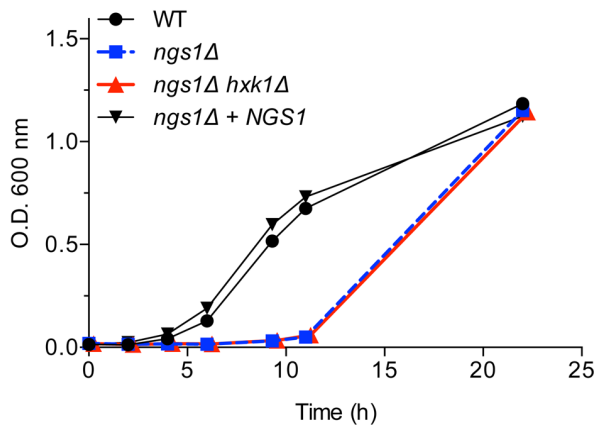
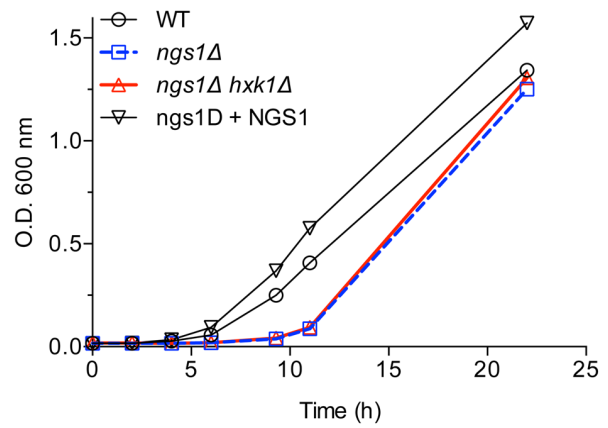


