

S6 Fig

Phenylalanine auxotrophy mutation pheA10 is allelic to PHA2 (A) Short read alignment. (B) Sanger resequencing. Red frame, TAA nonsense mutation appearing at codon 161. (C) 33G-D373 plated on selective media immediately after transformation with low copy number plasmids bearing indicated PHA2 alleles. Vector, pRS316. (D) Asp⁺ and Asp⁻ variants of 6P-33G-D373 spotted for nonsense suppression and copper resistance. Series of 5-fold dilutions are shown.