

Self-Cloning Baker's Yeasts That Accumulate Proline Enhance Freeze Tolerance in Doughs[∇]

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We constructed self-cloning diploid baker's yeast strains by disrupting *PUT1*, encoding proline oxidase, and replacing the wild-type *PRO1*, encoding γ -glutamyl kinase, with a *pro1(D154N)* or *pro1(I150T)* allele. The resultant strains accumulated intracellular proline and retained higher-level fermentation abilities in the frozen doughs than the wild-type strain. These results suggest that proline-accumulating baker's yeast is suitable for frozen-dough baking.

Ordinary commercial baker's yeast (mostly strains of *Saccharomyces cerevisiae*) is generally susceptible to damage during the freeze-thaw process and frozen storage and does not retain sufficient leavening ability after thawing (8, 11). Freeze tolerance is a necessary characteristic of baker's yeast used in frozen doughs because postthaw leavening activity is essential prior to baking. It is known that the laboratory yeast *S. cerevisiae* induces trehalose synthesis during various stresses (1, 10, 34) and that induced trehalose functions as a stress protectant (7, 16, 23).

We previously found that proline and charged amino acids, such as arginine and glutamate, have cryoprotective activities nearly equal to that of glycerol or trehalose in baker's yeast (26). It was shown previously that yeast cells with higher levels of proline are more tolerant of freezing than wild-type cells (13, 14, 22, 26, 28, 30). Proline and trehalose were reported to preserve membrane structure and function during freezing (20). It has been suggested that proline can prevent ice nucleation and dehydration by forming strong hydrogen bonds with intracellular free water. γ -Glutamyl kinase (GK; the *PRO1* gene product) is the key enzyme in proline biosynthesis in *S. cerevisiae*, and GK activity is subjected to feedback inhibition by proline (22, 32). Proline oxidase (PO; the *PUT1* gene product) catalyzes the first step of the proline degradation pathway in *S. cerevisiae* (36). Interestingly, the D154N and I150T mutant GKs are less sensitive to proline feedback inhibition than the wild-type enzyme, and yeast cells expressing these mutated GKs accumulate proline and show higher tolerance for freezing than wild-type cells (22). With respect to industrial yeast, we previously constructed novel proline-accumulating sake yeast by disrupting the *PUT1* gene and introducing the *pro1(D154N)* mutant allele and found that the resultant strain accumulated proline and was more tolerant of ethanol stress than the control strain (29). Furthermore, self-cloning (SC) diploid sake yeasts that accumulate proline have been developed, and their fermentation profiles during sake brewing have

been analyzed (27). For the application of recombinant yeasts for commercial use, an SC yeast, which has no foreign genes or DNA sequences except for yeast DNA, may be more acceptable to consumers than a genetically modified (GM) yeast.

Our objectives in this study were (i) to construct SC diploid baker's yeast strains that accumulate proline and (ii) to determine if intracellular proline enhances the freeze tolerance of baker's yeast strains in doughs. We report here that baker's yeasts that accumulate proline showed increased leavening abilities in the frozen doughs.