A. YPD cultures shifted to YNB + Dextrose

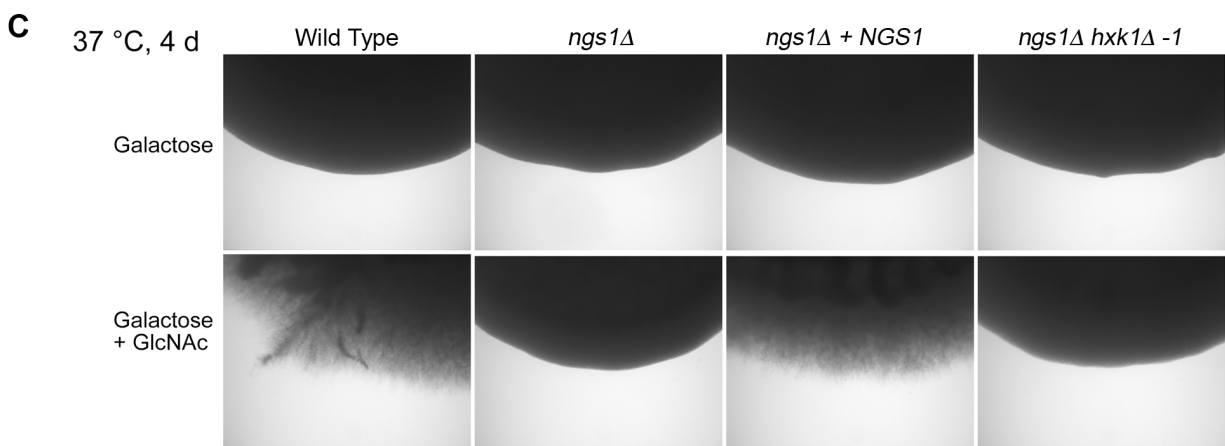
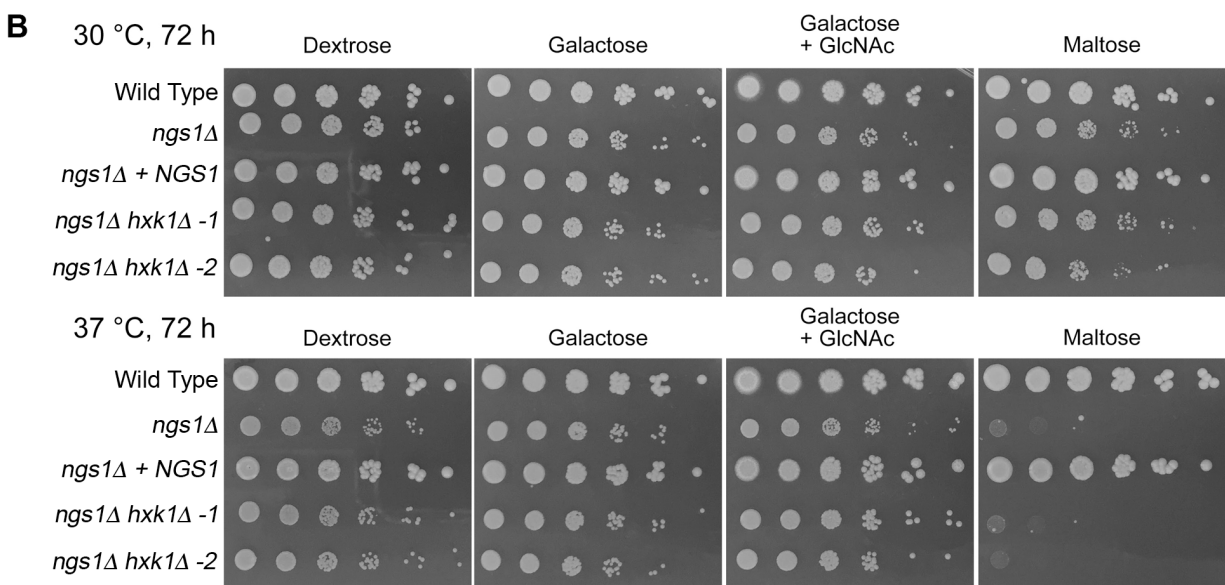
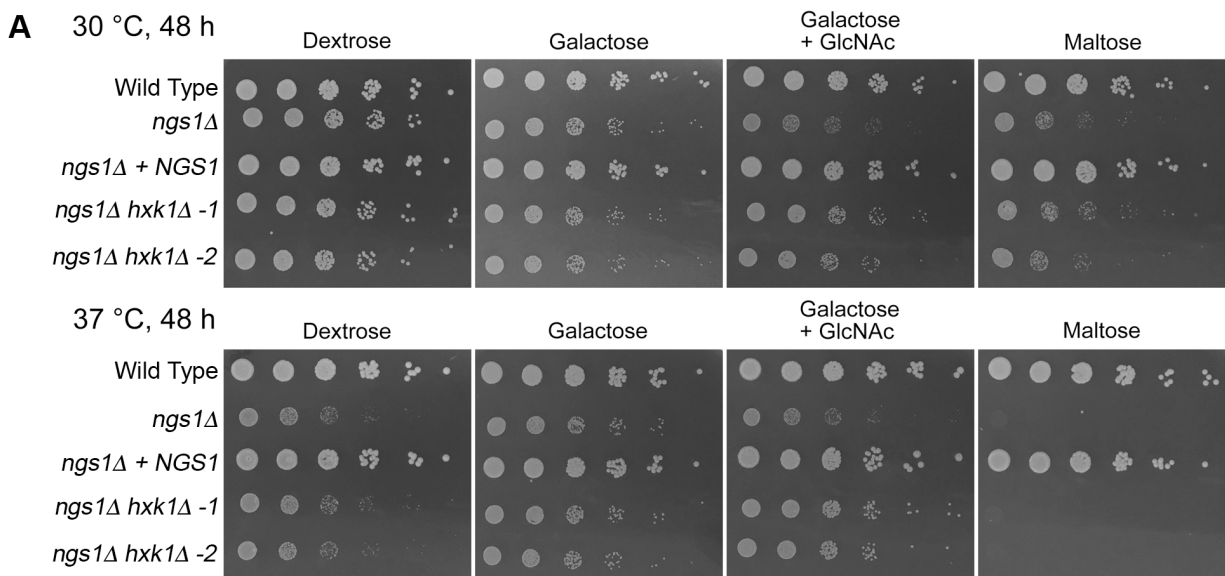


