

Supporting Information

Dissection of Hydrogen Bond Interaction Network around an Iron-sulfur Cluster by Site-specific Isotope Labeling of Hyperthermophilic Archaeal Rieske-type Ferredoxin

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Table S1. PCR primers used for construction of new BL21(DE3) strain RF4RIL

PCR primers	Purpose	Oligonucleotide sequences (5'→3')	from ^a	to ^a
aspC UpF	Up.F	ATGAACGCCTGCATAATCCCTTCTCTGC	989332	989358
aspC UpR ^b	Up.R	CTTCGAAGCAGCTCCAGCCTACACGCATTAATAACAATGAAGCCCGC	989580	989602
aspC DownF ^c	Down.F	GGAATAGGAACTAAGGAGGATATTCATATGGACGAGGTTCCATTATGGTTACAGAAGG	990794	990821
aspC DownR	Down.R	GTCCGTCCATCGCTTACACCAAATCTAAAG	991163	991192
aspC ext.F ^d	deletion of <i>aspC</i>	TTGATGACAGCGGCCTGACACTGATGCAG	988858	988886
aspC ext.R ^d	deletion of <i>aspC</i>	ACTCCAACCTCTTTGGTCTGGTTGATGG	991552	991579
tyrB UpF	Up.F	GCGAACGTGATACCCGCTTGGCGTAGTC	4173966	4173994
tyrB UpR ^e	Up.R	GAACCTCGAAGCAGCTCCAGCCTACACGCGATGGTTCTCCAGGTTTACGGGCAG	4174490	4174516
tyrB DownF ^f	Down.F	GGAATAGGAACTAAGGAGGATATTCATATGTGCAGGAAAGCAGGCTGGAG	4175711	4175730
tyrB DownR	Down.F	TGCCGAGGAGGTTAAAGGTGATTATTC	4176130	4176156
tyrB ext.F ^d	deletion of <i>tyrB</i>	GCGAAGGCAAACCTGGTCAACGTTC	4173729	4173754
tyrB ext.R ^d	deletion of <i>tyrB</i>	GATTGACCAGCCCCCTACCTACAATGG	4176451	4176478

^a Positions in the genomic sequence of BL21(DE3) strain (NCBI GenBank code, AM946981). *aspC*, 989603-990793; *tyrB*, 4174517-4175710.

^b This PCR primer consists of the oligonucleotide sequences from the BL21(DE3) *aspC* gene plus flanking region (23 bp) and the kanamycin resistance cassette gene (24 bp) (**Fig. 2**, step 1).

^c This PCR primer consists of the oligonucleotide sequences from the BL21(DE3) *aspC* gene plus flanking region (28 bp) and the kanamycin resistance cassette gene (30 bp) (**Fig. 2**, step 1).

^d Used for confirmation of the deletion of target chromosomal genes.

^e This PCR primer consists of the oligonucleotide sequences from the BL21(DE3) *tyrB* gene plus flanking region (27 bp) and the kanamycin resistance cassette gene (27 bp) (**Fig. 2**, step 1).

^f This PCR primer consists of the oligonucleotide sequences from the BL21(DE3) *tyrB* gene plus flanking region (20 bp) and the kanamycin resistance cassette gene (30 bp) (**Fig. 2**, step 1).