

Fig. S1. *ec/3⁺* was induced by phosphate starvation in the absence of Zfs1. (A) JY333 (WT) and JZ974 ($\Delta zfs1$) (Kanoh et al. 1995) cells were grown in YE medium to OD₆₀₀ = 0.5, and the mRNA levels of *ec/3⁺* were measured by real-time PCR assay ($n = 3$). (B) JY333 (WT) and JZ974 ($\Delta zfs1$) cells were grown in EMM to OD₆₀₀ = 0.5 and then transferred into EMM without Na₂HPO₄ ($n = 3$).

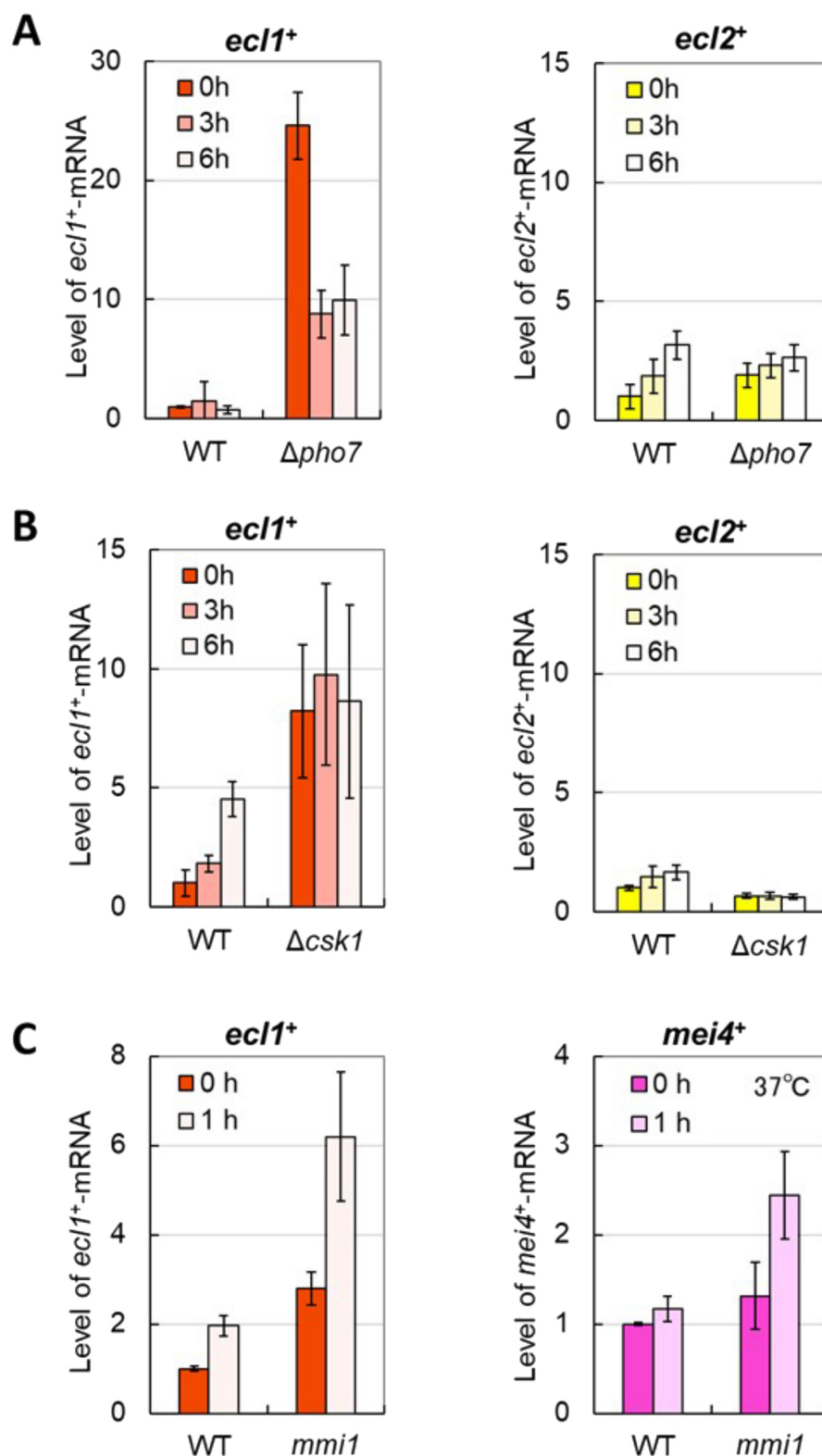


Fig. S2. The mRNA levels of *ecl1*⁺ and *ecl2*⁺ were measured by real-time PCR assay. (A) JY333 (WT) and JY333 $\Delta pho7$ ($\Delta pho7$) cells were grown in EMM to OD₆₀₀ = 0.5 and then transferred into EMM without Na₂HPO₄ ($n = 3$). (B) JY333 (WT) and JY333 $\Delta csk1$ ($\Delta csk1$) cells were grown in EMM to OD₆₀₀ = 0.5 and then transferred into EMM without Na₂HPO₄ ($n = 3$). (C) JY450 (WT) and JV579 (*mmi1*)(Yamashita et al. 2012) cells were grown in YE to OD₆₀₀ = 0.5 at 25°C and then transferred to 36°C for 1 h ($n = 3$). For cultures of JY333, JY333 $\Delta pho7$, JY333 $\Delta csk1$, JY450, and JV579 cells, 40 mg L⁻¹ adenine and 60 mg L⁻¹ leucine were added.

