

# Purification and properties of fatty acid esterases from yam (*Dioscorea batatas* Decne) tuber

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## Abstract

ng  $\beta$ -naphthyl myristate (C14 fatty acid ester) as a screening substrate, fatty acid esterases (FAEs) were purified from *D. batatas* tubers. Two FAE fractions (named FAE I and FAE II) were obtained after DE-52 ion exchange chromatography and Sephadex G-75 gel filtration purification steps, and then further purified by Con A Sepharose 4B affinity chromatography. The FAE I and II fractions contained the same 3 protein bands of about 50-64 kDa, corresponding to esterase activity bands on SDS-PAGE gels. Among  $\beta$ -naphthyl esters determined at pH 4.0, 5.0 and 6.0 the best substrate for both FAE fractions was a C10-containing one with a maximum pH at 5.0. The  $K_m$  and  $V_{max}$  for  $\beta$ -naphthyl caprate (C10 fatty acid ester) of FAE I and II at 37°C and pH 5.0 were 0.338 and 0.959 mM and 0.405 and 0.585 nmole  $\beta$ -naphthol/min  $\mu$ g protein, respectively. FAE activity was stable below 50°C and lost completely above 65°C.