Construction of the proline-accumulating baker's yeast. The baker's yeast strains 3346-ura3 and 3347-ura3 used in this study have only *URA3* as a useful marker gene. Therefore, we first integrated a mutated *pro1* gene by homologous recombination using *URA3* as the selection marker, excised the plasmid region containing the *URA3* marker, and then reused this marker with disruption of the *PUT1* gene according to the method in our previous study (27). In order to construct a *pro1(D154N)* or *pro1(I150T)* mutant strain, the linearized plasmid pRS406-D154NPRO1 or pRS406-I150TPRO1, which expresses the D154N or I150T mutant GK, respectively, with extreme desensitization to feedback inhibition by proline (22), was introduced into strain 3346-ura3 or 3347-ura3. The *Ura*⁺ transformants grown on synthetic complete medium lacking uracil (20 g of glucose/liter, 6.7 g of Bacto yeast nitrogen base without amino acids [Difco Laboratories, Detroit, MI]/liter, 2 g of dropout mixture/liter) were cultured in YPD medium (20 g of glucose/liter, 10 g of Bacto yeast extract/liter, 20 g of Bacto peptone/liter) at 30°C for 24 h with shaking, diluted in the same medium, and incubated for several days to obtain strains 3346-D154N-ura3, 3346-I150T-ura3, 3347-D154N-ura3, and 3347-I150T-ura3, which have excised the plasmid and lost one of the two copies of the duplicated region by homologous crossover, from 5-fluoroorotic acid-containing plates (4, 25). The overproduction of proline is believed to dilute the toxic proline analogue azetidine-2-carboxylate (AZC; Sigma-Aldrich, St. Louis, MO) (14, 26), which is incorporated into proteins in competition with proline (18, 33). We examined the growth of yeast strains on agar plates containing SD medium (20 g of glucose/liter, 6.7 g of Bacto yeast nitrogen base without amino acids/liter) with 2 mg of AZC/ml. These strains were capable of growing on SD medium-AZC plates due to proline

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TABLE 1. *S. cerevisiae* strains used in this study

Strain	Genotype	Background and/or description
3346	<i>MATa</i>	Haploid
3347	<i>MATα</i>	Haploid
3346-ura3	<i>MATa ura3</i>	Haploid
3347-ura3	<i>MATα ura3</i>	Haploid
3346-D154N-ura3	<i>MATa ura3 pro1(D154N)</i>	3346-ura3; <i>pro1(D154N)</i> mutant
3346-I150T-ura3	<i>MATa ura3 pro1(I150T)</i>	3346-ura3; <i>pro1(I150T)</i> mutant
3346-WT/Δput1	<i>MATa put1::URA3</i>	3346-ura3; <i>put1</i> disruptant
3346-D154N/Δput1	<i>MATa pro1(D154N) put1::URA3</i>	3346-D154N-ura3; <i>put1</i> disruptant
3346-I150T/Δput1	<i>MATa pro1(I150T) put1::URA3</i>	3346-I150T-ura3; <i>put1</i> disruptant
3347-D154N-ura3	<i>MATα ura3 pro1(D154N)</i>	3347-ura3; <i>pro1(D154N)</i> mutant
3347-I150T-ura3	<i>MATα ura3 pro1(I150T)</i>	3347-ura3; <i>pro1(I150T)</i> mutant
3347-WT/Δput1	<i>MATα put1::URA3</i>	3347-ura3; <i>put1</i> disruptant
3347-D154N/Δput1	<i>MATα pro1(D154N) put1::URA3</i>	3347-D154N-ura3; <i>put1</i> disruptant
3347-I150T/Δput1	<i>MATα pro1(I150T) put1::URA3</i>	3347-I150T-ura3; <i>put1</i> disruptant
AY13	<i>MATa/MATα</i>	Diploid derived from 3346 and 3347
PRO1-WT-ura3	<i>MATa/MATα ura3/ura3</i>	Diploid derived from 3346-ura3 and 3347-ura3
pro1-D154N-ura3	<i>MATa/MATα ura3/ura3 pro1(D154N)/pro1(D154N)</i>	Diploid derived from 3346-D154N-ura3 and 3347-D154N-ura3
pro1-I150T-ura3	<i>MATa/MATα ura3/ura3 pro1(I150T)/pro1(I150T)</i>	Diploid derived from 3346-I150T-ura3 and 3347-I150T-ura3
PRO1-WT/Δput1	<i>MATa/MATα put1/put1</i>	Diploid derived from 3346-WT/Δput1 and 3347-WT/Δput1
pro1-D154N/Δput1	<i>MATa/MATα put1/put1 pro1(D154N)/pro1(D154N)</i>	Diploid derived from 3346-D154N/Δput1 and 3347-D154N/Δput1
pro1-I150T/Δput1	<i>MATa/MATα put1/put1 pro1(I150T)/pro1(I150T)</i>	Diploid derived from 3346-I150T/Δput1 and 3347-I150T/Δput1

overproduction. We confirmed the correct gene replacement and the loss of the extraneous plasmid DNA by the direct sequencing of PCR products that contained the *PRO1* locus amplified from the chromosomal DNA of these strains.