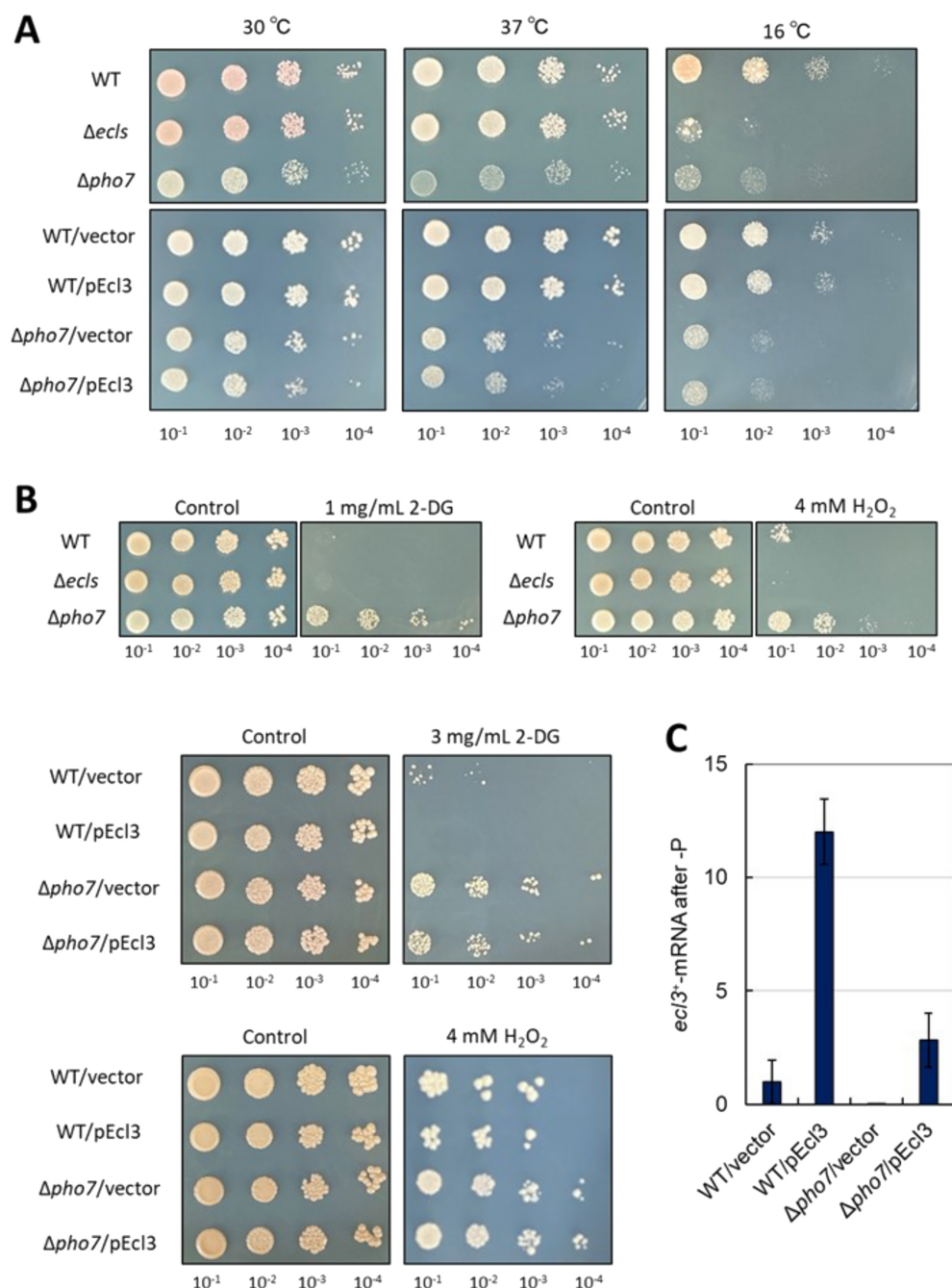


Fig. S3. Effects of *ecl* family genes and *pho7*⁺ on various stress sensitivities. (A) JY333 (WT), JY333 $\Delta ecl1/2/3$ ($\Delta ecl3$), and JY333 $\Delta pho7$ ($\Delta pho7$) cells were grown in YE to OD₆₀₀ = 1.0 and then spotted on YE plates with serial dilution (upper panel). JY333/pLB-Dblet (WT/vector), JY333/pEcl3 (WT/pEcl3), JY333 $\Delta pho7$ /pLB-Dblet ($\Delta pho7$ /vector), and JY333 $\Delta pho7$ /pEcl3 ($\Delta pho7$ /pEcl3) cells were grown in EMM to OD₆₀₀ = 1.0 and then spotted on EMM plate with serial dilution (lower panel). The plates were incubated at 16°C, 30°C, and 37°C. (B) JY333 (WT), JY333 $\Delta ecl1/2/3$ ($\Delta ecl3$), and JY333 $\Delta pho7$ ($\Delta pho7$) cells were grown in YE to OD₆₀₀ = 1.0 and then spotted on YE plates containing 2-DG or H₂O₂ with serial dilution (upper panel). JY333/pLB-Dblet (WT/vector), JY333/pEcl3 (WT/pEcl3), JY333 $\Delta pho7$ /pLB-Dblet ($\Delta pho7$ /vector), and JY333 $\Delta pho7$ /pEcl3 ($\Delta pho7$ /pEcl3) cells were grown in EMM to OD₆₀₀ = 1.0 and then spotted on EMM plates containing 2-DG or H₂O₂ with serial dilution (lower panel). (C) JY333/pLB-Dblet (WT/vector), JY333/pEcl3 (WT/pEcl3), JY333 $\Delta pho7$ /pLB-Dblet ($\Delta pho7$ /vector), and JY333 $\Delta pho7$ /pEcl3 ($\Delta pho7$ /pEcl3) cells were grown in EMM to OD₆₀₀ = 0.5. The mRNA level of *ecf3*⁺ was measured (*n* = 3).

