleotides were purchased from QIAGEN. The synthetic oligonucleotides used in this study are summarized in Table 1. The mutations were confirmed by nucleotide sequencing with an ABI PRISM 310 genetic analyzer (Perkin-Elmer), using a DNA sequencing kit, BigDye Terminator Cycle Sequencing Ready Reaction version 1.0 (Applied Biosystems), and two M13 primers, RV-N and M4 (Takara Shuzo Co., Ltd.).

Cultivation and Purification. Mutated cyt c_3 species were expressed in *Shewanella oneidensis* TSP-C transformed with a mutated plasmid, as described previously (17). The [$C_e^2H_3$]Met proteins were obtained in a medium comprising 31 mM sodium lactate, 31 mM disodium fumarate, 11 mM glucose, 10 mM sodium citrate, 93 mM NH_4Cl , 50 mM Na_2HPO_4 , 22 mM KH_2PO_4 , 12 mM $NaCl$, 500 μM $MgSO_4$, 100 μM $CaCl_2$, 20 μM $FeSO_4$, 1 μM $MnCl_2$, 20 mg/mL

Table 1: Synthetic Oligonucleotides Used for Site-Directed Mutagenesis^a

mutation (ligand ^b)	synthetic oligonucleotide
H34M (L_5^1)	5'(289)-GGC GAC TGT <u>ATG</u> CAC CCG GTC AAC-3'(312)
H52M (L_5^2)	5'(336)-C GCC ACC GCC GGT TGC <u>ATG</u> GAC AAC ATG G-3'(364)
H83M (L_5^3)	5'(433)-TGC GTG GGC TGC <u>ATG</u> CTT GAA ACC GCC-3'(459)
H106M (L_5^4)	5'(502)-GGC TCC AAG TGC <u>ATG</u> AGC TAA GAA TCC-3'(528)
H22M (L_6^1)	5'(250)-GTG GTC TTC AAC <u>ATG</u> TCG ACC CAC AAG-3'(276)
H35M (L_6^2)	5'(286)-TGT GGC GAC TGT CAT <u>ATG</u> CCG GTC AAC GCC-3'(315)
H25M (L_6^3)	5'(259)-AAC CAC TCG ACC <u>ATG</u> AAG GCC GTG AAG-3'(285)
H70M (L_6^4)	5'(393)-C TAC CAC GCC ATG <u>ATG</u> GAC AAG GGC ACC-3'(420)
H34K (L_5^1)	5'(285)-G TGT GGC GAC TGT <u>AAG</u> CAC CCG GTC AAC GG-3'(314)
H34Q (L_5^1)	5'(285)-G TGT GGC GAC TGT <u>CAG</u> CAC CCG GTC AAC GG-3'(314)
H34Y (L_5^1)	5'(285)-G TGT GGC GAC TGT <u>TAC</u> CAC CCG GTC AAC GG-3'(314)
T24V	5'(253)-GTC TTC AAC CAC TCG <u>GTC</u> CAC AAG GCC GTG-3'(282)

^a The sequencing numbers in parentheses are the sequence numbers according to ref 16. ^b L_i^n stands for the i th ligand at heme n . H, M, K, Q, Y, T, and V stand for histidine, methionine, lysine, glutamine, tyrosine, threonine, and valine, respectively.

adenosine, 20 mg/mL guanosine, 20 mg/mL cytidine, 20 mg/mL thymidine, 20 mg/mL uridine, 20 mg/mL thiamine hydrochloride, 20 mg/mL α -biotin, and an amino acid mixture. The amino acid mixture (all per liter of culture) consisted of 300 mg of Ala, 200 mg of Arg, 400 mg of Asn, 400 mg of Asp, 50 mg of Cys, 400 mg of Gln, 1000 mg of Glu, 300 mg of Gly, 200 mg of His, 400 mg of Ile, 400 mg of Lue, 450 mg of Lys, 200 mg of Phe, 400 mg of Pro, 500 mg of Ser, 200 mg of Thr, 100 mg of Trp, 300 mg of Tyr, 300 mg of Val, and 150 mg of [d_3 -methyl]Met (98% d_3 -methyl, Cambridge Isotope Laboratories, Inc.). The pH was adjusted to 7.5. To obtain each mutant protein for this work, a 24–36 L, total, cell culture was used. The mutated cyt c_3 species were purified by the procedure used for the wild type (19). To prepare ferrous cyt c_3 forms, [NiFe] hydrogenase was purified from *DvMF* cells according to the reported method (20).

Mass Spectrometry. The mass number was determined with a matrix-assisted laser desorption ionization time-of-flight (MALDI-TOF) mass spectrometer (Autoflex, Bruker Daltonics). Cyt c_3 solutions were dialyzed against ultrapure water. Aliquots (1 μL) were deposited on an MTP 384 target plate of polished steel TF (Bruker Daltonics) followed by deposition of 1 μL of matrix, which comprised 10 mg/mL sinapinic acid (Bruker Daltonics) in 50% acetonitrile and 0.1% trifluoroacetic acid. Mass data were calibrated with Protein Calibration Standard I (Bruker Daltonics).

Circular Dichroism Measurements. Circular dichroism (CD) spectra were acquired with a Jasco J-720WI CD spectropolarimeter at 30 °C. The sample concentration for the far-UV region (190–250 nm) was 5 μM in 30 mM sodium phosphate buffer (NaP_i) (pH 7.0). The solvent spectrum obtained under the same conditions was subtracted.

CD spectra were recorded using a quartz cell with a path length of 1 mm.

Electrochemical Measurements. Differential pulse polarograms were recorded at 30 °C with a Perkin-Elmer 394 digital electrochemical trace analysis system (Perkin-Elmer) using a dropping mercury electrode and a Ag/AgCl reference electrode. The modulation amplitude, sweep rate, and drop time were 20 mV, 2 mV s⁻¹, and 2 s, respectively. To obtain macroscopic reduction potentials, the polarograms were fitted to an analytical equation for four and three consecutive one-electron reversible electrode reactions (6).

