



Fungal biodegradation of pomegranate ellagitannins

<u>J. A. Ascacio-Valdés¹</u>, J. J. Buesrostro¹, L. Sepúlveda¹, R. De la Cruz¹, A. F. Aguilera-Carbó², A. Prado³, J. C. Contreras¹, R. Rodríguez-Herrera¹ & C. N. Aguilar¹.

¹Department of Food Science and Technology. School of Chemistry. Universidad Autónoma de Coahuila. Blvd. Venustiano Carranza, col. República. Saltillo, 25280, Coahuila, México. E-mail: alberto_ascaciovaldes@uadec.edu.mx.

²Department of Food Science and Technology. Universidad Autónoma Agraria "Antonio Narro". Buenavista, Saltillo, 25000, Coahuila, México.

³Department of Biotechnology, Universidad Autónoma Metropolitana Unidad Iztapalapa. 09340, México, D. F.

Key words: Aspergillus niger GH1, solid state culture, ellagitannins

Introduction. At this time, there is scarce about information ellagitannins biodegradation. Several authors mention that it is necessary to carry out further studies to elucidate the mechanism of ellagitannins biodegradation (Scalbert, 1991; Vivas et al 2004). There are some studies that describe the ellagitannins biodegradation by fungal enzymes, such as, tannase enzyme (Yoshida et al 1999), β-glucosidase (Vattem & Shetty, 2002, 2003), polyphenoloxidase (Shi et al 2005). In 2009 Aguilera-Carbó et al suggested the existence of an enzyme responsible for the ellagitannins degradation, produced by Aspergillus niger GH1, by solid state culture. The authors established the existence of the enzyme by electrophoresis analysis (SDS-PAGE), which had a molecular weight around 200 kDa. It is suspected that this enzyme, recently reported, is responsible for the ellagitannins degradation, however, it is necessary to generate new information in order to understand the ellagitannins degradation mechanism, therefore, the aim of this study was to associate an enzyme produced by Aspergillus niger GH1, by solid culture. with the ellagitannins state degradation and to identify the intermediate compounds of this degradation.

Methods. A solid state culture was carried out using Pontecorvo media culture (Aguilera-Carbó et al 2009), Aspergillus niger GH1, pomegranate ellagitannins and PUF as support at 0 h to 36 h in 250 mL reactors. The enzymatic extracts were recovered with citrate buffer (pH 5.0, 50 mM) and centrifuged in Nanosep® tubes (1.5 mL) for the partial purification. After, the ellagic acid accumulation was measured by HPLC, the ellagitannase enzymatic activity was determined (Mireles-Ramírez et al 2008), and the intermediate compounds was identified by LC/MS.

Results. The obtained results are showed in the following figures.



Fig.1. Enzymatic activities evaluated.



Fig.2. Enzymatic activity (blue), ellagic acid accumulation (red), punicalagina (A), punicalina (B), gallagic acid (C) and ellagic acid (D).

Conclusions. It was demonstrated that *Aspergillus niger* GH1, under solid state culture using ellagitannis as substrate, was able to produce an enzyme related with the pomegranate ellagitannins degradation.

Acknowledgements. This work was funded by the project: SEP-CONACYT-(51360)-2005-24348.

References.

Vattem, D., Shetty, K. (2002). Solid-state production of phenolic antioxidants from cranberry pomace by *Rhizopus oligosporum. Food Biotechnol.* (16) 189-210. Vivas, N., Laguerre, M., Pianet de Biossel, I., Vivas de Gaulejac, N., Nonier, M. (2004). Conformational interpretation of vescalagin and castalagin physicochemical properties. *J. Agric. Food Chem.* (52) 2073-2078.