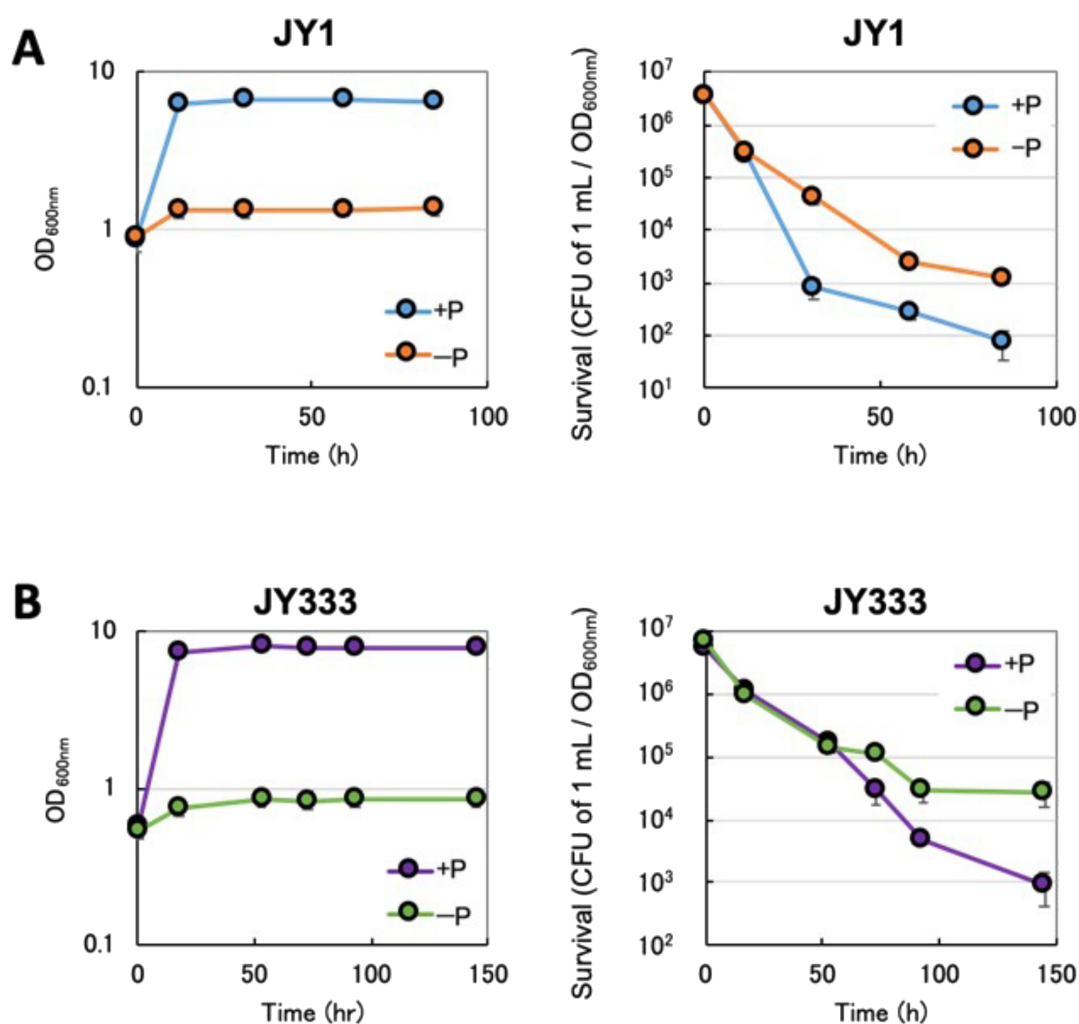


Fig. S4. The growth (left panels) and CLS (right panels) were measured during phosphate starvation. **(A)** JY1 cells were cultured in synthetic SD medium (Ohtsuka et al. 2017) with or without KH₂PO₄ ($n = 3$). **(B)** JY333 cells were cultured in synthetic SD medium with or without KH₂PO₄ ($n = 3$). Colony-forming units were used to measure survival (Ohtsuka et al. 2017).

