Author's Accepted Manuscript

Purification and biochemical characterization of peroxidase isoenzymes from *Ficus carica* latex

Alshaimaa M. Elsayed, Usama M. Hegazy, Marwa G.A. Hegazy, Somia S. Abdel-Ghany, Walaa H. Salama, Ahmed M.H. Salem, Afaf S. Fahmy



 PII:
 S1878-8181(18)30381-5

 DOI:
 https://doi.org/10.1016/j.bcab.2018.07.009

 Reference:
 BCAB807

To appear in: Biocatalysis and Agricultural Biotechnology

Received date: 30 May 2018 Revised date: 2 July 2018 Accepted date: 4 July 2018

Cite this article as: Alshaimaa M. Elsayed, Usama M. Hegazy, Marwa G.A. Hegazy, Somia S. Abdel-Ghany, Walaa H. Salama, Ahmed M.H. Salem and Afaf S. Fahmy, Purification and biochemical characterization of peroxidase isoenzymes from *Ficus carica* latex, *Biocatalysis and Agricultural Biotechnology*, https://doi.org/10.1016/j.bcab.2018.07.009

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting galley proof before it is published in its final citable form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

Purification and biochemical characterization of peroxidase isoenzymes from Ficus

carica latex

Alshaimaa M. Elsayed^{1*}, Usama M. Hegazy¹, Marwa G. A. Hegazy², Somia S. Abdel-Ghany¹, Walaa H. Salama¹, Ahmed M. H. Salem², Afaf S. Fahmy¹

¹Molecular Biology Department, Genetic Engineering and Biotechnology Division, National Research Centre, Dokki, 12622 Giza, Egypt,

²Biochemistry Department, Faculty of Science, Ain Shams University, P.O. box 11381, Abbassia, Cairo, Egypt.

*Corresponding Author: shaimaa.1182@yahoo.com

Abstract

Three peroxidase isoenzymes were isolated from *Ficus carica* latex using CM-Sepharose, DEAE-Sepharose and Sephacryl S-200. The complete purification was carried out for FP1 only due to the low level of activity and protein concentration of FP2 and FP3. The purified isoenzyme FP1 was found to be monomeric with a molecular weight of 30 kDa. FP1 and FP3 isoenzymes had the same pH and temperature optima at pH 5.5 and 40°C, whereas the optimum values of pH and temperature were at pH 7.0 and 30 °C for FP2. On the other hand, FP1, FP2 and FP3 were stable at 50°C, 40°C and 30 °C respectively, whereas FP3 had low thermostability. FP1 isoenzyme was found to be stable between pH 5.0 and 7.5, and FP2 was stable from pH 4.0 to 8.0, while FP3 was found to be stable in acidic range between pH 4.5 and pH 5.5. The activity of both FP1 and FP2 peroxidase isoenzymes was increased by the high concentration of Ca²⁺ (10mM). The three peroxidase isoenzymes have a broad specificity towards some phenolic substrates and *O* - Phenylenediamine showed higher affinity towards the three peroxidase isoenzymes,

Download English Version:

https://daneshyari.com/en/article/8405571

Download Persian Version:

https://daneshyari.com/article/8405571

Daneshyari.com