The reduction potentials of Met-coordinated hemes were determined with a 3-mercapto-1-propanol (MPOH)-coated gold electrode, using a computer-controlled three-electrode potentiostat (Hokuto Denko, HZ-5000). MPOH (Tokyo Chemical Ind. Ltd.) was used for preparation of the electrode without further purification. A mica sheet (Nilaco) was baked at 580 °C prior to the vapor deposition and maintained at 580 °C during the positioning of Au (99.99%). The substrates were then annealed at 530 °C for 8 h at ambient atmosphere to obtain large Au(III) terraces on the surface (21). SAM-MPOH-coated gold substrates were prepared by immersing Au-deposited mica substrates in a 1 mmol dm⁻³ ethanol solution of MPOH for 24 h. For voltammetry measurements, a substrate with SAM-MPOH was mounted at the bottom of a cone-shaped cell using an elastic O-ring (22). The surface area of the electrode was estimated to be 0.126 cm² from the diameter of the O-ring.

A solution of either 30 mM NaPi (pH 7.0) or 30 mM borate buffer (pH 9.0) in a cell was deaerated via Ar bubbling for 15 min. Ten milligrams of cyt c_3 was then added to the cell (2 mg/mL), followed by dissolution with further Ar bubbling for less than 1 min. The reference electrode used was a Ag/AgCl/saturated KCl one equipped with a Vycor glass for the liquid junction. A platinum wire was used as the counter electrode. All voltammetry measurements were made at 21 ± 1 °C. At first, cyclic voltammograms were recorded at 100, 50, and 20 mV/s. Reversibility was confirmed for those at 50 and 20 mV/s. Then, polarograms were recorded at 20 mV/s. Reduction potentials are referred to the standard hydrogen electrode.

Nuclear Magnetic Resonance Measurements. Samples [1 mM cyt c_3 dissolved in 30 mM NaPi (p²H 7.0) or in 30 mM borate buffer (p²H 9.0)] were prepared as described previously (23). Nuclear magnetic resonance (NMR) spectra at 500 and 600 MHz were recorded at 303 K with AVANCE DRX-500 and DRX-600 NMR spectrometers, respectively (Bruker Biospin). The size of the data set was 4096 × 512. The assignment of each heme in the fully reduced state and heme methyl signals in five macroscopic oxidation states was carried out as previously reported (13, 14, 24). Chemical shifts are presented in parts per million relative to 2,2-dimethyl-2-silapentane-5-sulfonate (DSS) as an internal reference.

Crystallization and Structure Determination. Crystals of H25M and T24V cyt c_3 were grown at 10 °C according to the method reported for the wild type (25). Here, H25M stands for the replacement of His25 with Met. T and V are threonine (Thr) and valine (Val), respectively; 150 μL of an H25M cyt c_3 solution [20 mg/mL, 10 mM borate buffer (pH 9.0)] containing 50% (v/v) 2-methyl-2,4-pentanediol (MPD) was equilibrated against 4 mL of the same buffer containing

Table 2: Summary of X-ray Crystallographic Data

	H25M cyt c_3	T24V cyt c_3
X-ray source	Rigaku RA-Micro7	BL41XU, SPring-8
wavelength (Å)	1.541	0.71000
space group	$P2_12_12_1$	$P2_12_12_1$
unit cell parameters (Å)	$a = 51.94$, $b = 67.16$, $c = 34.57$	$a = 52.43$, $b = 67.54$, $c = 34.41$
resolution range (outer shell) (Å)	11.50–1.50 (1.58–1.50)	10–1.00 (1.04–1.00)
no. of measured reflections	122630	465200
no. of unique reflections	18745	61485
completeness (%)	94.1 (89.9)	99.3 (98.9)
R_{merge}	0.068 (0.243)	0.058 (0.303)
$\langle I/\sigma(I) \rangle$	8.9	13.9
R	0.178	0.109
R_{free}	0.195	0.140

70% MPD. In the case of T24V cyt c_3 , 10 mM sodium phosphate buffer (pH 7.0) was used.

Diffraction experiments were carried out at 100 K for H25M and T24V cyt c_3 . The diffraction data set for H25M cyt c_3 was collected on an RAXIS-VII instrument (Rigaku Co. Ltd.) using Cu K α radiation generated by a rotating anode (RA-Micro7). The data set for T24V cyt c_3 was collected using a synchrotron radiation source ($\lambda = 0.71$ Å) at the BL41XU beamline with a Quantum315 detector system at SPring-8. The crystal-to-detector distance was 200 mm (Quantum315) or 75 mm (RAXIS-VII) with an oscillation range per image of 1.0°. Integration of the reflections was performed using HKL2000 (26) for T24V cyt c_3 and MOSFLM (27) for H25M cyt c_3 . Initial phase calculation and refinement for both mutants were carried out with CNS (28) using the atomic coordinates of the wild-type cyt c_3 without water molecules (16) as a starting model. The structures of T24V and H25M cyt c_3 , including water molecules, were refined with isotropic B factors using CNS. The structure of T24V cyt c_3 was further refined with anisotropic B factors using SHLXL (29). The R values for the final models of H25M and T24V cyt c_3 were 0.178 and 0.122, respectively. The X-ray crystallographic data are summarized in Table 2. The root-mean-square deviation (rmsd) values for the backbone atoms were calculated with Swiss-Pdb Viewer 3.7 (30). The coordinates and structure factors for T24V and H25M cyt c_3 have been deposited in the Protein Data Bank as entries 2EWK and 2YXC, respectively.

RESULTS

Effect of Mutation of an Axial Ligand on Cyt c_3 Expression. To elucidate the role of an axial ligand, every axial His of DvMF cyt c_3 was replaced with Met, one by one. The effects of the replacements were characterized by the yield, the ratio of absorbance at the α -band in the reduced state to that at 280 nm in the oxidized state ($A_{\alpha}^{\text{red}}/A_{280}^{\text{ox}}$), and the mass number, all results being summarized in Table 3. The yield of a mutant of the fifth ligand (L₅ mutant) was lower than that of the L₆ ones except for the heme 4 mutant. The heme content of each mutant was estimated on the basis of $A_{\alpha}^{\text{red}}/A_{280}^{\text{ox}}$. Although it was almost the same as that of the wild type for every L₆ mutant, it was reduced to 60–85% for every L₅ mutant. Additionally, the mass numbers of the L₅ mutants were reduced by ~610, which accounts for that of