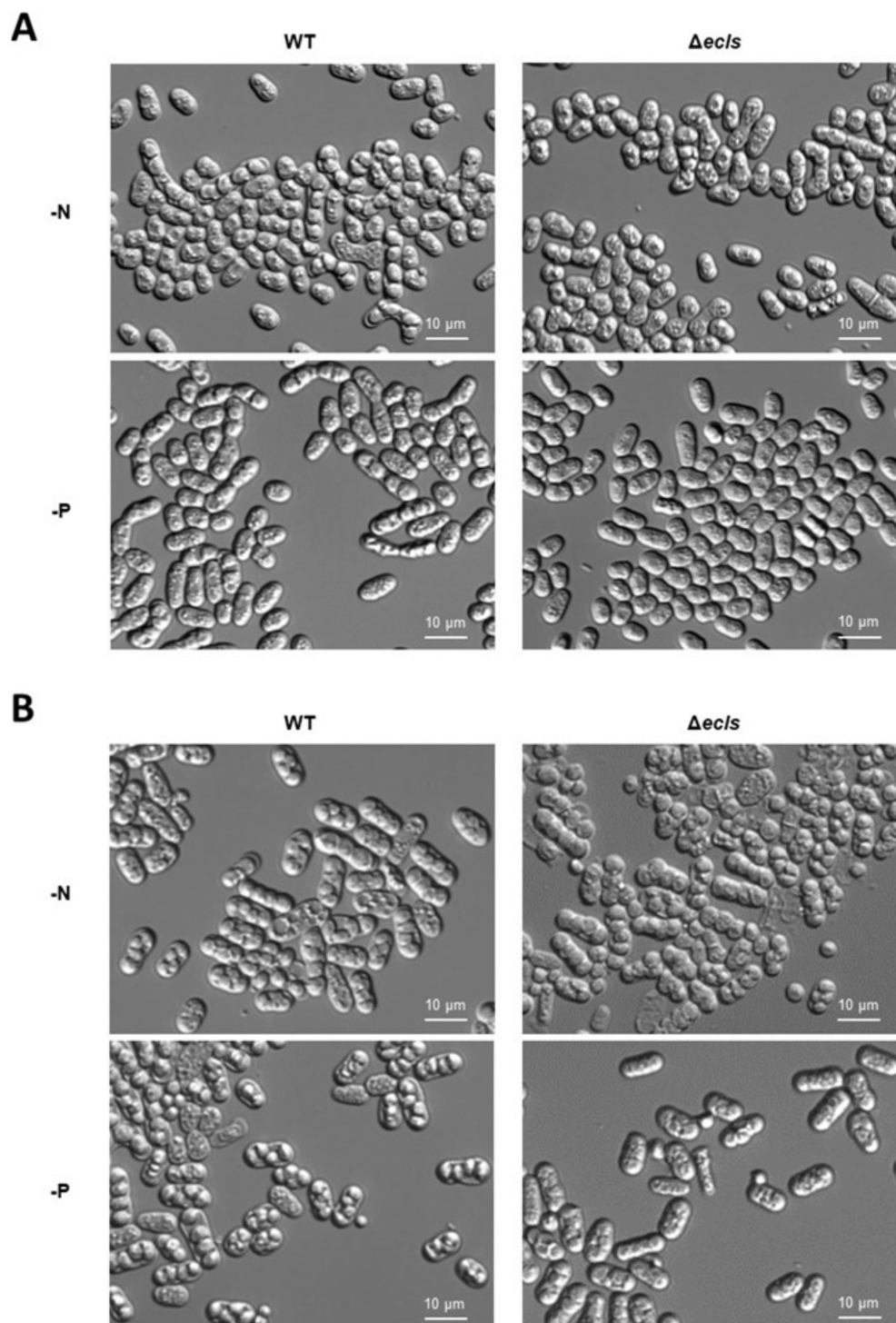


Fig. S5. Sexual development under phosphate depletion. (A) JY808 (WT) and JY808 $\Delta ecf1/2/3$ ($\Delta ecf1$) cells were grown in EMM to OD₆₀₀ = 0.5 and then transferred into nitrogen- and phosphate-depleted EMM for 25 h. (B) The diploid cells (JY333 \times HM3802) (WT) and (JY333 $\Delta ecf1/2/3$ and HM3802 $\Delta ecf1/2/3$) ($\Delta ecf1$) were grown in EMM to OD₆₀₀ = 0.5 and then transferred into nitrogen- and phosphate-depleted EMM for 25 h.