To disrupt the *PUT1* gene on the chromosome of each strain, we amplified the *URA3* fragments which had 60-bp *PUT1* sequences at both ends by PCR performed with genomic DNA from strains 3346 and 3347 using the PUT1disURA3(+) and PUT1disURA3(-) primers. The sequence of the forward primer [PUT1disURA3(+)] was 5'-GCT TGC TGG GAA CCG AAC ACA AAC TCC ACA AGT CCG TAG CAG CTC TTC TCT TTT GTC TTT AAT GTG GCT GTG GTT TCA GG-3', and the sequence of the reverse primer [PUT1disURA3(-)] was 5'-GCT ATG GCC TTG ATT AAT GGC CAG CCA TTA TCA GAT CTC ACA GCA TCC CCG TTT TCT TGC GTT CTG GCG AGG TAT TGG AT-3' (the underlining indicates the sequence 50 bp upstream of the ATG initiation codon and the sequence 30 bp upstream of the TGA termination codon of the *PUT1* gene, respectively). The sequences of *PUT1* and *URA3* were based on the sequences corresponding to GenBank accession numbers M18107 and K02206, respectively. A unique 1.24-kb amplified band containing the *URA3* gene was purified and then integrated into the *PUT1* locus in strains 3346-D154N-ura3, 3346-I150T-ura3, 3347-D154N-ura3, and 3347-I150T-ura3 by transformation. The transformants, which exhibited the Ura⁺ phenotype, were selected, and we confirmed that these strains failed to grow on SD medium containing proline (1 g/liter), instead of ammonium sulfate, as the sole source of nitrogen (data not shown). As a result, we constructed the following six haploid strains by an SC method: 3346-D154N/Δput1, 3346-I150T/Δput1, 3347-D154N/Δput1, 3347-I150T/Δput1, 3346-WT/Δput1, and 3347-WT/Δput1. As controls, strains 3346-WT/Δput1 and 3347-WT/Δput1 were constructed from strains 3346-ura3 and 3347-ura3, respectively.

In general, commercial baker's yeast is a diploid or polyploid strain. For industrial applications, the diploid strain is preferable to the haploid strain in terms of its growth characteristics

and fermentation abilities (17, 21). For application to dough fermentation and freeze tolerance in doughs, six diploid strains, PRO1-WT-ura3, pro1-D154N-ura3, pro1-I150T-ura3, PRO1-WT/Δput1, pro1-D154N/Δput1, and pro1-I150T/Δput1, were constructed by mating 3346-ura3 and 3347-ura3, 3346-D154N-ura3 and 3347-D154N-ura3, 3346-I150T-ura3 and 3347-I150T-ura3, 3346-WT/Δput1 and 3347-WT/Δput1, 3346-D154N/Δput1 and 3347-D154N/Δput1, and 3346-I150T/Δput1 and 3347-I150T/Δput1, respectively. When the zygote formation was observed with a microscope, the culture was plated onto YPD plates. The larger colonies were selected, and the cells were confirmed to sporulate on sporulation medium at 25°C (19). These diploid strains will be investigated for approval as SC strains, due to a lack of extraneous DNA derived from other organisms (Table 1). As expected, PRO1-WT/Δput1, pro1-D154N/Δput1, and pro1-I150T/Δput1 failed to grow on proline-containing plates, while strains carrying the wild-type *PUT1* gene (AY13, PRO1-WT-ura3, pro1-D154N-ura3, and pro1-I150T-ura3) could utilize proline as their nitrogen source (data not shown). Furthermore, we measured the activity of PO, which is the *PUT1* gene product (3, 13). The *put1*-disrupted strains, PRO1-WT/Δput1, pro1-D154N/Δput1, and pro1-I150T/Δput1, exhibited reduced PO activity (>0.2 U/mg of protein), although such activity was readily detected in AY13, PRO1-WT-ura3, pro1-D154N-ura3, and pro1-I150T-ura3 (1.97, 1.23, 1.98, and 1.04 U/mg of protein, respectively). These results showed that the disruption of *PUT1* was confirmed on the basis of gene construction, proline utilization, and enzymatic activity. Next, we examined the growth of diploid strains on plates containing SD medium with AZC (2 mg/ml). Strains pro1-D154N-ura3, pro1-I150T-ura3, pro1-D154N/Δput1, and pro1-I150T/Δput1 clearly showed AZC resistance, whereas the rest of the strains were sensitive to AZC (data not shown). To examine whether greater resistance to AZC reflects a higher level of intracellular proline, AY13, PRO1-WT/Δput1, pro1-D154N/Δput1, and pro1-I150T/Δput1 were cultivated in SD liquid medium and the cellular proline

