

hxl108	5'-ATTGTTGACGTTCCACCAGGGTTA-3'
hxl109	5'-TGAAAATCTCGCCAAGCTTGTCAAGCCAGGCTAATTGATAGAGTAA-3'
hxl110	5'-TTACTCTATCAATTAGCCTGGCTTGACAAGCTTGGCGAGATTTTCA-3'
hxl111	5'-CAAATAGGGGCAGGACGGGAATTCAGGCGTAGCAACCAGGCGTTTA-3'
hxl112	5'-TAAACGCCTGGTTGCTACGCCTGAATTCCCGTCCTGCCCCTATTTG-3'
hxl113	5'-CCCCGTCTTTCGCTTCCTGGTCATT-3'
hxl114	5'-ACTAATTGTCATGCAGGGTTCTCC-3'
hxl115	5'-CTGAAAATCTCGCCAAGCTTGTCACTAACATGGGGGGTTGTGGGAAATT-3'
hxl116	5'-AATTTCCACAACCCCATGTTAGTGACAAGCTTGGCGAGATTTTCAG-3'
hxl117	5'-GCCAGATAATTCGACTTAAACTTTCAGGCGTAGCAACCAGGCGTTTAA-3'
hxl118	5'-TTAAACGCCTGGTTGCTACGCCTGAAAGTTTAAGTCGAATTATCTGGC-3'
hxl119	5'-GAGCAAAACCTACTGTCACAACCAT-3'

Table S1 Primer Details Used in Construction and isolation of Mutants

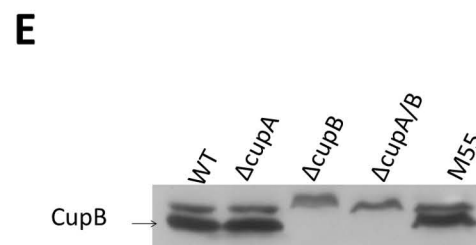
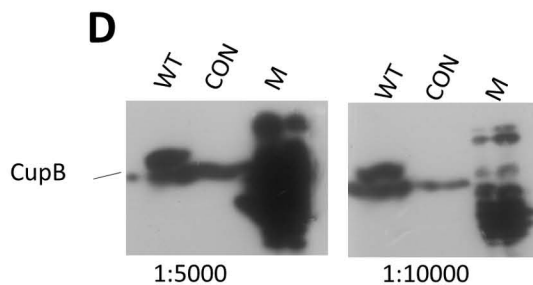
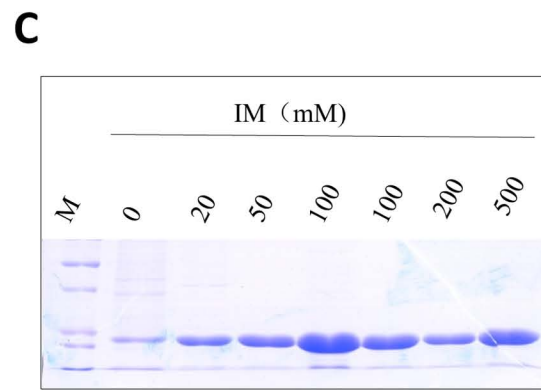
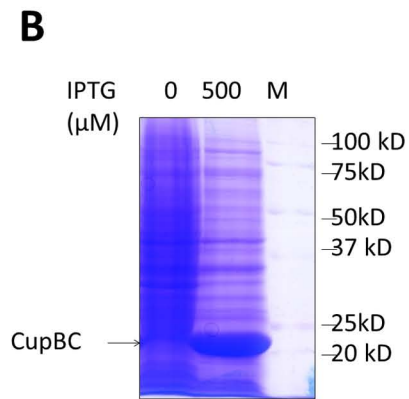
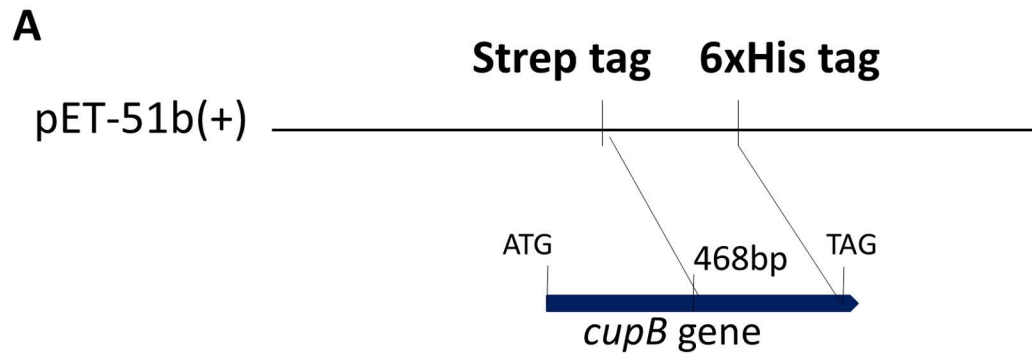