Table 3: Characterization of Mutant Cyt c_3 Species as Axial Ligands

sample (ligand ^a)	yield (mg/L of culture)	$A_{\alpha}^{\text{red}}/A_{280}^{\text{oxb}}$	mass number ^c
wild type	9.01	3.00	14056 (+12)
H34M (L ₅ ¹)	0.04	1.94 (65%)	13440 (−610)
H52M (L ₅ ²)	0.88	2.55 (85%)	13437 (−613)
H83M (L ₅ ³)	0.03	2.31 (77%)	13438 (−612)
H106M (L ₅ ⁴)	0.79	2.43 (81%)	13437 (−610)
H22M (L ₆ ¹)	0.85	2.88 (96%)	14066 (+16)
H35M (L ₆ ²)	4.76	2.98 (99%)	14048 (−2)
H25M (L ₆ ³)	4.76	2.92 (97%)	14046 (−4)
H70M (L ₆ ⁴)	0.71	2.94 (98%)	14053 (+3)
H34Q (L ₅ ¹)	0.01	1.80 (60%)	13437 (−610)
H34K (L ₅ ¹)	0.02	1.92 (64%)	13437 (−610)
H34Y (L ₅ ¹)	0.02	1.94 (65%)	13472 (−610)

^a L_iⁿ stands for the *i*th ligand at heme *n*. ^b The heme content estimated as the absorbance ratio is shown in parentheses, that for the wild type being 100%. ^c The difference (expected mass − observed mass) is given in parentheses. The standard errors are ±1%.

a single heme. It can be concluded that one heme is not linked to the polypeptide chain in the L₅ mutants, although the apparent heme content is 60–80%.

Since the number of covalent bonds between the coordinated N_{ε2} and C_α atoms of His (four bonds) is larger by one than that between the S_δ and C_α atoms of Met (three bonds), H34Q (N_ε or O_ε), H34K (N_ε), and H34Y (O_η) mutations were also introduced to maintain the chain length at the fifth coordination site (the expected coordinated atom being given in parentheses). Q, K, and Y stand for glutamine, lysine, and tyrosine, respectively. They exhibited similar results, which can be seen in Table 3. This indicates that no amino acid residues but His at this site can introduce covalent bonds between heme and Cys residues. Namely, only His in the heme-binding motif is recognized as the fifth ligand by the cytochrome *c* maturation system (Ccm) in *Shewanella* cells. This coincides with the observation that the *Escherichia coli* Ccm recognized only imidazole as the fifth ligand (31). In the case of *E. coli*, His in the CXXCH motif created in cyt *b*₅₆₂ was replaced with Met, arginine, or Lys. In the case of a multiheme cytochrome *c* such as a cyt *c*₃, the maturation of the whole protein is not blocked by the inhibition of heme linking at one site, suggesting that the heme linking process at each heme-binding motif is independent. However, the lack of one heme significantly suppresses the efficiency of the protein production. The heme content should be 75% of that of the wild type, if L₅ mutants have only three hemes. However, it is in the range of 60–85%, suggesting the presence of a *b*-type heme, as in the case of *E. coli* (31). A lower value such as 60–65% can be ascribed to broadening of the band due to structural inhomogeneity. Thus, the preparations of L₅ mutants are inhomogeneous in terms of heme number and structure.

Structural characterization of all mutants was carried out by CD spectroscopy. The spectra are illustrated in Figure 1. There are only small changes in the secondary structure contents for the L₆ mutants, in accordance with the previous work on DvH (32). In contrast, the secondary structure contents of the L₅ mutants were distinctly reduced. Nevertheless, the polypeptide could fold into a globular structure even with three hemes. Further characterizations of the L₅ mutants were not carried out because of their inhomogeneity in heme number and structure.

Coordination Structure at the Mutated Sites. To examine the effect of a mutation on its structure, we tried to crystallize

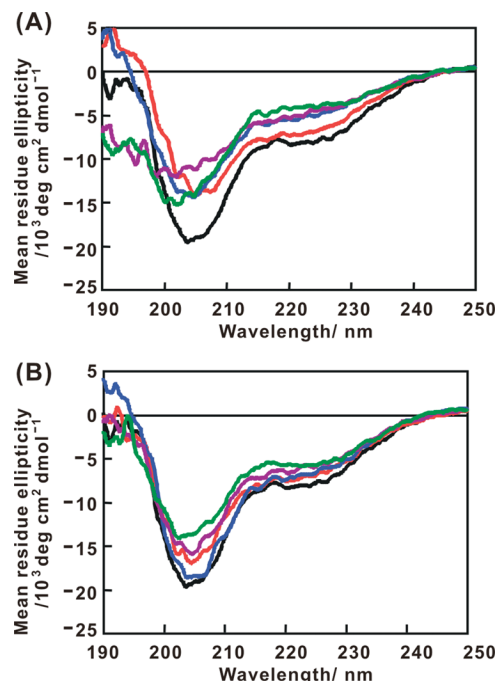


FIGURE 1: Circular dichroism spectra of H34M, H52M, H83M, and H106M cyt *c*₃ (A) and H22M, H35M, H25M, and H70M cyt *c*₃ (B) at pH 7.0. (A and B) Mutations at the fifth and sixth coordination sites, respectively. Black, red, blue, purple, and green show data for the wild type and heme 1–4 axial ligand mutants, respectively.

every L₆ mutant. Since only ferric H25M(L₆³) cyt *c*₃ was successfully crystallized at pH 9.0, its structure was determined (green in Figure 2A). L₆³ stands for the L₆ mutant of heme 3. There was no significant difference in the polypeptide folding in comparison with that of the wild type (PDB entry 1J00; red in Figure 2A). The rmsd value for the backbone atoms was 0.35 Å. The effect of the axial ligand replacement at heme 3 was local. The sixth ligand of heme 3 was S_δ of Met25, as expected (Figure 2B). The HisN_{ε2}–iron (Fe) and MetS_δ–Fe distances at heme 3 were 1.99 and 2.34 Å, respectively. The S_δ–Fe length is consistent with those found in other cyt *c* species with Met–His coordinations. For instance, those for human and horse cyt *c* and *Saccharomyces cerevisiae* iso-1-cyt *c* are 2.38, 2.66, and 2.19 Å, respectively (PDB entries 1J3S, 1AKK, and 1YIC, respectively). It can be concluded that S_δ is the sixth ligand of H25M cyt *c*₃.

