

Table 2 Transformation efficiency of non-frozen cells and cells after freezing with CaCl₂, using various yeast strains. Cells of all strains were frozen in 0.6 M sorbitol with 10 mM CaCl₂ and 10 mM Hepes, pH 7.5. Cell concentrations during the electric pulse were 1×10⁹ cells/ml and 2×10⁹ cells/ml for non-frozen cells and

cells after freezing with calcium, respectively. The electric pulse settings (for 25 μF, 200 Ω) were 10.0 kV/cm for *Sch. pombe* and 11.0 kV/cm for *Saccharomyces cerevisiae*, for both non-frozen cells and cells after freezing. Data represent averages ±SD for five independent experiments

Strain	Plasmid	Transformants/μg DNA	
		Non-frozen	After freezing with CaCl ₂
<i>Sch. pombe</i> (<i>leu</i> ⁻)	pAL7 (0.5 ng)	1.4 ± 0.6 × 10 ⁶	5.6 ± 1.0 × 10 ⁶
<i>Sch. pombe</i> (<i>ura</i> ⁻)	pAU5 (0.5 ng)	3.2 ± 2.9 × 10 ⁵	1.5 ± 1.1 × 10 ⁶
<i>Sch. pombe</i> TK107	pAL7 (10 ng)	2.8 ± 1.2 × 10 ⁴	4.3 ± 1.1 × 10 ⁴
<i>Sac. cerevisiae</i> AH22	YEp13 (5 ng)	1.7 ± 0.6 × 10 ⁵	4.2 ± 1.2 × 10 ⁵
<i>Sac. cerevisiae</i> D13-1A	YCp19 (2 ng)	2.7 ± 0.9 × 10 ⁵	5.9 ± 1.7 × 10 ⁵
<i>Sac. cerevisiae</i> 108-3C	YCp19 (100 ng)	2.1 ± 1.5 × 10 ²	3.9 ± 2.4 × 10 ²