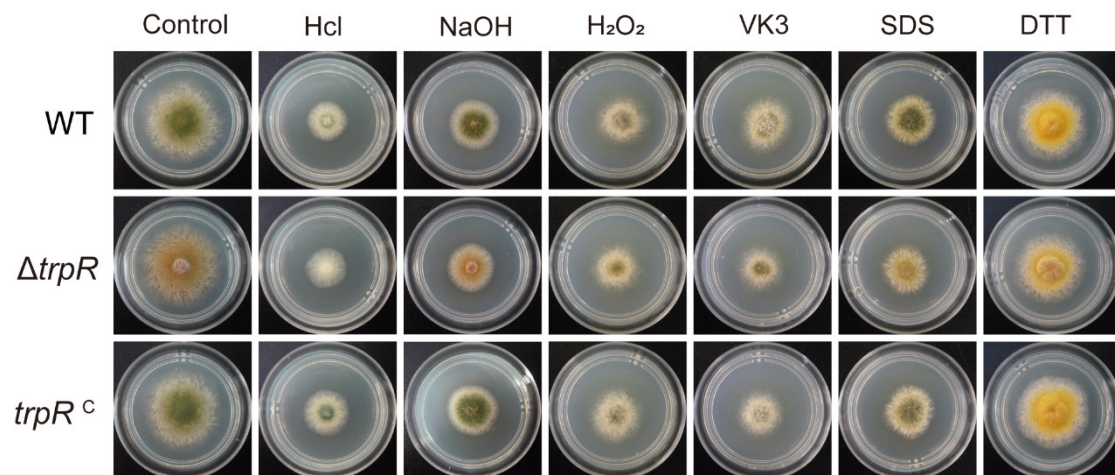
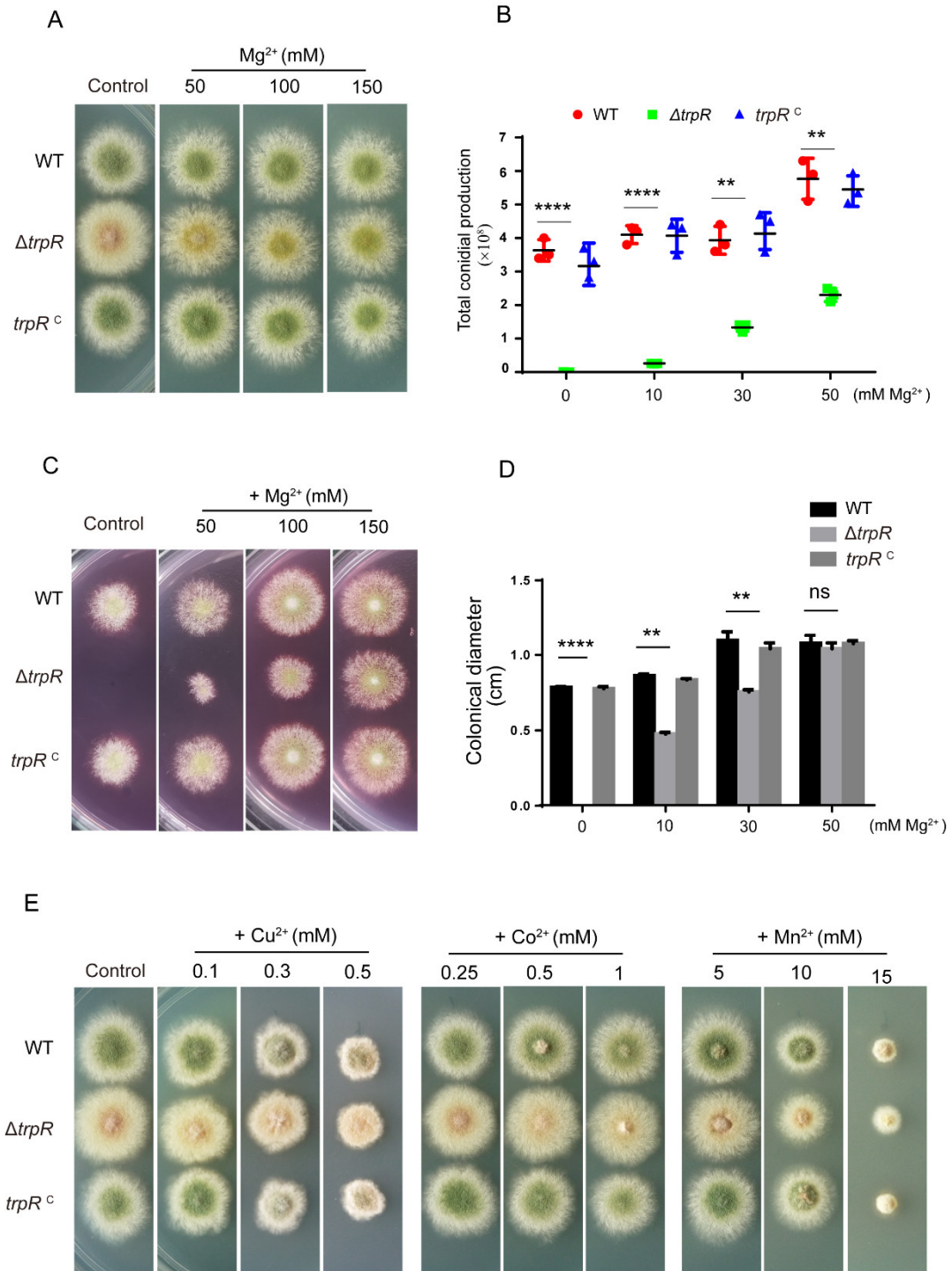


