

## عنوان مقاله:

Kinetic Study of Erythrose Reductase Extracted from *Yarrowia lipolytica*

## محل انتشار:

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تعداد صفحات اصل مقاله: 5

## نویسندگان:

Masoud Mohammadi Farsani - *Department of Biology, Faculty of Science, Shahid Chamran University of Ahvaz, Ahvaz, Iran*

Mohammadreza Mohammadi - *Department of Biology, Faculty of Science, Shahid Chamran University of Ahvaz, Ahvaz, Iran*

Gholam Reza Ghezelbash - *Department of Biology, Faculty of Science, Shahid Chamran University of Ahvaz, Ahvaz, Iran*

Ali Shahriari - *Department of Basic Sciences, Faculty of Veterinary Medicine, Shahid Chamran University of Ahvaz, Ahvaz, Iran*

## خلاصه مقاله:

Erythritol as a non-caloric and non-cariogenic sweetener is safe for diabetics. Both microbial fermentation and chemical methods can be used to produce erythritol, but chemical methods failed to be industrialized due to their low efficiency. *Moniliella tomentosa*, *Aureobasidium* sp. and *Yarrowia lipolytica* are industrial producers of erythritol. Erythrose reductase (ER) is a key enzyme in the biosynthesis of erythritol and catalyzes the final step in this pathway. Enzyme extract was obtained from *Y. lipolytica* by grinding cells with 0.5mm glass beads and ER activity was performed using 10  $\mu$ l enzyme extract, 7.5 mM NADPH and 12 mM D-erythrose in potassium phosphate buffer (pH 7.5). Reaction was monitored with decreasing of NADPH absorbance in OD340 at 37 °C for 8min by a microplate analyzer. In order to determine the activation energy ( $E_a$ ), activity of enzyme was measured in 4-45 °C and results were analyzed with Kinetic software according to Arrhenius equation. The best enzyme activity of ER was 6.268 mU. One unit of ER activity was defined as the amount of enzyme that catalyzes the oxidation of 1  $\mu$ mol of NADPH per minute. Specific activity of enzyme was equal to 3.24U/mg and finally the  $E_a$  was determined to be 29.6208 KJ. ER specific activity in this study was lower than the only similar study that used *Y. lipolytica*. Purification, overexpression and optimizing the reaction can help to increase enzyme performance.

## کلمات کلیدی:

Erythrose reductase, *Yarrowia lipolytica*, Enzyme kinetics

## لینک ثابت مقاله در پایگاه سیویلیکا:

<https://civilica.com/doc/1004888>



