

Supplementary material

S1 Figure. The effect of different stresses on strains expressing Rha1-GOF. Overnight cultures of SC5314 and Rha1-GOF were spotted under the indicated stress conditions. Rha1-GOF cultures grow normally under the different stress conditions including, the metal stressors, 100mM FeSO₄, 20 mM FeCl₃, and 5mM CuSO₄, the oxidative stress-inducing agent hydrogen peroxide (H₂O₂) at 5mM and the osmotic stressor 1M sorbitol, 0.4M CaCl₂, 0.15mM Menadione, 38mM hydroxyurea, pH 10, Glycerol 250mM, Hygromycin B 100µg/ml MMS 0.02%.

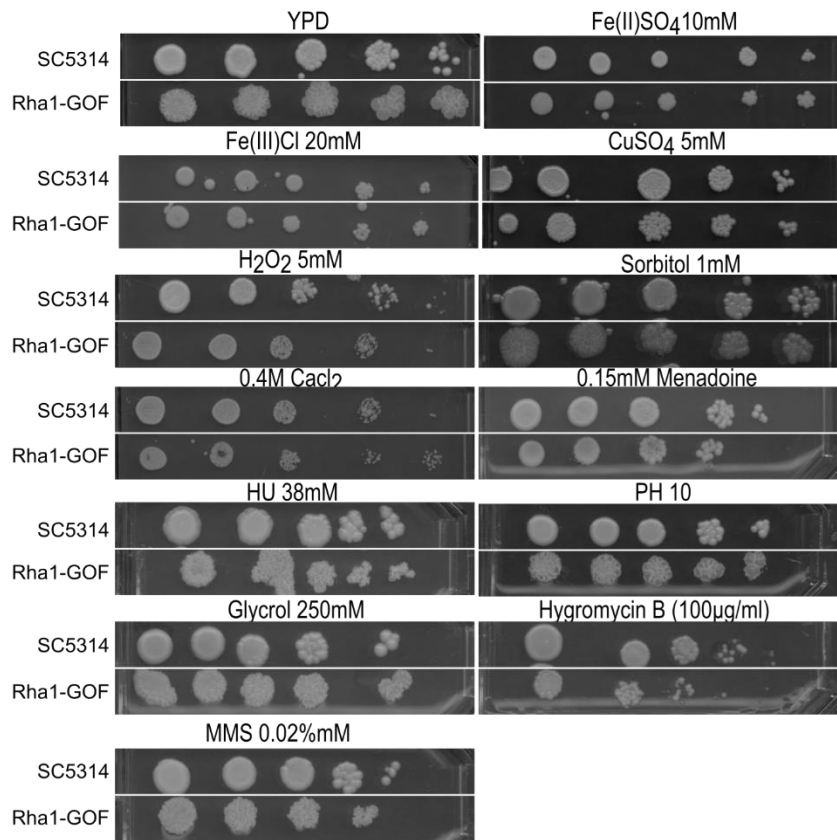
S2 Figure. RNAseq analysis of Rha1-GOF. (A) Heatmap of up-regulated genes (log₂fold >4, P_{adj} <0.001) versus the WT under non hyphal conditions. (B) Top 35 down-regulated genes P-values <0.03 in the Rha1-GOF compared with the WT under non-hyphal inducing conditions.

S3 Figure. Rha1 strains grow normally on media without arginine. Growth curves (OD₆₀₀ of *C. albicans* SC5314 WT, Rha1 GOF and *rha1*^{-/-} to determine influence of Rha1 function in arginine biosynthesis. The overnight strains were cultured in liquid SC without arginine and incubated at 30°C for 72hrs.

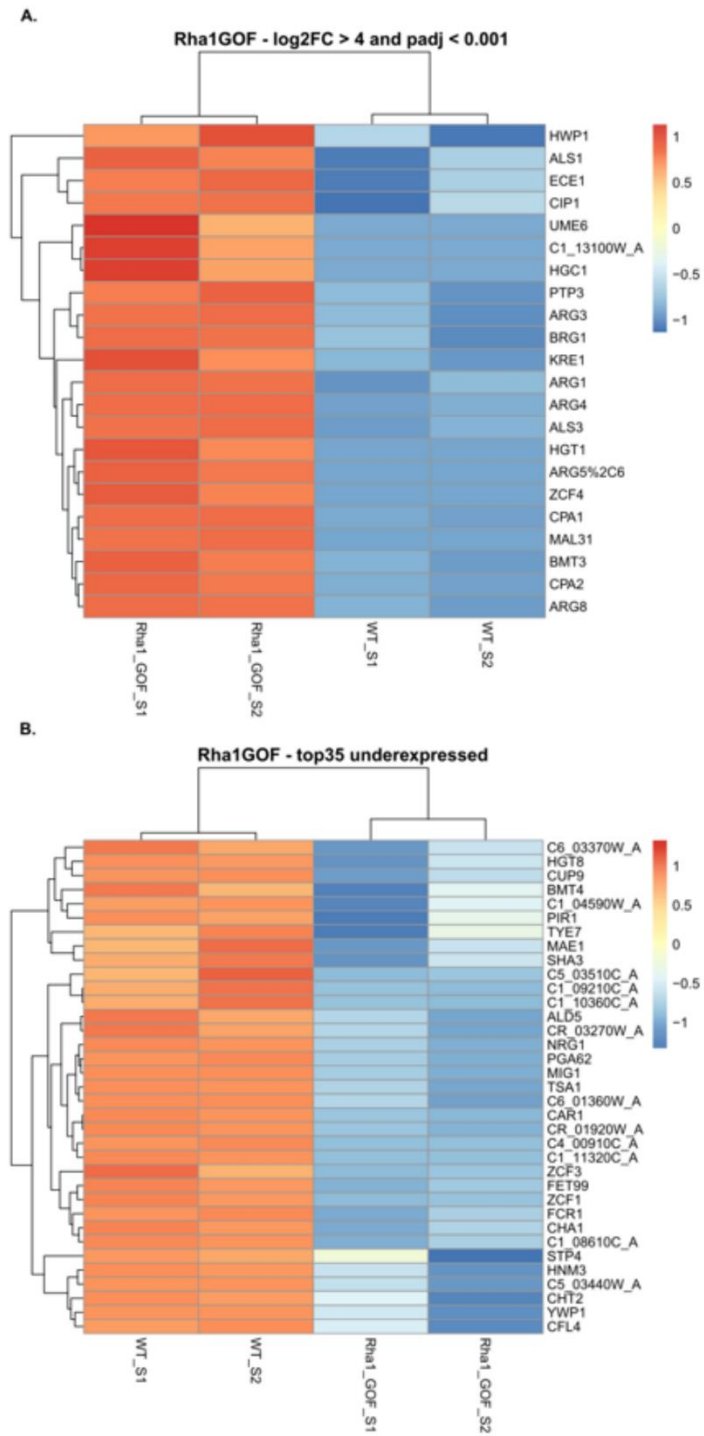
S4 Figure. One copy of Rha1 restores the filamentation morphology in *rha1* Δ/Δ. (A) The WT SC5314, SC1604M4B, (*rha1* Δ/Δ), SC1604MK2B (*rha1* Δ/Δ::RHA1) were (B) The DIC images of the indicated strains grown in the liquid Spider medium, 4 hrs at 37°C.