FIG.S1 Polyclonal antibody of CupB preparation and immunological characterization of CupB in mutants.

A. Construction of expression vector of CupB C-terminal (CupBC). B. expression analysis of CupB C-terminal. CupBC: CupB C-terminal, M: protein marker. C. purification of CupB C-terminal with Ni²⁺-NTA-agarose resin. IM: imidazole. D. immunological characterization of CupB. CON: CupB-His-Myc. M: protein marker. E. immunological characterization of CupB in different mutants.

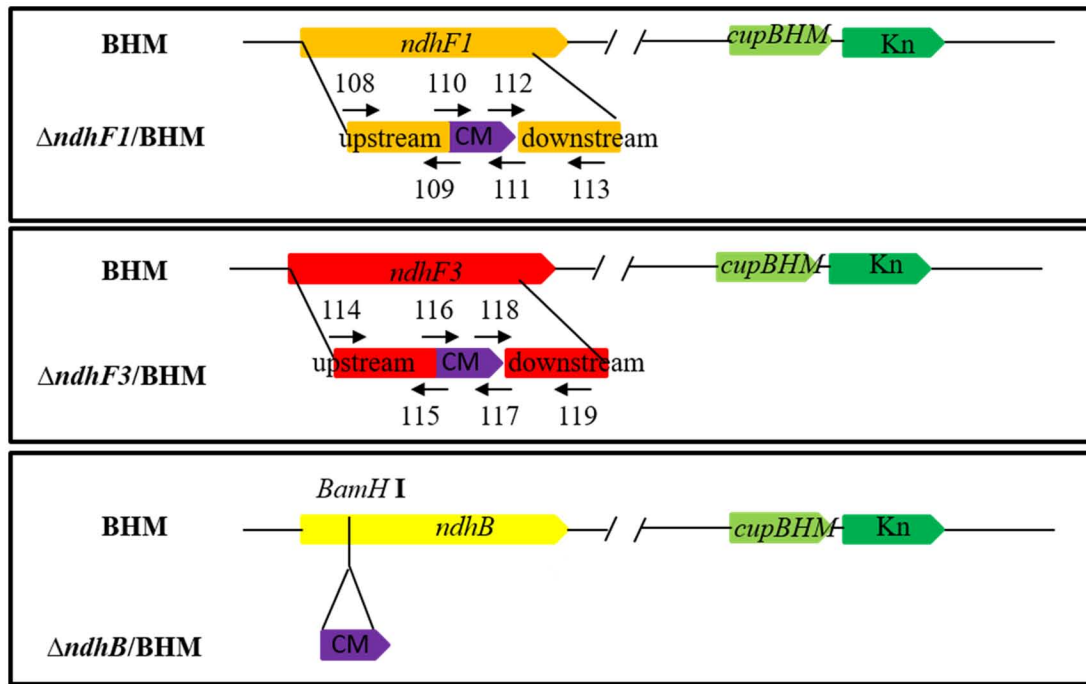
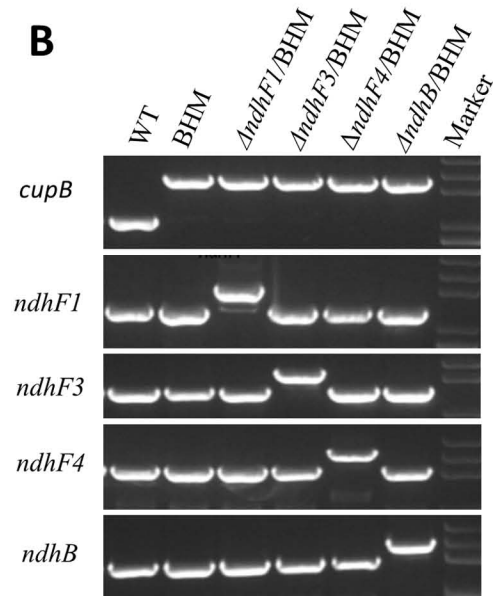
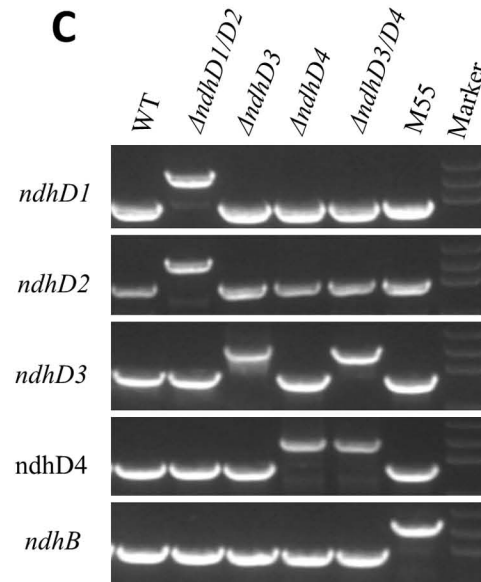
A

