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5mmminipage15cm A Morphologically Structured Model for Mycelial Growth and Secondary Metabolite Formation

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摘要 A morphologically structured model is proposed to describe the batch fermentation of lovastatin according to the growth kinetics of filamentous microorganisms. Three kinds of hyphae are considered in the model: actively growing hyphae, non-growing hyphae and deactivated hyphae. Furthermore, actively growing hyphae consist of three morphological compartments: apical compartment which gives rise to hyphal tip extension; subapical compartment which is related to hyphal branching; and hyphal compartment which is only responsible for secondary metabolite formation. The kinetics of mycelial growth mechanism is summarized and applied in modeling lovastatin fermentation. A Michaelis-Menten kinetic model with substrate inhibition is proposed for product formation. As expected, the model simulations fit well with experimental data obtained either from a laboratory scale 10 L fermenter or from a pilot-plant scale fermenter.

关键词 [lovastatin](#) [Aspergillus terreus](#) [filamentous microorganism](#) [morphologically structured model](#) [kinetics](#)

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Key words [lovastatin](#); [Aspergillus terreus](#); [filamentous microorganism](#); [morphologically structured model](#); [kinetics](#)

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