

The antimutagen capacity of synthesized polyphenol glycoside through transglycosylation by CGTase enzyme of bacilluspolymyxa D4

Abstract

Polyphenol glycoside was synthesized through enzymatic transglycosylation by cyclodextrin glucanotransferase (1,4- α -D-glucan 4- α -D-1,4-glucano-transferase) or CGTase EC 2.4.1.19, of *Bacillus polymyxa* D4. Soluble starch and resorcinol were used as the substrate and the acceptor respectively. The transfer product was detected using thin layer chromatography as resorcinol glucoside. Purification of transfer product was carried out using column chromatography and resorcinol glucoside was collected in fraction of 20% methanol. The bioassay of mutagenesis was detected by formation of mutation induced by aflatoxin B₁ 1 μ g/ml in *Salmonella typhimurium* TA98. The effect of antimutagenesis was evaluated using this culture on L-histidine deficiency medium containing resorcinol glucoside. Results show that resorcinol glucoside like arbutin and resorcinol can inhibit mutagenesis at concentration of 25 mM.