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蛋白质组学方法分析不同苯酚浓度下菌株Acinetobacter sp. EDP3的应激机理

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摘要 Strain EDP3 was isolated from an industrial-activated sludge. It belonged to the ga group of Proteobacteria with an identity of 97.0% to Acinetobacter calcoaceticus accordi the 16S rRNA gene sequences. It can tolerate up to 1000mg•L-1 phenol at roor temperature with a much longer lag phase. This indicates that higher phenol concentrat has induced some physiological and genotypic changes in the bacterium. The aim of this is, therefore, to investigate these responses to phenol concentration variations in strain EDP3. Proteome analysis is conducted by means of a two-dimensional polyacrylamide ge electrophoresis (2D PAGE) and matrix-assisted laser desorption ionization time of flight r spectrometry (MALDI-TOF-MS) was conducted to obtain a deeper insight into the adaptiv responses inside the bacterium. Comparative analysis of the proteome profiles of strain grown in 400mg• L-1 and 1000mg• L-1 phenol allowed us to identify that among all the proteins up-regulated under the higher phenol concentration, oxidative st proteins were dominant. The synthesis of a heat shock protein, 60000 chaperonin GroEl also amplified. In addition, the expression of one membrane protein, adenosine 5'triphosphate (ATP)-binding cassette (ABC) type sugar transporter, was found up-regulated The inhibition of adenosine 5'-triphosphate (ATP) and RNA/protein synthesis was also observed.

关键词 <u>adaptation phenol-degrading bacteria</u> <u>Acinetobacter sp.</u> proteome 分类号

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## Proteome analysis of the adaptation of a phenol-degrading bacterium *Acinetobacter* sp. EDP3 to the variation of phenol loadings

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Abstract Strain EDP3 was isolated from an industrial-activated sludge. It belonged to the gamma group of Proteobacteria with an identity of 97.0% to Acinetobacter calcoaceticus according to the 16S rRNA gene sequences. It can tolerate up to 1000mg•L-1 phenol at room temperature with a much longer lag phase. This indicates that higher phenol concentration has induced some physiological and genotypic changes in the bacterium. The aim of this study is, therefore, to investigate these responses to phenol concentration variations in strain EDP3. Proteome analysis is conducted by means of a two-dimensional polyacrylamide gel electrophoresis (2D PAGE) and matrix-assisted laser desorption ionization time of flight mass spectrometry (MALDI-TOF-MS) was conducted to obtain a deeper insight into the adaptive responses inside the bacterium. Comparative analysis of the proteome profiles of strain EDP3 grown in 400mg•L-1 and 1000mg•L-1 phenol allowed us to identify that among all the proteins up-regulated under the higher phenol concentration, oxidative stress proteins were dominant. The synthesis of a heat shock protein, 60000 chaperonin GroEL, was also amplified. In addition, the expression of one membrane protein, adenosine 5'-triphosphate (ATP)-binding cassette (ABC) type sugar transporter, was found up-regulated. The inhibition of adenosine 5'-triphosphate (ATP) and RNA/protein synthesis was also observed.

Key words adaptation; phenol-degrading bacteria; Acinetobacter sp.; proteome

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