



## Role of *Schizosaccharomyces pombe git1* gene in oxidative stress response

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### Abstract

Glucose is the preferred energy and carbon source for many organisms. Glucose sensing and signal transduction in yeast is generally accomplished through a system of heterotrimeric G-protein and G protein-coupled cell surface receptors. Signal transduction pathway to cAMP/protein kinase is activated with glucose sensing. In this pathway, *Git1* is a C2 domain protein that is directly linked to adenylate cyclase and it is one of the 6 proteins required for the activation of adenylate cyclase. The 3' end of *git1* gene contains "Mammalian uncoordinated homology 13, domain 2." It plays role membrane trafficking, exocytosis, vesicle secretion. This study aims to find out whether the *git1* gene, which is one of the genes involved in glucose signaling, and the 3' end of *git1* gene, are related to oxidative stress response. In this study, *Schizosaccharomyces pombe* wild type (972h-) and *git1*- (*git1Δ*) mutant with *Escherichia coli DH5α* were used. Genomic DNA of *S. pombe* 972h- was used as a template to obtain *git1* and 3' deletion *git1* genes. These genes were cloned into plasmid pSGP572 containing the GFP reporter gene in the cloning site. The resulting recombinant vectors were transfected into super-efficient *E. coli DH5α* and then isolated. These isolated vectors were transformed into the *S. pombe git1Δ* mutant. Cell morphologies of transformants in the selective media were stained DAPI and then examined under confocal microscope. Transformants carrying recombinant plasmids were confirmed by GFP luminescence detected in a confocal microscope. There was no statistically significant difference in superoxide dismutase and catalase enzyme activities in H<sub>2</sub>O<sub>2</sub> induced oxidative stress conditions in *S. pombe* recombinants and *S. pombe git1Δ* mutant. These results make think that cells probably select the different pathway alternatives in the stress response.

**Keywords:** *Schizosaccharomyces pombe*, *git1* gene, oxidative stress, glucose metabolism.