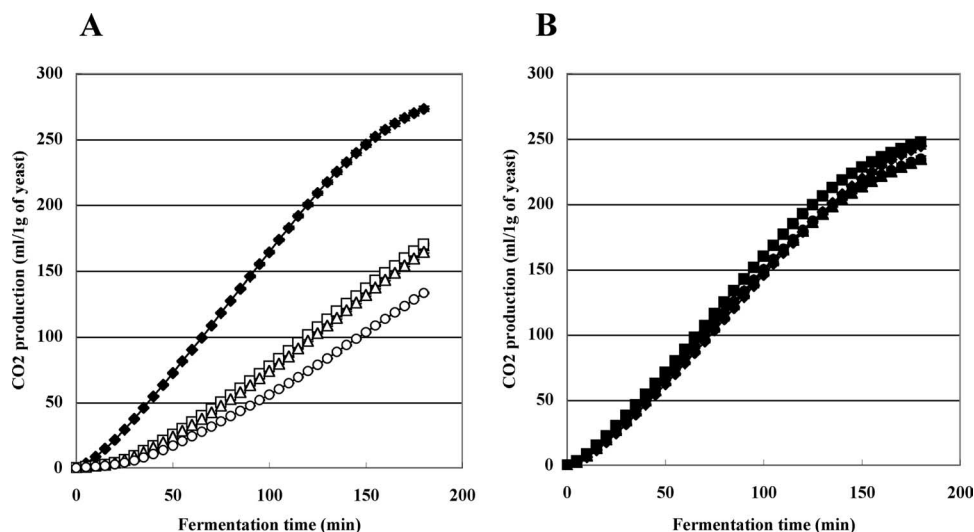


FIG. 1. Fermentation abilities of the diploid baker's yeast strains. The bread doughs containing AY13 (closed diamonds), PRO1-WT-ura3 (open squares), pro1-D154N-ura3 (open circles), and pro1-I150T-ura3 (open triangles) (A) or AY13 (closed diamonds), PRO1-WT/ Δ put1 (closed squares), pro1-D154N/ Δ put1 (closed circles), and pro1-I150T/ Δ put1 (closed triangles) (B) were prepared, and then CO₂ gas production was monitored every 5 min for 180 min. The data shown are from one experiment. Similar results were seen in replicates of this experiment.

