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Partial purification and some properties of a-glucosidase from *Trichoderma longibrachiatum*

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ABSTRACT: The use of hydrolase enzyme plays an important role in the industrial production of a-D-glucose from carbohydrate sources. This study investigated partial purification and characterization of a-glucosidase from *Trichoderma longibraciliatum* with a view to enhancing its potentials in biotechnological processes. Strains of *Trichoderma longibraciliatum* were cultured on rice bran medium at 30°C for 96 hour for the production of a-glucosidase. The enzyme was partially purified by eluting the ammonium sulphate (70%) saturation precipitated sample on Sephadex G-75 and Sephadex G-25. Enzyme assay was carried out using p-nitrophenyl-a-D-glucopyranoside (PNP- a-G) as the substrate and protein concentration was determined. Kinetica parameters, molecular weight, pH effect, temperature and thermostability were also determined. The activity of enzyme in the presence of arylglucosides: and different cations were monitored. The partially purified protein, migrated as a single band in 10% SDS-Polyacrylamide gel-electrophoresis. The enzyme presented a relative molecular weight of about 58KDa as estimated by PAGE. The extracellular a-glucosidase showed typical a-glucosidase activity, hydrolyzing p-nitrophenyl-a-D-glucopyranoside and exhibited optimum catalytic activity (4.89pmol/ml/min), at 40°C and pH 4.5. The enzyme was stable at 40°C for 150 minutes. Carboxymethylcellulose was also hydrolyzed by this enzyme. The K_m and V_{max} with p-nitrophenyl- *a* -D-glucopyranoside were 33.33mM and 20.00 pmol/min/mg protein, respectively. This study therefore revealed the presence of a-glucosidase enzyme.

KEYvVORDS: Enzyme purification, characterization, a-glucosidase, Trichoderma longibrachiatum.