Then, coordinated atoms in the sixth axial ligands of other mutants were identified on the basis of one-dimensional ¹H NMR spectra of the ferrous forms at pH 9.0 (Figure 3). Since the coordinated Met80 C_εH₃ signal of recombinant human cytochrome *c* appears at −3.28 ppm (BioMagResBank entry 5406), the signals indicated by black arrows in Figure 3 are candidates for the C_εH₃ signal of the coordinated Met. To confirm the assignment, Met was specifically deuterated using [C_ε²H₃]Met. The intensity of the tentatively assigned signal was significantly diminished on deuteration for every mutant (colored red in Figure 3). The Met coordination in the crystal structure of H25M cyt *c*₃ at pH 9.0 also justifies the assignment of the coordinated Met C_εH₃ signals. This is also the case for the reduced mutants at pH 7.0, because similar signals were observed at pH 7.0 (spectra not shown).

¹H NMR spectra for the heme methyl signal regions of the fully oxidized cyt *c*₃ species are shown in Figure 4.

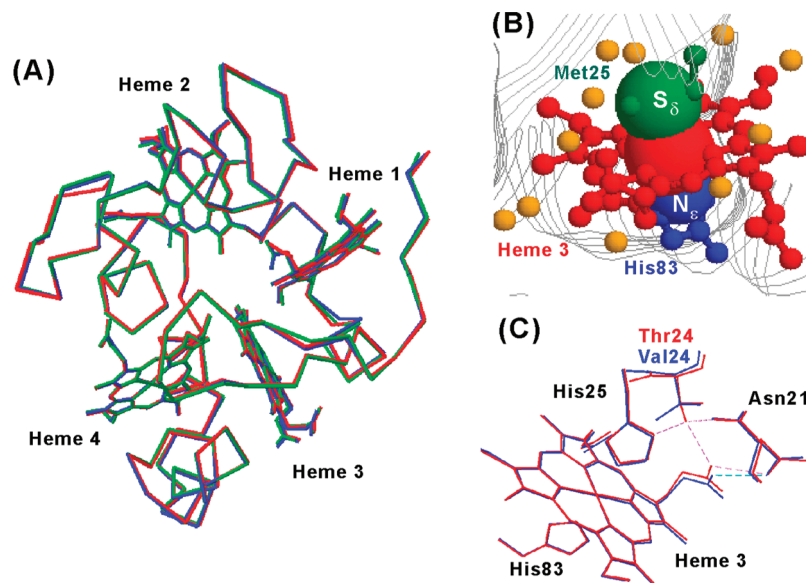


FIGURE 2: (A) Overall structures (backbone) and heme architectures of the wild-type (red), H25M (L_6^3) (green), and T24V (blue) cyt c_3 in crystal. (B) Crystal structure of H25M cyt c_3 around heme 3. The backbone, heme 3, His83 (fifth ligand), Met25 (sixth ligand), and water molecules within 10 Å of the Fe of heme 3 are represented as gray lines, and red, blue, green, and orange spheres, respectively. This figure was prepared with RasMol (47). (C) Comparison of the crystal structures of the wild-type (red) and T24V (blue) cyt c_3 around heme 3. The hydrogen bond in the wild type and that in the T24V mutant are colored pink and light blue, respectively. Panels A and C were prepared with Swiss-PdbViewer (30).

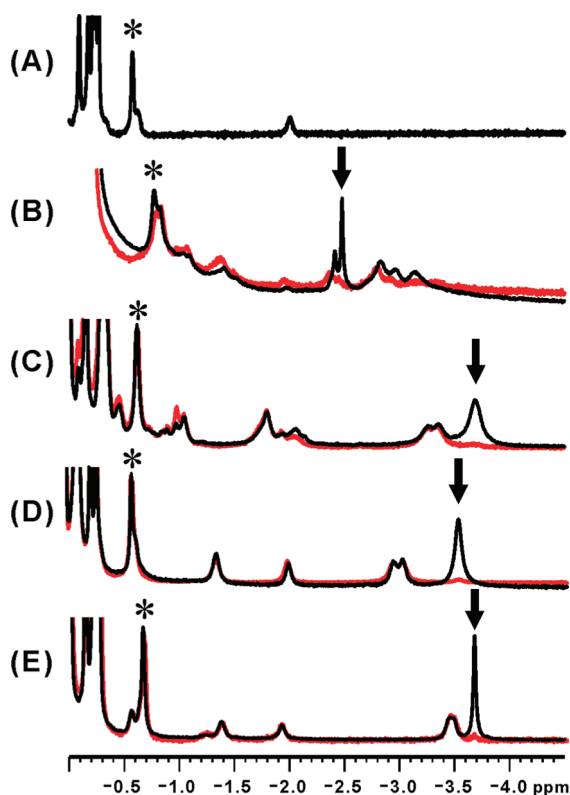


FIGURE 3: ^1H NMR spectra of wild-type (A), H22M (B), H35M (C), H25M (D), and H70M (E) ferrous cyt c_3 in a low-frequency region at pH 9.0, 600 MHz, and 303 K. All but spectrum A are L_6 mutants of hemes 1–4, respectively. The spectra of $[\text{C}_\epsilon^2\text{H}_3]\text{Met}$ -labeled proteins are superimposed on those of the relevant mutants in red. Asterisks and arrows indicate Ala68 βCH_3 and Met $\text{C}_\epsilon\text{H}_3$ signals, respectively.

Typical low-spin spectra were observed at pH 9.0, as expected from Figure 3. In contrast, broad peaks appeared in the region from 35 to 60 ppm and at 39 ppm at pH 7.0 for H22M (L_6^1) and H70M (L_6^4) cyt c_3 , respectively. This

could be assigned to a high-spin or intermediate-spin state. A much smaller peak was also observed at ~ 40 ppm for the H35M (L_6^2) mutant. Although the low-spin state also prevails at pH 7.0, the molecular species are heterogeneous. The coordination structure can be examined in the lowest-frequency region (Figure 5). The coordinated His $\text{C}_{\delta 2}$ proton signals appear in this region (33). The coordinated Met $\text{C}_\epsilon\text{H}_3$ signal of horse cytochrome c appears at -27.6 ppm (BioMagResBank entry 317) (34) due to a paramagnetic effect. In Figure 5, the spectra of nonlabeled (black) and $[\text{C}_\epsilon^2\text{H}_3]\text{Met}$ -labeled (red) L_6 mutants were superimposed. Coordinated Met $\text{C}_\epsilon\text{H}_3$ signals can be seen at approximately -5.5 and -11.5 ppm for H25M and H35M ferric cyt c_3 , respectively, at pH 7.0, and only for the H25M one at pH 9.0. The weaker paramagnetic effect, compared to that for horse cyt c , indicates that the coordination in solution is weaker in spite of a normal coordination bond length for H25M cyt c_3 in the crystal. Since the ligand at every mutated site is Met in the reduced state, the ligand should be replaced during oxidation for H22M, H35M, and H70M cyt c_3 at pH 9.0, and for H22M and H70M cyt c_3 at pH 7.0. In view of the low-spin state, a non-Met ligand in the oxidized state should be a strong one. However, there is no candidate amino acid residue in the proximity. An obvious candidate is a hydroxide ion. A small fraction of mutated hemes would be in the high-spin or intermediate-spin state, judging from Figure 4. The rate of exchange between different states is slow on the NMR time scale.

