Year: February 1987.

Abstract

Out of 217 fungal isolates, Absidia sp. strain BA16 was found to be able to convert lithocholic acid into a derivative which had similar characteristics on thin layer chromatography

acid, the two bile acids which possess the property of solubilizing

The morphological and cultural chalracteristics of Absidia

plate to those of ursodeoxycholic acid and chenodeoxycholic

sp. BA16 was studied in comparison with two reference strains,

Absidia corymbifera IFO 4009 and Absidia butleri (Gongronella

Screening for Microorganisms Capable

Associate Professor Dr. Naline Nilubol

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to Transform Lithocholic Acid into

Useful Bile Acids.

Miss Oytip Kanjanapanjapol

Miss Sunanta Cajesanun

Project Title:

Name of the Investigator:

cholesterol gallstone.

butleri) IFO 8080. Though some characteristics of the strain BA16 were resembled to the reference strains, other different characteristics were also observed. Therefore, Absidia sp. BA16 might be a new species of the genus "Absidia".

The derivative of lithocholic acid produced by strain BA16

The derivative of lithocholic acid produced by strain BA16 was extracted from the culture broth with ethyl acetate and chomatographed on silica gel column. Then it was crystallized by

identified as $3 \propto$, 15β -dihydroxy-5 β - cholanic acid on the basis of elemental analysis, IR, C^{13} NMR, H'-NMR.

The optimal conditions for the product formation in the

adding the mixture of ethyl acetate and hexane. The product was

fermentor were the production medium containing 40 g/l casava starch and 5 g/l sodium nitrate as carbon and nitrogen sources, cultivation temperature at 30°c, initial pH at .6.5 without pH

control during cultivation, agitation speed at 300 rpm/min., aeration rate at 1.2 vvm, feeding lithocholic acid in 1% dioxane after 56 hour of cultivation followed by the addition of lithocholic at 72 and 84 hour. The maximal yield obtained under these

conditions was 2.87 g/l at 90 hour of cultivation.