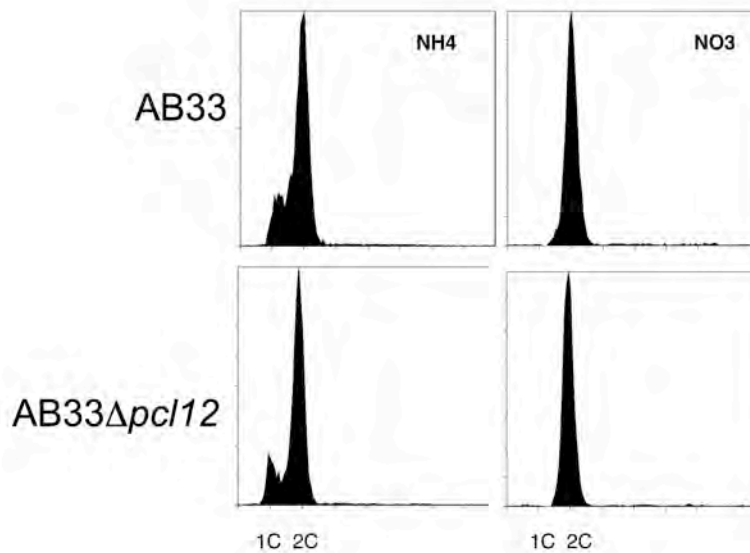


Supplemental Figure 1. FACS analysis of AB33 and AB33 Δ *pci12* cells. Cultures of the indicated strains were grown for 8 hours in non-inducing conditions (minimal medium with ammonium as nitrogen source, NH₄) or inducing conditions (minimal medium with nitrate as nitrogen source, NO₃). Observe that both control and Δ *pci12* cells accumulate with a 2C DNA content in inducing conditions.



Supplemental Figure 2. Confrontation assay. We followed the procedure described by Snetselaar et al., 1996. Briefly, cultures of wild-type and mutant cells were grown overnight in CMD, washed once in distilled sterile water and resuspended to a density of about 5×10^7 cells/ml. On a glass microscope slide covered with 2% water agar, 0,5 μ l drops of each compatible cell suspensions were placed 100 to 200 μ m apart. The pairs of drops were then covered with paraffin oil and incubated at 28°C. Wild-type combinations produced conjugation tubes that were directed towards the compatible partner. In contrast, $\Delta pcl12$ were severely impaired in the formation of conjugation tubes. Bar: 20 μ m

