

Open access • Journal Article • DOI:10.1007/S00449-015-1432-5

Enhancing enzyme stability and metabolic functional ability of β -galactosidase through functionalized polymer nanofiber immobilization — Source link \square

Mailin Misson, Mailin Misson, Bo Jin, Binghui Chen ...+1 more authors

Institutions: University of Adelaide, Universiti Malaysia Sabah, Xiamen University

Published on: 24 Jun 2015 - Bioprocess and Biosystems Engineering (Springer Berlin Heidelberg)

Topics: Enzyme binding and Functional ability

Related papers:

- A rapid and sensitive method for the quantitation of microgram quantities of protein utilizing the principle of protein-dye binding
- Manipulation of nanofiber-based β-galactosidase nanoenvironment for enhancement of galacto-oligosaccharide production.
- Galacto-oligosaccharides Synthesis from Lactose and Whey by β -Galactosidase Immobilized in PVA
- Interfacial biocatalytic performance of nanofiber-supported β-galactosidase for production of galacto-oligosaccharides
- Development of a Hybrid Bioinorganic Nanobiocatalyst: Remarkable Impact of the Immobilization Conditions on Activity and Stability of β-Galactosidase



Enhancing enzyme stability and metabolic functional ability of βgalactosidase through functionalized polymer nanofiber immobilization

Abstract

A functionalized polystyrene nanofiber (PSNF) immobilized β -galactosidase assembly (PSNF-Gal) was synthesized as a nanobiocatalyst aiming to enhance the biocatalyst stability and functional ability. The PSNF fabricated by electrospinning was functionalized through a chemical oxidation method for enzyme binding. The bioengineering performance of the enzyme carriers was further evaluated for bioconversion of lactose to galacto-oligosaccharides (GOS). The modified PSNF-Gal demonstrated distinguished performances to preserve the same activity as the free β -galactosidase at the optimum pH of 7.0, and to enhance the enzyme stability of PSNF-Gal in an alkaline condition up to pH 10. The PSNF assembly demonstrated improved thermal stability from 37 to 60 °C. The nanobiocatalyst was able to retain 30 % of its initial activity after ninth operation cycles comparing to four cycles with the unmodified counterpart. In contrast with free β -galactosidase, the modified PSNF-Gal enhanced the GOS yield from 14 to 28 %. These findings show the chemically modified PSNF-based nanobiocatalyst may be pertinent for various enzyme-catalysed bioprocessing applications.