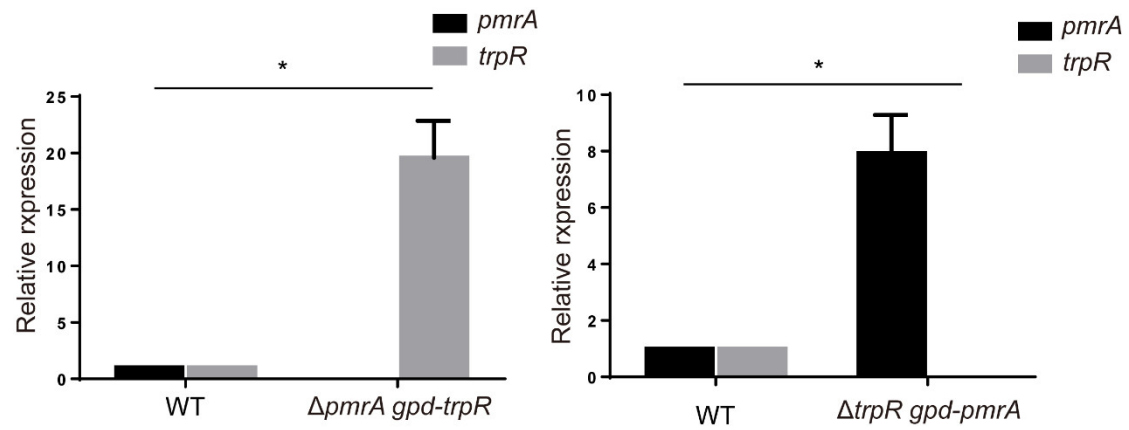
**Figure S1.** Diagnostic PCR analyses of the indicated strains. (A) The TrpR-GFP genomic DNA was used as template for diagnostic PCR with primer pairs *trpR-gfp* P1/*gfp* R for line 1, and *pyrG* F/*trpR-gfp* P6 for line 2. (B) As for the  $\Delta trpR$  mutant, line 1 and line 2 were performed with primer pair *trpR* P1/ *pyrG* R and *pyrG* F/ *trpR* P6, respectively. Line 3 was performed with primer pair *trpR* diag F/*trpR* diag R suggesting the successful deletion for *trpR*. Line 4 showed the PCR products with primer pair *trpR* diag F/ *trpR* diag R using wild-type genomic DNA as template.



**Figure S2.**  $\Delta trpR$  mutant showed no hypersensitivity to various stress agents. Colony morphology for the indicated strains grown on solid PDRUU in the absence or presence of 100 mM Hcl, 10 mM NaOH, 0.5 mM  $H_2O_2$ , 10  $\mu$ M vitaminK3 (VK3), 0.005% SDS or 3 mM DTT at 37 °C for 2.5 days.



**Figure S3.** The defects in the  $\Delta trpR$  mutant can be restored by excess magnesium. (A) Colony morphology for the indicated strains grown on solid PDRUU in the absence or presence of 50, 100 and 150 mM  $MgCl_2$  at 37 °C for 2.5 days. (B) Quantitative total conidial production for the strains shown in panel A. (C) Colony morphology for the indicated strains grown on solid PDRUU supplemented with 5  $\mu g/ml$  CR and in the absence or presence of 50, 100, 150 mM  $MgCl_2$  at 37 °C for 2.5 days. (D) Quantitative total colonial diameter for the strains shown in panel C. Values represent mean  $\pm$  SD from three replicates. (ns, not significant; \*\*,  $P < 0.001$ ; \*\*\*,  $P < 0.001$ ; \*\*\*\*,  $P < 0.0001$ ). (E) Colony morphology for the indicated strains grown on solid PDRUU supplemented with  $Cu^{2+}$ ,  $Co^{2+}$  or  $Mn^{2+}$ .



**Figure S4.** Verification of the overexpression strains. qRT-PCR analysis was performed to verify the  $\Delta pmrA$  *gpd-trpR* (left) and  $\Delta trpR$  *gpd-pmrA* (right) strains after growth of cultures in liquid PDR or PDRUU medium for 18 h at 37 °C. The *tubA* gene was used as an internal control. Values represent mean  $\pm$  SD from three replicates. (\*,  $P < 0.05$ ).