

E337Purification and Characterization of Superoxide Dismutases
from *S. coelicolor*김 은자*, 김 형표, 하 영칠, 노 정혜
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S. coelicolor (Müller) contains two distinct superoxide dismutase (SOD) activities detected on native PAGE. The level of each SOD changed differently depending on growth media, and scarcely responded to paraquat, a superoxide-generating agent. These SODs have been purified from *S. coelicolor*.

One SOD (SOD1) was a tetramer of 13.5 kD subunits. It was found to be a novel SOD containing 0.75 atom of Ni per subunit as determined by atomic absorption spectroscopy. The other SOD (SOD2) was composed of 27.5 kD subunits. It was found to contain 0.4 to 0.5 atom of Fe and about 0.3 atom of Zn per subunit. Electron spin resonance spectroscopy also revealed the existence of Ni and Fe as cofactors of SOD1 and SOD2, respectively.

The N-terminal amino acid sequences of both SODs were determined. SOD2 showed high similarity to MnSOD and FeSOD from *E. coli*. SOD1 was less similar to known SODs but still contained several conserved amino acids.

E338Production and Characterization of Cathepsin B
Inhibitor Produced from *Streptomyces aburabiensis* S175박상진*, 이현숙, 최영출, 김인섭, 이계준
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The aim of the present study was to develop strains of actinomycetes producing low molecular weight cathepsin B inhibitor. An economical and effective screening method was developed for the screening of cathepsin B inhibitor. Among 700 isolates isolated from soil samples, a strain of *Streptomyces aburabiensis* S175 was selected. Optimum culture condition for the production of cathepsin B inhibitor was determined. Glucose and soytone were selected as good carbon and nitrogen sources, respectively. The production of inhibitor was related with mycelial growth. The inhibitor in culture filtrate was purified by adsorption on activated charcoal, butanol extraction, silica gel chromatography, ion exchange using Dowex-1 (Cl⁻ form) and Amberlite IRC-50 (H⁺ form), and preparative TLC. The physico-chemical characteristics were studied and the chemical structure of the inhibitor is being determined from UV spectroscopic analysis, IR spectroscopic analysis, Mass spectroscopic analysis, ¹H-NMR, and ¹³C-NMR studies.