



## Study of 2-mercaptobenzoic acid and 2-pyridinethiol as inhibitors on the cresolase and catecholase reactions of mushroom tyrosinase

Nematollah Gheibi<sup>1</sup>, Mehran Sheikhzadegan<sup>2</sup>

- 1) Cellular and Molecular Research Center, Qazvin University of Medical Sciences, Qazvin, Iran
- 2) Department of Biophysics, Islamic Azad University, Science and Research Branch, Tehran, Iran

### ABSTRACT

Catecholase and cresolase activities of mushroom tyrosinase (MT) were studied in the presence of 2-mercaptobenzoic acid (thiosalicylic acid) and 2-pyridinethiol inhibitions. Caffeic acid and p-coumaric acid were used as natural substrate for the enzyme for the catecholase and cresolase reactions, respectively. The catecholase and cresolase activities of MT in the presence of 2-pyridinethiol as inhibitor achieved in the concentrations of (1.5, 3, 4.5 and 6  $\mu\text{M}$ ). In addition, the cresolase and catecholase activities of MT in the presence of 2-mercaptobenzoic acid used in presence of (2.5, 5, 10 and 15  $\mu\text{M}$ ). The inhibition constant ( $K_i$ ) values of 2-pyridinethiol obtained 0.84 and 5.37  $\mu\text{M}$  for catecholase and cresolase reactions, respectively, with noncompetitive pattern. But for the 2-mercaptobenzoic acid revealed a competitive mode of inhibition with the inhibition constants of 5.45 and 9.35  $\mu\text{M}$ , for catecholase and cresolase reactions, respectively. Thus, the results showed that the carboxyl and sulfhydryl functional group of these organosulfur compounds play a crucial role in the inhibition of MT. Their  $K_i$  values showed that they are among the good inhibitors of enzyme.

**Key words:** Mushroom tyrosinase; 2-mercaptobenzoic acid; 2-Pyridinethiol; Inhibition; Sulfur