

SUPPLEMENTAL DATA

Table S1. Cytochrome *c* maturation (Ccm) components and their role

Ccm protein	Structural Features	Proposed Funtion
CcmA	Peripheral membrane protein with ATP-binding domain (Walker A and B motifs)	Part of the CcmABCD complex, a predicted ABC-type transporter; required for the ATP-dependent release of holoCcmE from the CcmABCDE complex.
CcmB	Six TM helices and a conserved FXXDXXDGSL motif	Part of the CcmABCD complex, a predicted ABC-type transporter; required for the release of holoCcmE from the CcmABCDE complex
CcmC	Six TM helices with a tryptophan-rich (WGX[F,Y,W]WXWDXRLT) motif, and two conserved His residues	Part of the CcmABCD complex, a predicted ABC-type transporter; transfers heme to apoCcmE, providing axial ligands to the heme iron; acts as a CcmE-specific heme lyase.
CcmD	Small single TM helix protein with no conserved domains	Part of the CcmABCD complex, a predicted ABC-type transporter; required for the release of holoCcmE from the CcmABCDE complex
CcmE	Single TM helix with a periplasmic domain containing a conserved HXXXY motif	Heme chaperone binding covalently vinyl-2 of heme via the conserved His residue; delivers heme to the apocytochrome <i>c</i> substrates
CcmF	Eleven TM helices with a tryptophan-rich (WGGXWFWDPVEN) motif, and four conserved His residues (two of them are ligands of a <i>b</i> -type heme)	Part of the CcmFHI heme ligation complex; interacts with apocytochromes <i>c</i> and CcmE, and suggested to reduce heme in holoCcmE for its transfer to the apocytochrome
CcmG	Single TM helix with a periplasmic domain containing a thioredoxin motif (CXXC)	Binds poorly apocytochromes <i>c</i> , and suggested to reduce the disulfide at the heme binding site of apocytochrome <i>c</i> either directly or via CcmH
CcmH	Single TM helix with a periplasmic domain containing a thioredoxin-like motif (LRCXXC)	Part of the CcmFHI heme ligation complex; interacts with apocytochrome <i>c</i> and suggested to reduce its disulfide bond for the stereo-specific heme ligation
CcmI	Two TM helices, linked by a cytoplasmic loop with a leucine-zipper-like motif, and a large periplasmic domain containing three TPR repeats	Part of the CcmFHI heme ligation complex; is an apocytochrome <i>c</i> chaperone binding to the C-terminal portion of the apocytochrome <i>c</i> via its periplasmic domain
CcdA	Six TM helices with two conserved redox active Cys residues	Thiol-reduction of CcmG

Table S2. Nucleotide sequences of the oligonucleotide primers used in this study.

Designation	Constructs	Nucleotide sequence (5' to 3')
CcmE-NdeI-Fwd CcmE-BamHI-Rv	pAV4	CGG CAT ATG AGA AAC CTG AAG AAA ACG CGG ATT GGA TCC TTA TGC CGC TCA GCC CTC GGG CT
CcmEH123A-Fwd CcmEH123A-Rv	pAV4H123A	CGA AAT CCT GGC CAA GGC TGA CGA AAA CTA CAT G CAT GTA GTT TTC GTC AGC CTT GGC CAG GAT TTC G
FLAG-Fw FLAG-Rv	pFLAG-CcmI	TAC CAT GGG CGA CTA CAA GGA CGA CGA CGA CAA GAG CAG CGG CCA TAT CG GGC CGC TGC TCT TGT CGT CGT CGT CCT TGT AGT CGC CCA TGG TAT ATC TCC

Figure S1. Verissimo *et al.*,

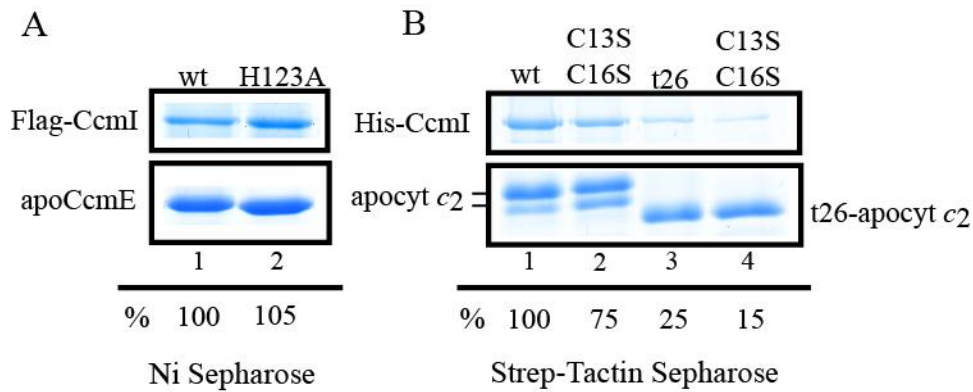


Figure S1. A) Co-purification of Flag-CcmI with His-apoCcmE (lane 1) and its His-apoCcmE-H123A mutant (lane 2) using Ni Sepharose resin. **B)** Co-purification of FlagCcmI with Strep-apocytochrome c_2 (lane 1), its C13SC16S mutant (lane 2), the t26-truncated Strep-apocytochrome c_2 (lane 3) and its cysteine-less derivative (lane 4) using the Strep-Tactin Sepharose resin. The data show that apoCcmE interacts specifically with CcmI and that the Flag-fused CcmI shows the same behavior towards apocytochrome c_2 that its His-derivative, indicating that the different tags do not interfere with the observed protein-protein interactions.