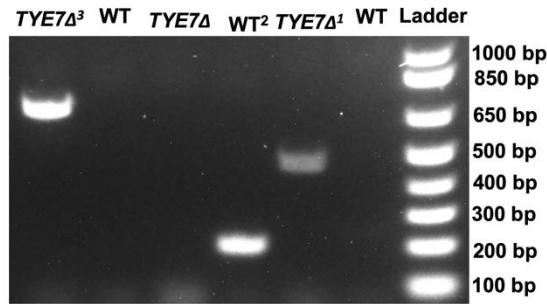
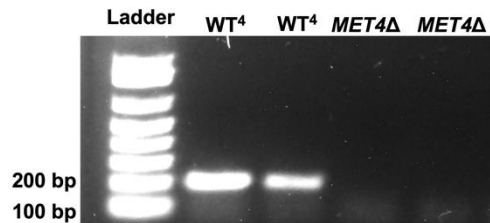


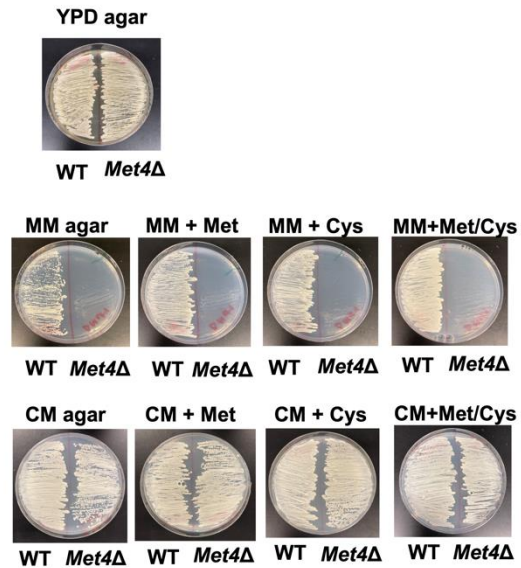
A) *TYE7*Δ genotype



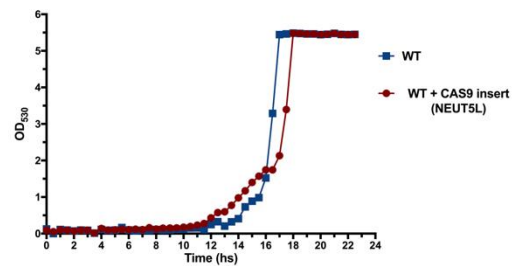
B) *MET4*Δ genotype



C) *MET4*Δ auxotrophy



D) Growth of *Candida albicans* SC5314 with Cas9 insert



S3 Figure: Validation of the genotype of deletion mutants. **A)** Confirmation of generation of *TYE7*Δ mutant by PCR and gel electrophoresis where the *TYE7*Δ¹ band (~ 478 bp) confirmed the replacement of *TYE7* gene sequence with the gene drive cassette and the WT² band (~ 221 bp) confirmed the presence of *TYE7* gene in wild type. The deletion of the *TYE7* gene sequence in the *TYE7*Δ mutant (*TYE7*Δ³ band; ~747 bp) confirmed the integration of Cas9 plasmid into *C. albicans* genome at the NEUT5L locus. **B)** Confirmation of *MET4*Δ genotype by PCR and gel electrophoresis where the WT⁴ band (~ 224 bp) confirmed the presence of *MET4* gene in wild type and the deletion of *MET4* gene sequence in *MET4*Δ mutant. **C)** Growth of *C. albicans* SC5314 wild type and *MET4*Δ mutant in rich agar media (YPD), Minimal Media (MM), and Complete Minimal (CM) dropout media (synthetic mix minus methionine and cysteine) with or without addition of methionine and/or cysteine at 40 μg/mL final concentration. **D)** The integrated-plasmid NEUT5L did not show a defective effect on fungal fitness.