

Abstract



Optimization of fermentation conditions for extracellular production of a therapeutic protein drug, uricase, by Aspergillus welwitschiae using response surface methodology

Noura El-Ahmady El-Naggar

City of Scientific Research and Technological Application, Egypt

Abstract:

Uricase is commonly used in clinical analysis for determination of urate in blood and other biological fluids. Microbial uricases as protein drug have been found effective in the treatment of hyperuricemia and gout, as well as in prophylaxis and treatment of tumor lysis hyperuricemia. A potential culture, Aspergillus sp. strain 1-4 displayed high extracellular uricase activity. This uricolytic fungal isolate was identified as Aspergillus welwitschiae strain 14 on the basis of phenotypic characteristics, together with ITS region sequence analysis. Sequencing product was deposited in the GenBank database under accession number MG323529. The Plackett-Burman experimental design with 20 runs was applied for screening of fifteen variables for their significances on uricase production by Aspergillus welwitschiae. Incubation time was the most significant variable affecting uricase production followed by yeast extract and inoculum size with significant P-values of 0.0002, 0.0083 and 0.0118; respectively. These variables were chosen for optimization studies using central composite design. It was found that, the maximum uricase production (59.01U/mL) by Aspergillus welwitschiae is achieved at the following fermentation conditions g/L: sucrose 30, uric acid 3, peptone 2, yeast extract 2, NaNO3 2, K2HPO4 1, MgSO4.7H2O 0.2, NaCl 0.2 and FeSO4.7H2O 0.03", incubation time 7 days, temperature 35, pH 6, inoculum size 4 mL/50 mL medium, inoculum age 72 h and medium volume 50 mL/250 mL conical flask. An overall 2.5 fold increase in uricase production by Aspergillus welwitschiae was achieved in the medium after statistical optimization as compared with the unoptimized basal medium (23.58U/mL) before applying Plackett-Burman.

Keywords: Uricase production, Aspergillus welwitschiae, identification, scanning electron microscope, ITS region sequence, optimization, Plackett-Burman design, central composite design



Biography:

Noura El-Ahmady El-Naggar is a microbiologist. She completed her Ph.D in microbiology from Botany Department, Faculty of Science, Mansoura University, Egypt, 1998. She is presently employed as professor in Bioprocess Development Department, Genetic Engineering and Biotechnology Research Institute, City for Scientific Research and Technology Application, Alexandria, Egypt. She has published more than 50 papers in reputed journals and has been serving as an editorial board member of the "Journal of Microbial & Biochemical Technology" and the Journal "Fermentation Technology". She is a reviewer of several journals such as "Applied Biochemistry and Biotechnology, World Journal of Microbiology and Biotechnology.

Recent Publications:

- 1. Hatijah MS, Ruhayu WRW. Preliminary study on factors that enhanced the production of uricase by Aspergillus Flavus, Int J Biosci Biochem Bioinformatics 2013; 3(5).
- 2. Abbas AA. Extraction, optimization of uricase from Aspergillus niger. Int J Adv Res. 2016; 4(3), 1865-1872.

3rd Webinar on Plant Science | December 14, 2020 | Paris, France

Citation: Noura El-Ahmady El-Naggar; Optimization of fermentation conditions for extracellular production of a therapeutic protein drug, uricase, by Aspergillus welwitschiae using response surface methodology; Plant Science Webinar; December 14, 2020; Paris, France pg-70