B. YPD cultures shifted to YNB + Maltose



**Supplemental Figure S1. The *ngs1Δ* mutant is delayed in growth after switching media.**

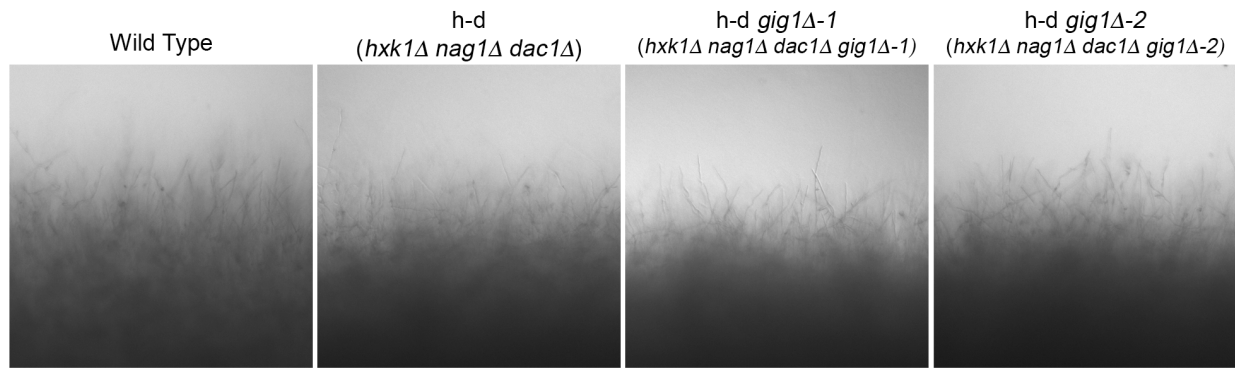
The indicated cells were grown first in rich YPD medium and then samples were washed and diluted into synthetic medium containing (A) dextrose or (B) maltose. Cells were grown at 30°C and then the O.D.600 was recorded at the indicated times. The cells included the wild type control strain (LLF100), the *ngs1Δ* strain (SN1429), the *ngs1Δ + NGS1* complemented strain (SN1430), and the *ngs1Δ hxx1Δ* strain (KM1433).



**Supplemental Figure S2. Deletion of HXK1 from *ngs1Δ* cells reduces the toxic effects of GlcNAc, but does not improve growth on maltose or invasive growth into GlcNAc agar medium.**

(A) Dilutions of cells indicated on the left were spotted onto synthetic agar medium containing the sugar shown at the top. Plates were incubated for the time and temperature indicated at the upper left side of each set.

(B) The indicated cells were spotted onto agar medium containing the sugar shown on the left, incubated for 4 d at 37°C, and then the edge of the spot was photographed to record the extent of invasive growth. The cells included the wild type control strain (LLF100), the *ngs1Δ* strain (SN1429), the *ngs1Δ* + *NGS1* complemented strain (SN1430), and the *ngs1Δ hxx1Δ* strain (KM1433).



**Supplemental Figure S3. *GIG1* does not influence the ability of *ron1Δ* cells to form hyphae.**

Cells were spotted onto medium containing 50 mM galactose + 50 mM GlcNAc and then incubated at 37°C for 4 d. The strains included the wild type (DIC185), *hpk1Δ nag1Δ dac1Δ* (AG738), *hpk1Δ nag1Δ dac1Δ gig1Δ-1* (KM1435) and *hpk1Δ nag1Δ dac1Δ gig1Δ-2* (KM1436).