Redox Properties of L_6 Mutant Cyt c_3 . Two-dimensional exchange spectra of L_6 mutants at different oxidation stages were recorded (spectra not shown). The assignment of heme methyl signals in the five macroscopic oxidation states was carried out as previously reported (13, 14, 24). The reduction fractions (R_i^j , where i and j are the heme number and j th reduction step, respectively) were calculated using the chemical shifts of signals 2- and 18- CH_3 for heme 1, 7- and

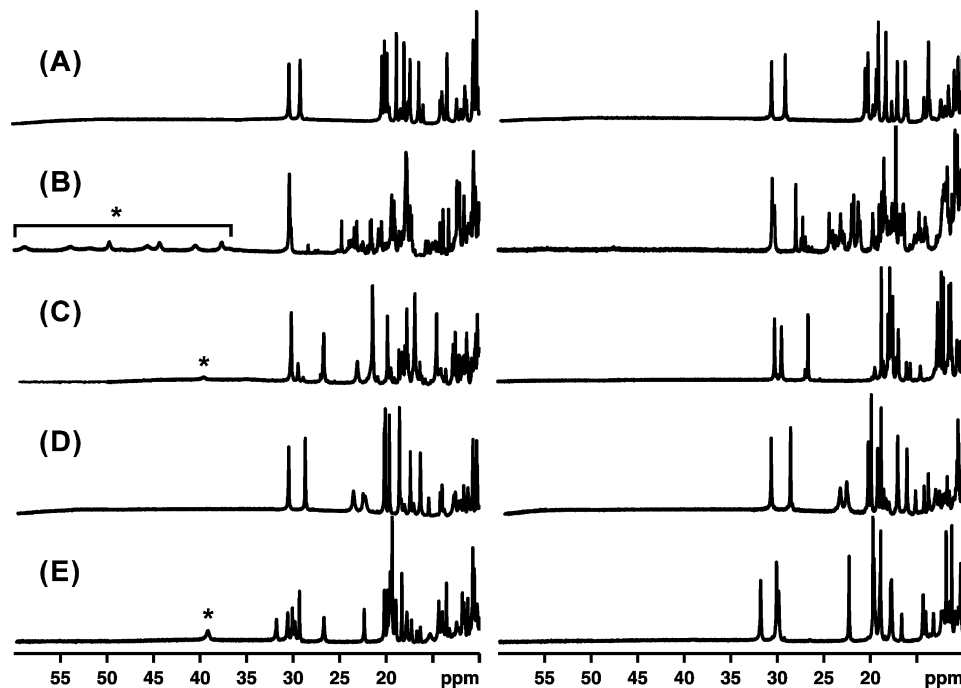


FIGURE 4: ^1H NMR spectra of wild-type (A), H22M (B), H35M (C), H25M (D), and H70M (E) ferric cyt c_3 at 600 MHz and 303 K: (left) p ^2H 7.0 and (right) p ^2H 9.0. Only the fingerprint regions for heme methyl groups are presented. Asterisks indicate putative high- or intermediate-spin species.

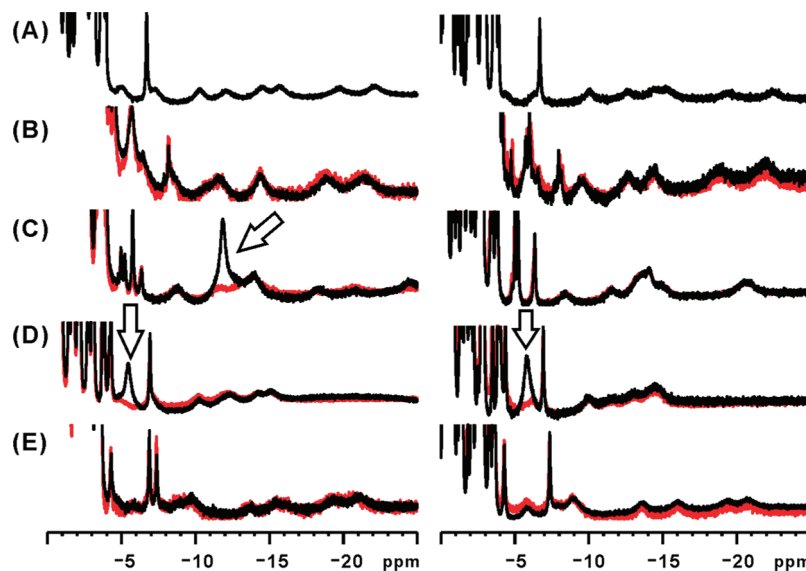


FIGURE 5: ^1H NMR spectra of wild-type (A), H22M (B), H35M (C), H25M (D), and H70M (E) ferric cyt c_3 in a low-frequency region at 600 MHz and 303 K: (left) p ^2H 7.0 and (right) p ^2H 9.0. The spectra of $[\text{C}_\epsilon^2\text{H}_3]\text{Met}$ -labeled proteins are superimposed on those of the relevant mutants in red. Arrows indicate Met $\text{C}_\epsilon\text{H}_3$ signals.

18- CH_3 for heme 2, 12- and 18- CH_3 for heme 3, and 2- and 18- CH_3 for heme 4 (5), which are summarized in Table S1 of the Supporting Information. The reduction fractions of H22M (L_6^1) and H70M (L_6^4) cyt c_3 were obtained at only pH 9.0 because there were at least two molecular species at pH 7.0 (Figure 4). The methyl chemical shifts of the mutated heme could not be directly assigned for every mutant because of the slow exchange. In this case, its reduction fraction was estimated by calculation under the condition of $\sum_i R_i^j = 1$. The obtained average R_i^j values are presented in Table S2 of the Supporting Information. In every case, the mutated heme was responsible for the first reduction step (nearly 90% reduction in R^1).