levels were examined. The proline content in PRO1-WT/ Δ put1 (0.069% of the dry weight of cells) was approximately threefold that in the control strain AY13 (0.023%), due to the deficiency of the degradation pathway (28). In agreement with the findings of our previous studies (22, 30), there was significant accumulation of proline in cells of pro1-D154N/ Δ put1 and pro1-I150T/ Δ put1 (0.32 and 0.73%, respectively). GK activity could be detected in the crude extracts prepared from yeast cells, but all GKs, even the wild-type enzyme, showed little inhibition in the presence of 50 mM proline (data not shown). Recently, we found that the purified GK is subject to feedback inhibition by proline and that the D154N and I150T mutant GKs are less sensitive than the wild-type enzyme (22). In fact, pro1-D154N/ Δ put1 and pro1-I150T/ Δ put1 clearly accumulated larger amounts of proline than PRO1-WT/ Δ put1 in both SD medium and cane molasses medium (30 g of sugar [calculated as sucrose]/liter, 1.93 g of urea/liter, and 0.46 g of KH₂PO₄/liter), in proportion to the diminished sensitivities to feedback inhibition by proline. Although the reason for the absence of feedback inhibition in the crude extracts is unclear, we consider that the D154N and I150T mutant GKs are functionally capable of overproducing proline *in vivo*. These results showed that the baker's yeast strains that carried the mutant *PRO1* gene *pro1(D154N)* or *pro1(I150T)* and had a disrupted *PUT1* gene accumulated large amounts of proline, in a manner similar to that of the laboratory or sake yeast strains (22, 27).

Fermentation ability of the proline-accumulating baker's yeast. We tested the fermentation abilities of diploid baker's yeast strains that accumulate proline for the application of bread baking. To determine the CO₂ gas production abilities as indicators of fermentation abilities, seven diploid strains were cultured in cane molasses medium as a carbon source, which simulates the commercial baker's yeast production process, and the amount of CO₂ produced by the yeast cells in the dough was measured according to the method of Nishida et al. (15) (Fig. 1). The fermentation abilities differed between the two groups, the Ura⁻ strains and the Ura⁺ strains. As shown

in Fig. 1A, the gassing powers of the Ura⁻ strains, PRO1-WT-ura3, pro1-D154N-ura3, and pro1-I150T-ura3, were approximately 40 to 50% lower than that of the Ura⁺ strain AY13. In contrast, the Ura⁺ strains PRO1-WT/ Δ put1, pro1-D154N/ Δ put1, and pro1-I150T/ Δ put1 exhibited fermentation abilities equivalent to that of strain AY13 (Fig. 1B). Incompatible auxotrophic markers among the strains probably affected the cell growth (5). Therefore, we used four Ura⁺ strains, AY13, PRO1-WT/ Δ put1, pro1-D154N/ Δ put1, and pro1-I150T/ Δ put1, for further analyses.

Amino acid content and stress tolerance of the proline-accumulating baker's yeast. After cultivation in the molasses medium, the amounts of intracellular amino acids in the baker's yeast strains were analyzed by a method described previously (10, 22). As on the SD medium, the *put1*-disrupted strain PRO1-WT/ Δ put1 contained a higher proline level (0.20%) than the control strain AY13 (0.096%). Strains pro1-D154N/ Δ put1 (3.4%) and pro1-I150T/ Δ put1 (6.3%) accumulated much higher proline levels than AY13, as expected from the above-described results. In contrast, the proline-accumulating strains showed slightly decreased levels of glutamate and arginine, both of which are proline precursors, compared to the wild-type strain (data not shown).

We previously found that yeast strains that accumulate proline are tolerant of osmotic stress factors, such as NaCl and sorbitol, in addition to freezing stress (22, 26). Yeasts used in bread making are exposed to high concentrations of sucrose during sweet-dough fermentation (2). Such high sucrose concentrations exert severe osmotic stress on yeast (35). Therefore, the growth phenotypes of the proline-accumulating baker's yeast strains under freezing and high-sucrose stress conditions were examined. As shown in Fig. 2, the proline-accumulating strains PRO1-WT/ Δ put1, pro1-D154N/ Δ put1, and pro1-I150T/ Δ put1 were more resistant to high-sucrose stress than AY13. Contrary to our expectations, however, there were no significant differences in growth between the proline-

