

Development and proteom analysis of recombinant *E.coli* strain Crds-C(836bp) which produces soluble beta-1,3-glucan

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Polysaccharides, particularly glucans, have a long history as immuno-modulators¹). β -1,3-glucans have been found to induce the activation of different defence processes in both plants and animal²). β -1,3-glucans are a heterogeneous group of glucose polymers found in the cell walls of plant, yeast, bacteria and fungi.

Following β -1,3-glucan synthesis metabolic pathway, UDP-glucose is the exclusive precursor for the synthesis of β -1,3-glucan. To produce β -1,3-glucan in *E. coli*, recombinant *E. coli* strains carrying β -1,3-glucan synthase genes and producing β -1,3-glucan synthase which catalyzed the reaction from UDP-glucose to β -1,3-glucan were developed. As a result, recombinant *E. coli* strains[Crds-F(1964bp), Crds-C(836bp) and CN-termH6(1124bp), CC-termH6(1676bp)] were obtained and their glucan production characteristics were studied. *E.coli* mutant strain Crds-C(836bp) produces amount 5.9g/L soluble β -1,3-glucan. And then we analyzed the proteome of the *E.coli* mutant strain Crds-C(836bp) with 2D-electrophoresis.

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