At first, we tried to determine the four macroscopic reduction potentials by measuring a differential pulse polarogram (DPP) with a mercury electrode. The observed DPPs are presented in Figures 6 and S1 (Supporting Information). The fitting of experimental data was carried out as described in Materials and Methods. However, the macroscopic reduction potentials obtained with the assumption of four consecutive one-electron reversible electrode reactions contradicted the reduction fractions obtained via NMR. The Met coordination structure might be susceptible to the interaction with the mercury electrode. A MPOH-coated gold electrode was utilized, because it should prevent a direct metal–protein interaction. Actually, responses were

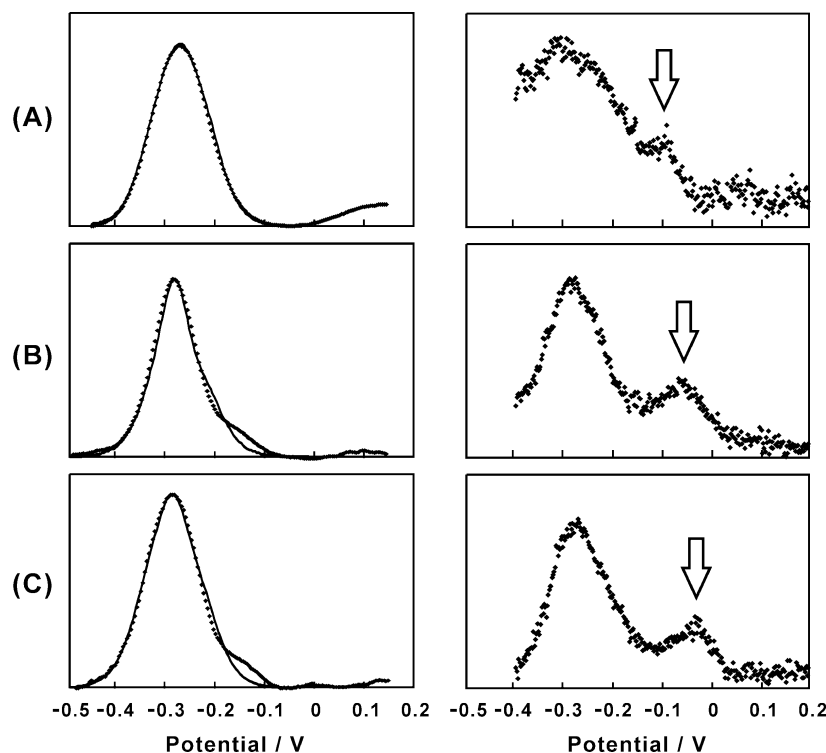


FIGURE 6: Differential pulse polarograms and voltammograms (DPPs and DPVs, respectively) of H35M (L_6^2) (A) and H25M (L_6^3) (B and C) cyt c_3 at pH 7.0 (A and B) and pH 9.0 (C). Left and right panels show data for DPPs and DPVs, respectively. The squares represent experimental data, and the lines are the best-fit theoretical curves. The arrows indicate responses ascribed to the Met-coordinated heme. The potentials are referred to the standard hydrogen electrode.

Table 4: Four Macroscopic Reduction Potentials of the Wild-Type and Mutated Cyt c_3 Species in 30 mM Sodium Phosphate Buffer (pH 7.0) or 30 mM Borate Buffer (pH 9.0) at 30 °C

sample (ligand ^a)	$E_I^{o'}$ ^b	$E_{II}^{o'}$	$E_{III}^{o'}$	$E_{IV}^{o'}$
pH 7.0				
wild type ^c	−242	−296	−313	−358
H22M (L_6^1)	nd ^d	−220 (+76)	−242 (+71)	−297 (+61)
H35M (L_6^2)	−95 (+147)	−240 (+56)	−285 (+28)	−331 (+27)
H25M (L_6^3)	−63 (+179)	−240 (+56)	−293 (+20)	−315 (+43)
H70M (L_6^4)	nd ^d	−210 (+86)	−269 (+44)	−312 (+46)
T24V	−221 (+21)	−273 (+23)	−305 (+8)	−340 (+18)
pH 9.0				
wild type	−240	−292	−324	−363
H22M (L_6^1)	nd ^d	−210 (+82)	−238 (+87)	−296 (+67)
H35M (L_6^2)	nd ^d	−230 (+62)	−297 (+27)	−337 (+26)
H25M (L_6^3)	−41 (+199)	−240 (+52)	−301 (+23)	−339 (+24)
H70M (L_6^4)	nd ^d	−210 (+82)	−259 (+65)	−312 (+51)

^a L_i^n stands for the i th ligand at heme n . ^b $E_i^{o'}$ ($i = I-IV$) is the macroscopic reduction potential at the i th reduction step relative to the standard hydrogen electrode (SHE). The standard errors are ± 2 mV. The differences (mutant − wild type) are given in parentheses. ^c From ref 16. ^d Not determined.

observed at approximately −100 and −50 mV in the differential pulse voltammograms (DPVs) of H35M (L_6^2) and H25M (L_6^3) cyt c_3 , respectively, as shown in Figure 6. They were also observed in the cyclic voltammograms (CVs, Figure S2). DPV of the buffer exhibited no signals in the relevant region (Figure S1), showing that the responses originated from the sample proteins. There is no macroscopic reduction potential in this potential range for the wild type (Table 4), and actually no response was observed in its DPV (not shown). These observations led to the assignment of the minor response to the Met-coordinated heme. Such a

response, however, was not observed for the other L_6 mutants (Figure S1), presumably due to the ligand conversion.

To analyze these data, we have to characterize the species responsible for the DPPs and DPVs. The peak current of CV measured with the MPOH-coated gold electrode is a linear function of the square root of the scan rate as shown in Figure S2 for all mutant cyt c_3 species at pH 9.0. This is the case not only for the major peak but also for the minor one, revealing that the species responsible for their electrochemical processes at the electrode are freely diffusing proteins. Furthermore, they should take on the intact structures, because every Met is coordinated to the heme in the potential range of interest as we will show below, and the structure of the Met-coordinated protein should be similar to that of the wild type in view of the crystal structure and CD and NMR spectra described above. The minor response was observed for only H35M (pH 7.0) and H25M (pH 7.0 and 9.0) cyt c_3 , which maintain the Met coordination throughout the redox process. In the potential range of the major response, every mutated heme should be reduced, since the reduction potential of the mutated heme is the highest among those of the four hemes as shown in the NMR results. We have shown that Met is coordinated to the heme in the reduced state at pH 9.0 and 7.0.