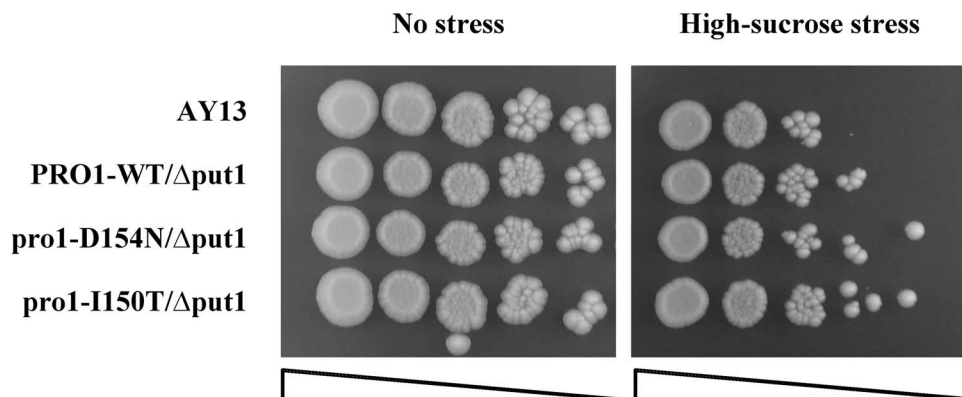


FIG. 2. Growth phenotypes of the diploid baker's yeast strains under sucrose stress. Yeast cells were precultured for 1 day at 30°C in 50 ml of SD medium. Harvested cells were washed, suspended in modified liquid fermentation medium containing (per liter) 500 g of sucrose and 50 g of maltose without glucose (7) at an OD_{600} of 6 (no stress), and then cultured for 4 days at 30°C with shaking (high-sucrose stress). Approximately 10^7 cells of each strain and 10^{-1} to 10^{-4} serial dilutions (from left to right, as indicated by triangles) were spotted onto YPD agar plates, and the plates were incubated at 30°C for 2 days.

accumulating strains and AY13 under freezing stress conditions (data not shown).

Freeze tolerance of the proline-accumulating baker's yeast in dough. Because freeze tolerance is an important characteristic in recent frozen-dough baking processes (2), we assayed the freeze tolerance of the proline-accumulating baker's yeast in doughs. The formula of dough was 100 g of bread-making flour, 5 g of sucrose, 2 g of NaCl, 4 g of yeast (66% moisture basis), and 68 ml of water. The ingredients were mixed for 3 min with a Swanson type mixer (National Manufacturing Co., Ltd., Sterling, IL). Mixed dough was divided into pieces (40 g each), placed into screw-cap bottles, stored at -20°C without prefermentation before freezing, and kept frozen for 3 weeks. The frozen dough was thawed for 30 min at 30°C, and the fermentation ability was assayed by measuring CO_2 gas production using a Fermograph AF-1000 (Atto, Tokyo, Japan). The freeze tolerance was expressed as the percentage of fermentation ability remaining after freezing relative to the ability before freezing (Fig. 3A). We found that pro1-D154N/ Δput1 and pro1-I150T/ Δput1 retained their fermentation abilities at

the same levels as those before freezing, although that of the parent strain AY13 fell to 85% of the prefreezing level. The trehalose contents of AY13, PRO1-WT/ Δput1 , pro1-D154N/ Δput1 , and pro1-I150T/ Δput1 after cultivation in the molasses medium were 2.7, 4.8, 4.9, and 3.5% of the dry weight, respectively. It appears that there is a weak correlation between the trehalose content and freeze tolerance, because PRO1-WT/ Δput1 , which contained more trehalose than AY13, exhibited the same level of freeze tolerance as AY13 (Fig. 3A). It is known that yeast cells do not accumulate trehalose in response to freezing stress (12). As for proline, there were no significant differences in freeze tolerance in doughs among the proline-accumulating strains. These results suggest that an appropriate proline level in yeast cells is important to induce a protective effect against freeze-thaw stress.

