

Evaluation of Proteins and Hydrolytic Enzymes from the Seeds of Genus *Caesalpinia*

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The study investigates the protein content in total and activity of esterase in soaked dehulled seeds of three species of *Caesalpinia*. Protein content varied between 11% and 15%, with *Caesalpinia mimosaoides* the highest at 15.04%. All species demonstrated esterolytic activity, with *C. mimosaoides* showing the highest ester hydrolyzing activity and specific activity. Conversely, *C. decapetala* displayed the lowest esterase activity and specific activity. These findings highlight species-specific variations in protein content and enzymatic activity within the *Caesalpinia* genus. Over the course of germination, protein content decreased from 17% at 72 hours to 9% at 120 hours. Esterase activity was detected in both soaked seeds and seedlings at all phases of germination. The highest esterase activity was observed on the 3rd day of germination. Polyacrylamide gel electrophoresis was conducted to study the banding patterns of esterases in the presence of an artificial substrate. The 1-Naphthyl acetate substrate revealed four esterolytic bands in *C. mimosaoides*. These findings provide insights into the dynamics of protein degradation and esterolytic enzyme activity during seed germination.

Keywords: *Caesalpinia* Seeds; Esterases; Germination; Proteins; Variation.

Esterases are a family of enzymes that accelerate the hydrolysis reaction of various types of endogenous and exogenous esters. These include phosphoric monoester hydrolases, phosphoric diester hydrolases, thioester hydrolases, sulfuric ester hydrolases, and carboxylic ester hydrolases.¹

They are found abundantly in nature, exist in various molecular forms, and display a wide range of substrate specificity. The physiological roles of these esterases are multifaceted. Esterases help in the maturation of fruits, reproductive process, abscission, cell expansion, and the hydrolysis of xenobiotic

molecules containing ester. These are widespread in the nature, occurring in microorganisms, plants, and animals. They are present in various tissues, such as the brain, muscle, small intestine, heart, liver, lung, kidney, adipose tissue, testes, nasal and respiratory tissues, blood and leukocytes.² However, the expression and activity are dependent on both the tissue and the organism.³ Esterases are crucial in the metabolism and subsequent detoxification of various agrochemicals and pharmaceuticals.^{4,5} Pyrethroids undergo hydrolysis through Carboxylesterases⁶⁻⁸ and bind to organophosphates in a stoichiometric manner.⁹⁻¹⁰ Carboxylesterases are crucial in the

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