FIG.S2 Construction and segregation of *ndhF* and *ndhD* mutants.

A. chloramphenicol resistance cassette was inserted into *ndhF1*, *ndhF3* and *ndhB*, separately. *cupBHM*: CupB tagged with 6xHis-cMyc at the C-terminal. CM: chloramphenicol resistance cassette.

Kn: Kanamycin resistance cassette.

108-119: hxl108-hxl119, see supplemental data table 1.

B**C**

B. The mutated genes in the transformants of *ndhB* and *ndhF* mutants were segregated to homogeneity as determined by PCR amplification. **C.** The mutated genes in the transformants of M55 and *ndhD*s were segregated to homogeneity as determined by PCR amplification.

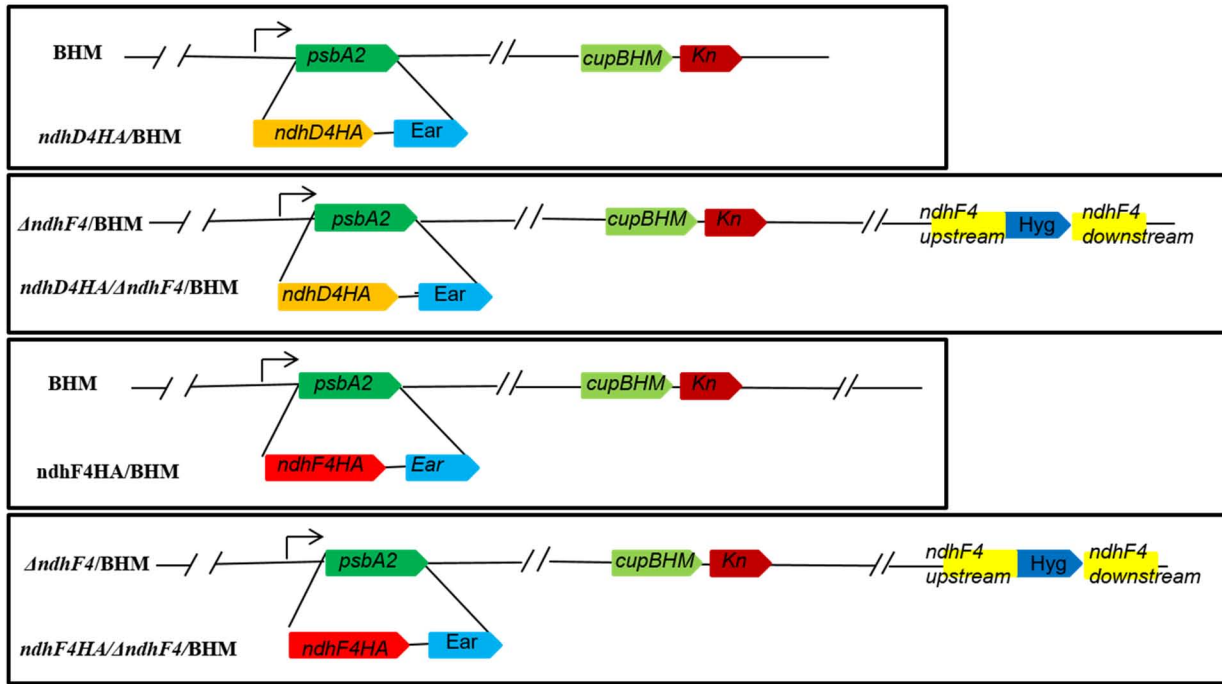
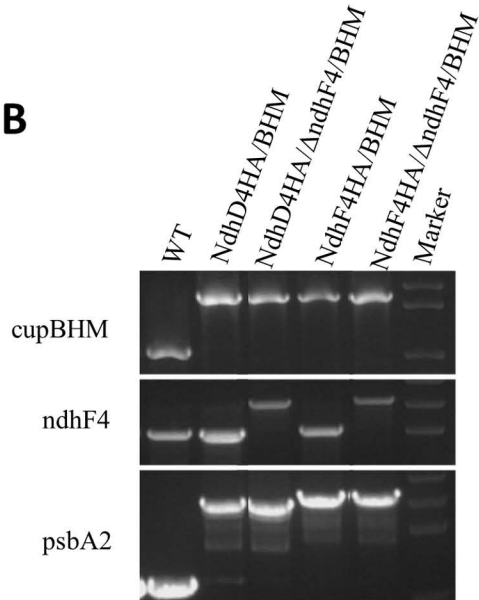
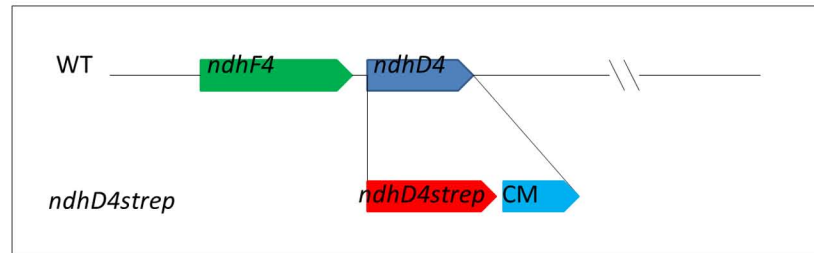
A**B****C**

FIG.S3 construction and segregation check of *ndhD4HA/BHM*, *ndhD4HA/ΔndhF4/BHM*, *ndhF4HA/BHM*, and *ndhF4HA/ΔndhF4/BHM*, and *ndhD4strep*.

A. *ndhD4HA/BHM*, *ndhD4HA/ΔndhF4/BHM*, *ndhF4HA/BHM*, and *ndhF4HA/ΔndhF4/BHM* were constructed as shown in **FIG.S3A**. PS II A2 gene was exchanged by *ndhD4* and *ndhF4* tagged with HA gene. **B.** The mutated genes in the transformants of *ndhD4HA/BHM*, *ndhD4HA/ΔndhF4/BHM*, *ndhF4HA/BHM*, and *ndhF4HA/ΔndhF4/BHM* mutants were segregated to homogeneity as determined by PCR amplification. **C.** *ndhD4strep* was constructed as shown in **FIG.S3C**. *ndhD4* gene was exchanged by *ndhD4* tagged with strep gene.

psbA2: photosystem II A2 gene.

cupBHM: CupB tagged with 6xHis-cMyc at the C-terminal.

