## PARTIAL PURIFICATION OF PECTINASES PRODUCED BY THE FILAMENTOUS FUNGUS Aspergillus niveus

Maller, A<sup>1</sup>.; Silva, T.M<sup>2</sup>.; Damásio, A.R.L<sup>1</sup>; Jorge, J.A<sup>2</sup>., Terenzi, H.F<sup>2</sup>. & Polizeli, M.L.T.M<sup>2</sup>

<sup>1</sup>Departamento Bioquímica e Imunologia, FMRP/USP, SP <sup>2</sup>Departamento Biologia, FFCLRP/USP, SP e-mail:alemaller@yahoo.com.br

Pectinases are important industrial enzymes which may be classified in esterases and depolymerases. These enzymes catalyze the degradation of pectic substances that occur in higher plants, and are constituted by a main chain of polygalacturonic acid branched with other sugars. Pectinases are mainly produced by several Aspergillus sp, however nothing is known about this enzyme from A. niveus. The aim of this work was to standardize cultivation conditions and physicochemical parameters for pectinase production and the partial purification of a crude filtrate by ion exchange chromatography. The assays were carried out with 1% polygalacturonic acid in 100 mM sodium acetate buffer, pH 6.0. The reducing sugar formed was quantified with 3',5' dinitrosalicylic acid. The optimized conditions were: Czapeck medium added of 1% pectin (Sigma), at 30°C, for 9 days under stationary conditions, or 2 days under agitation. Citric fruits peels were also good inducers of polygalacturonases and also low levels of pectin and pectate lyases. Regarding polygalacturonase, it was observed a maximum activity at pH 4.0 and 55°C. This enzyme was thermal stable for 90 min at 60°C. The enzyme was activated by 1mM Mn<sup>++</sup> (17%) and EDTA (10%). The polygalacturonase was partially purified by 80% ammonium sulfate precipitation, elution in DEAE cellulose, followed by Biogel P100. Support: FAPESP, CNPg