

## Cloning, Expression and Preliminary Characterization of the perosamine synthase Gene *per* from *E. coli* O157

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### Abstract

The *per* gene, encoding a perosamine synthase enzyme, has been cloned from *E. coli* O15. Per can synthesize GDP-D-perosamine from GDP-4-keto-6-deoxy-D-mannose. The D-perosamine is found as a main constituent of LPS in *E. coli* O157, *Brucella* spp., and *Vibrio cholerae* O1. And it is the intermediate of antibiotic perimycin of *Streptomyces coelicolor* var. *aminophilus*. In this study, Per from *E. coli* O15 was expressed and characterized. The molecular mass of His<sub>6</sub>-tagged Per, as determined by SDS-PAGE, was 44 kDa. The optimal temperature was 37 °C, and optimal pH was 7.5. It showed the high activity to L-glutamate, L-glutamine, L-aspartate, L-asparagine, and L-alanine as amino donor.

### Reference

1. Albermann, C., and W. Piepersberg. 2001. Expression and identification of the RfbE protein from *Vibrio cholerae* O1 and its use for the enzymatic synthesis of GDP-D-perosamine. *Glycobiology* 11:655-661.
2. Awram, P and J. Smit. 2001. Identification of lipopolysaccharide O antigen synthesis genes requires for attachment of the S-layer of *Caulobacter crescentus*. *Microbiology* 147:1451-1460.