The complementary strain SC1604MK2B induced hyphal formation in Spider medium. Scale Bar represents 10 μ m. C) Shown are the serum treated cells after 4 hrs at 37°C. Cells which represent one copy of Rha1 add back the filamentation function. Scale bar is 5 μ m.

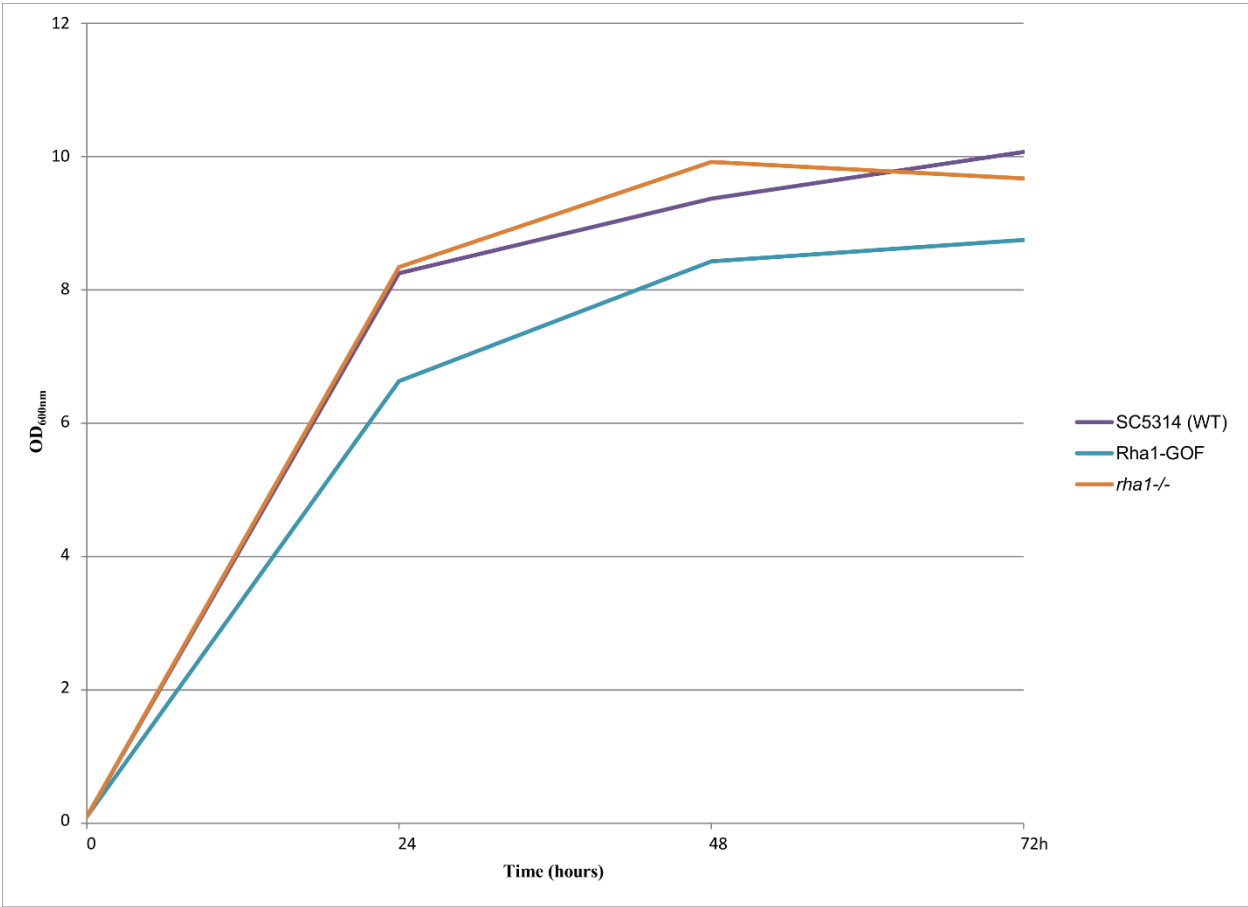
S1 Figure



S2 Figure



S3 Figure



S4 Figure