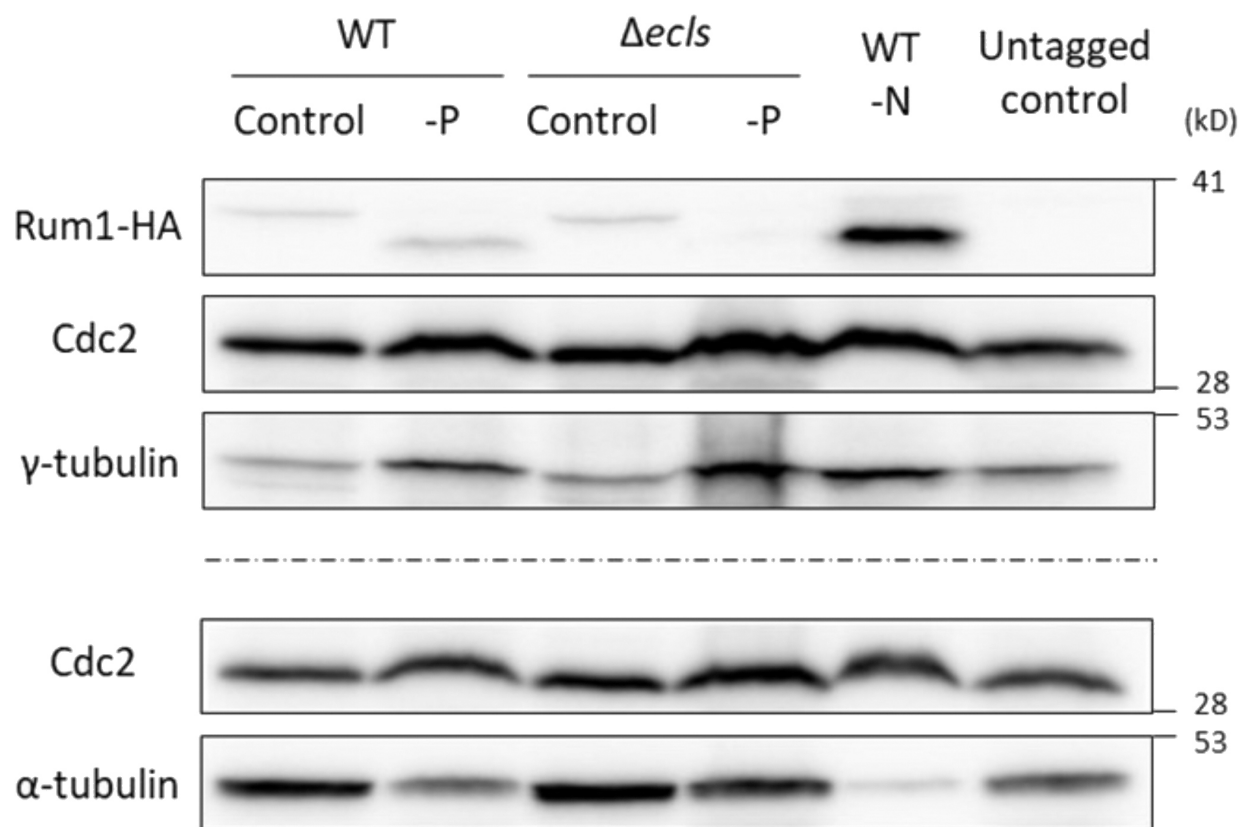


Fig. S6. Rum1 levels under phosphate depletion. WT (FY7288) and $\Delta ecls$ (FY7288 $\Delta ecl1/2/3$) were grown in EMM to OD₆₀₀ = 0.5 (Control) and then transferred into nitrogen-depleted (–N) or phosphate-depleted (–P) EMM (18 h). The level of Rum1-HA was measured using western blot assay. Both upper and lower bands of Rum1-HA were not observed in the untagged control, suggesting that these bands were derived from Rum1-HA.

Table S1. Yeast Strains and PCR Primers

[Click here to download Table S1](#)

References

- Kanoh, J., Sugimoto, A. and Yamamoto, M. (1995). *Schizosaccharomyces pombe* *zfs1* encoding a zinc-finger protein functions in the mating pheromone recognition pathway. *Mol. Biol. Cell* **6**, 1185-1195. doi:10.1091/mbc.6.9.1185
- Yamashita, A., Sakuno, T., Watanabe, Y. and Yamamoto, M. (2017). Analysis of *Schizosaccharomyces pombe* meiosis. *Cold Spring Harb. Protoc.* 2017, pdb.top079855. doi:10.1101/pdb.top079855