CM: Chloramphenicol resistance cassette.

Kn: Kanamycin resistance cassette.

Ear: Erythromycin resistance cassette.

Hyg: Hygromycin resistance cassette.

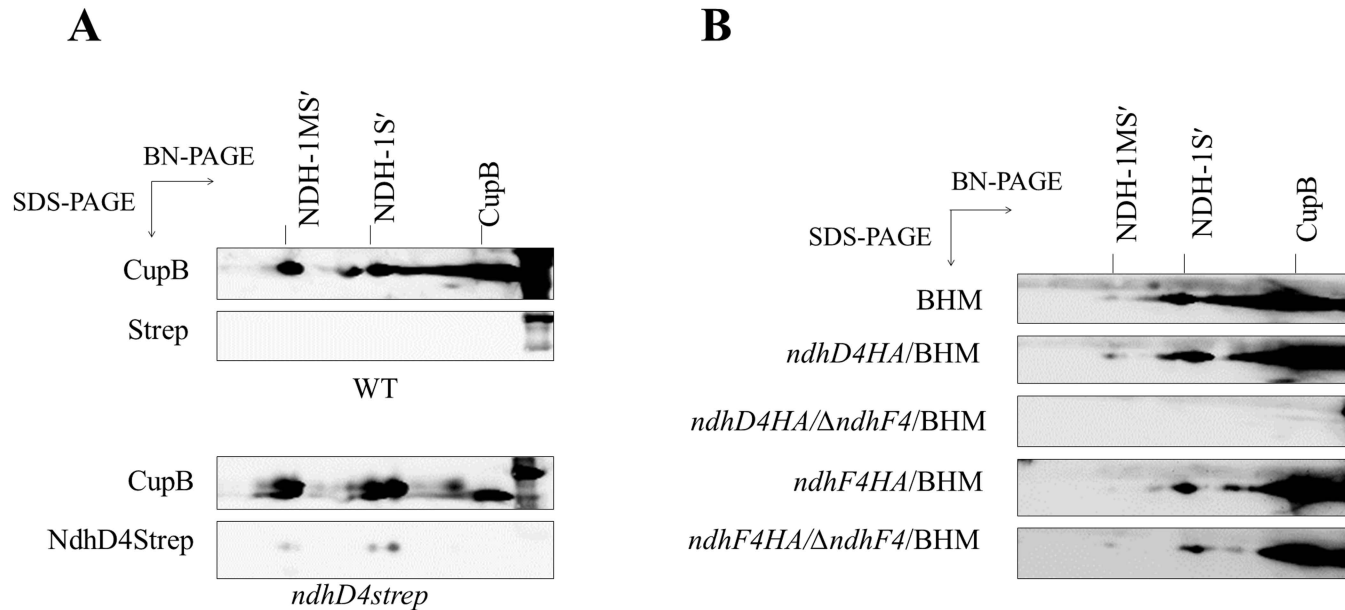


FIG. S4. Association of CupB with NdhD4 and NdhF4.

(A) The position of CupB and NdhD4 were detected by specific antibody. Total membrane proteins from wild type and *ndhD4strep* were separated and immunodetected with antibody against CupB and Strep, respectively. (B) Assembly of CupB complex in transformed mutants background. Membrane proteins from BHM, *ndhD4HA/BHM*, *ndhD4HA/ΔndhF4/BHM*, *ndhF4HA/BHM*, and *ndhF4HA/ΔndhF4/BHM* were separated by BN-SDS-PAGE, and were immunodetected with antibody against CupB.