

PARTIAL PURIFICATION AND BIOCHEMICAL CHARACTERIZATION OF AN
EXTRACELLULAR XYLANASE OF *FUSARIUM HETEROSPORUM* NEES

Henn, C².; Soster, C. C².; Costa, A. M¹.; Simão, R.C.G².; Osaku, C. A².; Peralta,
R, M¹.; Kadowaki, M. K².

¹Depto. Bioquímica – UEM/Maringá-PR/Brazil.

²Centro de Ciências Médicas e Farmacêuticas – UNIOESTE/Cascavel-PR/Brazil

E-mail: marinakk@unioeste.br

Xylanases are produced by a large variety of microorganisms. Fungal systems are excellent xylanase producers, but often co-secrete cellulases which can adversely affect pulp quality. Based on this, *Fusarium heterosporum* Nees presented cellulase-free xylanase activity induced by corn residues, mainly for corn straw and corn cob (21.15 and 13.26 U/mg protein), respectively. This inducible xylanase was partially purified from the extract of *F. heterosporum* Nees and biochemical characterization was performed. The enzyme was purified from an eight day old culture filtrated in submerged fermentation, by ethanol precipitation followed by gel filtration Sephadex G-100 and ion exchange chromatography CMCelulose columns. The partially purified sample showed maximum activity at pH 6.0, whereas the optimum temperature for enzyme activity was 60°C and thermal stability up to 55°C. The xylan hydrolysis activity was inhibited with 5mM Hg⁺², Cu⁺², Al⁺³, (48%, 17% and 75%), respectively. β-mercaptoethanol enhanced the xylanase activity in 24%, Ca⁺² and Mn⁺² activated 19% and 38%, respectively. The xylanase hydrolyzed beechwood xylan and birchwood xylan yielded mainly xylose as end products, suggesting it is exoxylanase. Thus, this microorganism can represent a useful tool for saccharify corn xylan, since commercially available enzymes are unable to degrade it.

Supported by: Unioeste/Cascavel-PR/Brazil