ABSTRACT

This present study deals with the Biotechnological applications of commercially important α-Amylase produced from *Penicillium digitatum* using surface culture fermentation. To enhance the output, the One Factor at a Time (OFAT) approach was used in the production of α -amylase by altering fermentation parameters such as incubation temperature, incubation period, pH, and Carbon and Nitrogen sources. To increase enzyme yield, different concentrations of sucrose, yeast extract, potato peel, KH2PO4, MgSO4, MnSO4, FeSO4, ZnSO4, and NaCl were tested. Optimized production was reached with 2.5 g Sucrose, 5 g Yeast Extract, 3.5 g Potato Peel, 0.25 g MgSO₄,0.75 g KH₂PO₄, 1 g MnSO₄, 0.75 g FeSO₄, 0.5 g ZnSO₄, and 1.25 g NaCl. The enzyme yield was 2.958 ± 0.1 U/mL for the extracellular extract and 2.147 ± 0.1 U/mL for the intracellular extract. Fermentation was carried out at pH 5.5 and 27°C for 5 days. The enzyme was partially purified by adding sodium sulphate to the crude enzyme generated from solid-state fermentation, which was then stirred at room temperature. The enzyme's stability was examined at a variety of temperatures, pH levels, and substrate concentrations. α-Amylase remains stable at 37°C and perform best at pH 5.5. Penicillium digitatum was treated to UV irradiation and ethyl methane sulfonate (EMS) mutagenesis to increase α-amylase synthesis. This resulted in the creation of mutant strains with greater enzymatic activity. Penicillium digitatum has been shown to be a sustainable and cost-effective source of α -amylase for industrial applications such as food processing, bioremediation, and bio-fertilizer production. This research reveals that α-amylase from P. digitatum can be effectively used in numerous biotechnological applications.