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Efficient Extraction of Ferulic Acid from Sugar Beet Pulp Using the Culture Supernatant of *Penicillium chrysogenum*

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We found a microorganism, Penicillium chrysogenum 31B, that has high ability to release ferulic acid from sugar beet pulp. Approximately 85% of alkaline-extractable ferulic acid in sugar beet pulp could be released using the culture supernatant of *P. chrysogenum* 31B. However, the culture supernatant did not efficiently extract ferulic acid from wheat bran, peel of corn seed, or sugar-cane bagasse. A ferulic acid esterase (FAE-1) was purified from the culture filtrate of *P. chrysogenum* 31B. The molecular mass of the enzyme was determined to be 62 kDa by SDS-PAGE. Optimum conditions for enzyme activity were 50°C and pH 6-7. The enzyme showed activity towards methyl esters of hydroxycinnamic acids including ferulic acid, p-coumaric acid, and caffeic acid, but was not active on methyl sinapinate or 3,4-dimethoxy cinnamate. The lack of activity of FAE-1 toward these substrates appears to be due to the presence of two methoxy groups on the benzene ring. The substrate specificity of FAE-1 seemed to be similar to that of ferulic acid esterase (CinnAE) of Aspergillus niger. However, there was a difference between FAE-1 and CinnAE in respect to activity towards methyl vanillate. It is remarkable that FAE-1 hydrolyzed methyl vanillate, which, to our knowledge, is the first report of a ferulic acid esterase hydrolyzing a hydroxybenzoic acid methyl ester.

Key words: ferulic acid, ferulic acid esterase, Penicillium chrysogenum, sugar beet pulp



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