

Biochemical characteristics of chitinase enzyme from *Bacillus* sp. of Kamojang Crater, Indonesia.

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Abstract

Chitinase and chitin deacetylase are enzymes capable of degrading chitin into chitooligomers and chitosan. The chitinases characterized and purified in this study were extracted from the acidophilic *Bacillus* sp. isolated from Kamojang Crater West Java Indonesia. When grown in liquid media containing colloidal chitin, the optimum chitinase activity of the acidophilic isolate was reached after 4-5 days of incubation. The optimum temperature and pH of the chitinase and chitin deacetylase were found at 37 degrees C and pH 5. When incubated at pH 5, the activity of chitin deacetylase was increased; after 3 h, the activity was 1.5 times of the control. The enzyme was stable at pH 4, after 2 h incubation, the activity was still 80% of the control. The chitinase and chitin deacetylase activities were not influenced by Mg(++) nor Ca(++), Ni(++), and Cu(++). Ni(++) inhibited the chitinase activity, while chitin deacetylase activity was not affected by Cu(++) addition. When 1 mM of EDTA was added, the enzyme activity was reduced 40 to 50%.