The DPP response curves are similar to the major response curves in DPV which can be seen in Figure 6 and Figure S1, revealing that they represent the same electrochemical processes. Therefore, the DPP responses should also come from the three nonmutated hemes of the freely diffusing intact proteins. The competence of the three nonmutated hemes was verified in the DPV experiments with H22M and H70M cyt c_3 (Figure S2). In view of these results, the polarograms were fitted to the three consecutive one-electron

Table 5: Microscopic Reduction Potentials at the First and Fourth Reduction Steps (e_1^I and e_1^{IV} for heme i , respectively) at pH 7.0 or 9.0 and 30 °C^a

sample	pH 7.0			
	e_1^I	e_2^I	e_3^I	e_4^I
wild type ^b	−308	−325	−286	−252
H35M (L ₆) ^c	nd ^d	−97 (+228)	nd ^d	nd ^d
H25M (L ₆) ³	nd ^d	nd ^d	−64 (+222)	nd ^d
T24V	−294 (+14)	−332 (−7)	−236 (+50)	−247 (+5)
	e_1^{IV}	e_2^{IV}	e_3^{IV}	e_4^{IV}
wild type	−293	−315	−344	−308
H35M (L ₆) ²	−295 (−2)	nd ^d	−316 (+28)	−281 (+27)
H25M (L ₆) ³	−277 (+16)	−295 (+20)	nd ^d	−285 (+23)
T24V	−285 (+8)	−319 (−4)	−291 (+53)	−307 (+1)
	pH 9.0			
	e_1^I	e_2^I	e_3^I	e_4^I
wild type	−309	−323	−286	−248
H25M (L ₆) ³	nd ^d	nd ^d	−42 (+244)	nd ^d
	e_1^{IV}	e_2^{IV}	e_3^{IV}	e_4^{IV}
wild type ^b	−297	−330	−345	−307
H25M (L ₆) ³	−308 (−11)	−323 (+7)	nd ^d	−295 (+12)

^a The differences (mutant − wild type) are given in parentheses. ^b Recalculated from the data reported in ref 14. ^c L_{*i*}^{*n*} stands for the *i*th ligand at heme *n*. ^d Not determined.

reversible electrode reactions (6). These macroscopic potentials can be ascribed to the Met-coordinated proteins as mentioned above. The obtained values are summarized in Table 4. A small shoulder found at approximately −0.13 V in the differential pulse polarograms of Figure 6 may be due to the denatured cyt c_3 adsorbed on the mercury surface. Although there are minor species for H22M and H70M cyt c_3 at pH 7.0, the observed DPPs and DPVs should represent the electrochemical reactions of the major low-spin species.

To clarify the effect of mutation on each heme, the microscopic reduction potentials were determined from the macroscopic reduction potentials and the reduction fractions (14) for H35M and H25M cyt c_3 , in which Met is stably coordinated during the redox process. Their microscopic reduction potentials at the first reduction step (e_1^I) and the fourth reduction step (e_1^{IV}) could be determined without any assumption. The former and the latter are those between the fully oxidized (S_0) and one-electron-reduced (S_1) states and between the three-electron-reduced (S_3) and fully reduced (S_4) states, respectively. They are summarized in Table 5. The reduction potentials of the relevant heme increased by 222–244 mV.

Structural and Redox Properties of T24V Cyt c_3 . H_{δ1} of His25 (sixth ligand of heme 3) is involved in hydrogen bonding with the carbonyl oxygen of asparagine 21 and O_γ of Thr24 in the oxidized state (16), while the latter is lost in the reduced state (2). The crystal structure of T24V DvMF cyt c_3 was determined at 1.0 Å resolution and is presented in panels A and C of Figure 2. It was similar to that of the wild type except for the mutated region. Therefore, the effect of the mutation on reduction potentials could be ascribed to elimination of the hydrogen bond with Thr24. The microscopic reduction potentials were calculated from reduction fractions (Table S2 of the Supporting Information) and macroscopic reduction potentials (Table 4) and are included in Table 5. The effect of the mutation was largest for heme

3, e_3^I and e_3^{IV} increasing by 50 and 53 mV, respectively. Those for DvH T24V cyt c_3 were 75 and 106 mV, respectively (35). These results indicate that Thr24 makes heme 3 the most effective electron donor in the reduced state with the lowest e^{IV} .

DISCUSSION

Two thioether linkages and the fifth axial His are the basic features of *c*-type cytochromes (18, 36). The essential role of the latter in formation of the former has been revealed for a genuine *c*-type and multiheme cytochrome gene in this work. We can conclude that the fifth axial His is essential not only for the redox regulation but also for the heme linking to *c*-type apocytochromes in *S. oneidensis* as well as in *E. coli* (31). There are three distinct types of Ccms, namely, types I, II, and III (36). The Ccms of *S. oneidensis* and *E. coli* are classified as type I. Since the *D. vulgaris* Ccm is also classified as type I (37), His should play the same role in the maturation of cyt c_3 in sulfate-reducing bacteria. A mechanism for the His recognition has been proposed on the basis of the crystal structure of *B. subtilis* ResA (a component of a type II Ccm) (38), where a special cavity called “a histidine clamp” was assumed to recognize His in the heme-binding motif (38). ResA exhibits a high degree of homology to CcmG of *E. coli* in terms of amino acid sequence (30% identical) (39). Thus, the involvement of the fifth axial His in heme linking would be a general mechanism in Ccms. Furthermore, since not only His in CXXCH but also that in CXXXXCH is recognized as the essential residue for heme linking, Ccm should be originated earlier than the establishment of oxygen respiration. Although the heme linking takes place independently among the four hemes in cyt c_3 , the formation of the heme architecture seems to be important for protein folding (40).