In commercial frozen-dough processes, prefermentation before freezing is desirable in terms of the texture and taste of the product (9, 31). We therefore performed the same experiment using the doughs prefermented for 120 min at 30°C before freezing, and the dough was kept frozen for 9 days (Fig.

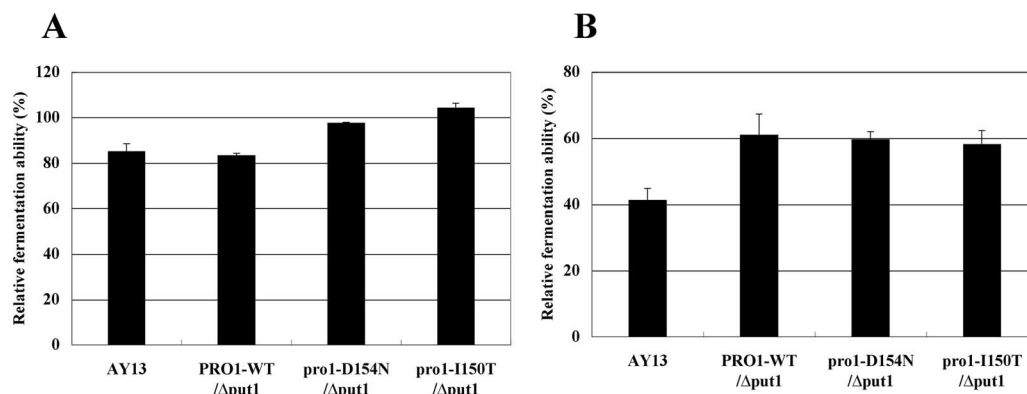


FIG. 3. Freeze tolerances of the diploid baker's yeast strains in doughs. Frozen doughs were prepared by using AY13, PRO1-WT/ Δput1 , pro1-D154N/ Δput1 , and pro1-I150T/ Δput1 . The doughs were not prefermented and were frozen at -20°C for 3 weeks (A) or were prefermented for 120 min at 30°C and then frozen at -20°C for 9 days (B). The frozen dough was thawed for 30 min at 30°C, and the remaining CO_2 gas production was measured. The gassing power before freezing was defined as 100%. The values are the means and standard deviations of results from three independent experiments.

3B). The remaining gassing power of AY13 was dramatically decreased to 41% of that before freezing. It is noteworthy that pro1-D154N/ Δ put1 and pro1-I150T/ Δ put1, and even PRO1-WT/ Δ put1, showed approximately 50% greater fermentation abilities than AY13. Prefermentation is an important process in bread baking, because yeast cells activated during prefermentation produce the metabolites, such as alcohols and organic acids, which influence the taste and flavor of the bread. The reason for the loss of the gassing power remains unclear; however, it is possible that prolonged prefermentation causes serious damage to the membranes of the yeast cells in the dough (11). It is also known that the intracellular amino acid pool decreases through prefermentation (24). Therefore, we consider that intracellular proline, as a cryoprotectant, was not degraded in PRO1-WT/ Δ put1 during prefermentation, leading to improved tolerance for freezing. These data strongly suggest that proline-accumulating baker's yeast has higher-level freeze tolerance than non-proline-accumulating yeast and is suitable for frozen-dough baking.

The present study is the first to report the construction of diploid baker's yeasts that accumulate proline. It is relatively difficult to breed an industrial baker's yeast strain with higher-level freeze tolerance than that of a laboratory strain. The process that involves adding proline externally to the cell or to the dough remains somewhat troublesome for practical applications. However, yeast strains with comparatively large amounts of proline already in the cells may overcome this problem. Another potential problem is that the strains mentioned above are GM yeasts, because the *S. cerevisiae*-*Escherichia coli* shuttle vector was used to complement the auxotrophic marker. According to the Japanese government guidelines, SC processes are considered to be the same as naturally occurring gene conversions, such as recombination, deletion, and transposition; thus, SC yeast need not be treated as recombinant GM yeast (6).

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