| Title | Purification and some properties of alpha-amylase from post-harvest Pachyrhizus erosus L. tuber |
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Abstract

 α -Amylase, a starch splitting enzyme, was purified to homogeneity from post-harvest *Pachyrhizus erosus* L. tuber by successive chromatography on DEAE- and CM-cellulose columns. Purification achieved was 110 fold from the crude extract with a yield of 22.8%. SDS-PAGE showed a molecular weight of 40 kDa for the enzyme. The enzyme is of α -type as it lost total activity in the presence of EDTA, a chelating agent. It is a glycoprotein that contains 2.6% sugar as estimated by the phenol-sulfuric acid method. The enzyme displayed optimum activity at pH 7.3 and 37 °C with an apparent $K_{\rm m}$ value of 0.29% for starch as substrate. The enzyme was strongly inhibited by Cu²⁺, Fe²⁺ and Zn²⁺, moderately by Li²⁺, Hg⁺ and Cd²⁺ and slightly by Ag⁺, Mg²⁺ and K⁺. Calcium ion almost doubled the activity whereas Fe³⁺, Mn²⁺ and Na⁺ enhanced it appreciably.