The effects of the L₆ mutations on the reduction potentials were significant for hemes 1 and 4 because their sixth ligands are converted during the redox process. It has been reported that the protein structure becomes more compact in the reduced state than in the oxidized state (2). This compactness would stabilize the Met coordination in the reduced state. In particular, conformational changes take place at the interfaces between hemes 1 and 2 and between hemes 2 and 4 during the oxidation. The effect of the conformational change should be smallest for heme 3. Furthermore, the environment around the sixth ligand of heme 3 may be flexible enough to adapt to the replacement of His with Met even in the oxidized state. A protein structure without Met coordination has less secondary structure and is less stable. The reduction fraction of H22M cyt c_3 (Table S2 of the Supporting Information) indicates that the order of the major reduction changed from hemes 4 → 1 → 2 → 3 to hemes 1 → 3 → 2 → 4 on mutation. In view of the similar secondary structure for wild-type and H22M cyt c_3 (Figure 1), the changes in the reduction potentials for all hemes should be due to a modification of the heme architecture.

To confirm the effect of the heme architecture on the reduction potentials, a differential pulse polarogram of the triheme architecture of H52M (L₅)² cyt c_3 (Figure S3 of the Supporting Information) was obtained. It showed an increase in the midpoint potential by ~50 mV compared to that for the wild-type tetraheme cyt c_3 , supporting the

contribution of the tetraheme architecture to the lowering of the reduction potentials. A similar tendency was also observed for cyt c_3 -like natural triheme cytochrome c species (cyt c_7 and PpcA). The midpoint potentials of those from *Desulfuromonas acetoxidans* and *Geobacter sulfurreducens* are -181 (41) and -146 mV (42), respectively.

The microscopic reduction potentials of *DvMF* cyt c_3 are pH-dependent in the neutral-pH region. However, the pK_a of propionate would not be affected directly by the replacement of an axial ligand because the behaviors of the heme methyl chemical shifts for H35M (L_6^2) and H25M (L_6^3) cyt c_3 between pH 7.0 and 9.0 were similar to those of the wild type (Figure 4 and Table S1 of the Supporting Information). Therefore, we can discuss the changes in the reduction potentials as the effects of the mutations. The most significant effect on the reduction potential of each heme was found for the mutated heme. The increases in e_3^1 of H25M cyt c_3 on replacement with Met were 222 and 244 mV at pH 7.0 and 9.0, respectively (Table 5). This is similar to the increase in the e_2^1 of H35M cyt c_3 at pH 7.0 (228 mV). In the case of *DvH* cyt c_3 , an isolated weak response was observed at -90 and -65 mV for H25M and H70M cyt c_3 , respectively (7, 8). Although the microscopic reduction potentials were not determined, the increases in e_3^1 and e_4^1 may be similar to those of *DvMF* cyt c_3 , judging from their highest macroscopic reduction potential. It should be noted that the Met-coordinated hemes are different for *DvMF* and *DvH* cyt c_3 . In the case of monoheme cyt c species, the replacement of the sixth axial Met with His reduced the reduction potentials, ΔE° , by 221, 390, and 205 mV in the cases of M80H horse mitochondrial cyt c (18), M78H *Aquifex* cyt c_{555} (43), and M61H *Pseudomonas* cyt c_{551} (44), respectively. The reported data indicate that the observed increase in the reduction potential induced by the replacement of His with Met in *DvMF* cyt c_3 is reasonable, although the observed potential itself is still much lower than those of mitochondrial cyt c species.

On the basis of this work and the investigations reported so far, we can summarize the factors contributing to the extremely low reduction potentials of *DvMF* cyt c_3 in comparison with a mitochondrial cyt c (~ 560 mV higher than the midpoint potential of cyt c_3). As shown in this work, bis-histidine coordination decreases the reduction potentials by more than 220 mV with respect to the Met–His coordination. The higher degree of heme exposure in cyt c_3 ($\sim 17\%$) is a well-known contribution (16). With simple estimation (45), one can expect a ~ 180 mV decrease in comparison with the reduction potential of a mitochondrial cyt c (4% exposure). The quadrilateral tetraheme architecture of cyt c_3 is another factor. The increase in the midpoint potential for a chain tetraheme architecture (small tetraheme cyt c) is more than 100 mV in comparison with that of cyt c_3 (46). This would be based on the difference in the mode of interheme interaction. We may regard ~ 100 mV as a global effect based on the heme architecture. These contributions account for ~ 500 mV out of 560 mV. In addition, hydrogen bonding with an axial ligand and an aromatic π – π interaction with the axial imidazole (5) decrease the reduction potential of a particular heme by 40–50 mV.

CONCLUSION

The fifth axial His was found to be directly involved in the heme linking in the process of maturation of c -type cytochromes. The three heme attachment sites (two thioether linkages and the fifth ligand) in the heme-binding motif CX_nCH ($n = 2-4$) yield a well-defined orientation of a heme in the protein. The sixth coordination will fix the folding of the polypeptide and refine the heme orientation. Since most c -type cytochromes are involved in electron transport, the fixed heme orientation should be important for the efficient electron transfer with a redox partner. This kind of heme-fixing device must have been particularly important for a small protein or a multiheme protein at an early stage of evolution, because the polypeptide was not long enough to stabilize its folding and heme cavity on its own. Thus, the fifth His was incorporated into the cytochrome c maturation reaction at the time of an anaerobic atmosphere, to which cyt c_3 can be traced back. This resulted in the evolutionary conservation of the heme-binding motif. Consequently, the fifth His became a strategically essential residue for c -type cytochromes. Bis-imidazole coordination at the heme iron could provide the best redox center for electron transport in a reduced environment because of its extremely low reduction potential. Then, the sixth ligand and protein environment were used for diversification of the electron transport system and for adapting to the oxidization of the atmosphere. To keep the reduction potential extremely low, the sixth axial His is also essential for cyt c_3 .

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SUPPORTING INFORMATION AVAILABLE

A table giving the chemical shifts of heme methyl signals in the five macroscopic oxidation states of mutated cyt c_3 species (Table S1), a table summarizing the reduction fractions of the four hemes at the four reduction steps (Table S2), a figure showing differential pulse polarograms and voltammograms of H22M, H25M, H35M, and H70M cyt c_3 (Figure S1), a figure showing cyclic voltammograms (CVs) of H25M and H35M cyt c_3 and peak current/(scan rate) $^{1/2}$ plots for CVs of H22M, H25M, H35M, and H70M cyt c_3 (Figure S2), and a figure showing a differential pulse polarogram of H52M cyt c_3 (Figure S3). This material is available free of charge via the Internet at <http://pubs.acs.org>.

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