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Expression of cspH, encoding a member of major cold shock protein CspA family in Salmonella typhimurium UK-1.

Bae Hoon Kim and Yong Keun Park Graduate School of Biotechnology, Korea University

In E.coli, cold shock proteins(Csp family) containing two cold-box domains(homologous to Y-box in eukaryotes: it has nucleotide binding property) at its amino acid sequence have been well known about induction conditions(at cold(10-25°C), stationary phase, starvation, etc) and functions (RNA chaperone, etc). Csp family known in E.coli is 9 proteins from CspA to CspI. But, in Salmonella typhimurium, scarcely has Csp family been known except cspA and cspB. The cspH, one of the Csp family, has not been known about an expression condition and its functions in both E.coli and S.typhimurium. In this study, we analyzed the promoter sequence of cspH between E.coli and S.typhimurium, and found the very different sequence composition. After the sequence analysis, we investigated the patterns of expression of its gene in S.typhimurium with Northern hybridization, translational promoter-lacZ fusion (pRS414), etc. And then, we found that like E.coli CspA, CspH is induced after temperature downshift(approximately 15°C) only in log phase(not induced in stationary phase), and expressed during growth under non-stress condition. We also found the transcriptional start site of cspH by primer extension analysis. Extraordinary, the untranslated region(UTR) of its gene is very short unlike cold shock genes of E.coli. Judging from our experiments, We suggest that the CspH may be required not only for adaptation at cold temperature but also efficient growth in early log phase. We also demonstrate that because of very short UTR, cspH in S.typhimurium may be regulated by transcriptional level.

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The snpA gene, a suppressor of npgA1, encodes a translation termination factor, RF1, of Aspergillus nidulans

Seong-Soo Cheong, Kyu-Yong Han, Young-Jea Jeong and Dong-Min Han Division of Life Science, College of Natural Sciences, Wonkwang University, Iksan 570-749, Korea

The *snpA* gene, originally identified as a suppressor of *npgA1* mutation which is defective in the arrangement of outer layer of cell wall and thus eventually results in the lack of pigments in hyphae, conidia and ascospores, has been known to be involved in differentiation since a temperature sensitive allele, *snpA6*, caused pleiotrophic phenotypes in growth and differentiation at high temperature. We cloned the gene from chromosome specific library III by complementation assay. The full complementing activity was retained in 2.5 kb EcoRV fragment. Nucleotide sequence analysis revealed that the gene encoded a translation termination factor, RF1. It showed a high similarity to that of *Podospora anserina*, higher than 80% in part. Mutation phenotypes of *snpA6* showed also great similarity to those in RF1 genes of other fungi, *Saccharomyces cerevisiae* and *P. anserina.*, which were identified as omni-potent suppressor and caused also pleiotrophic phenotypes related to differentiation, respectively.