

The 7th Conference on Bioinformatics, 3-5 January 2018





Designing specific primer to replication partial sequence of the *PgaA* gene responsible for the synthesis of endo-polygalacturonase enzymes in *Aspergillus niger*

Heydari Nezhad AM, PhD Student, University of Zabol, Zabol, Iran* Panjekeh N, Associate Professor, University of Zabol, Zabol, Iran Sabbagh SK, Associate Professor, University of Yazd, Yazd, Iran *amir.masoud_90@yahoo.com

Abstract: Pectin components are polysaccharides that are found in the middle lamella, plant cell wall and most groups of algae. Pectinases are one of the most important enzymes widely distributed in bacteria and fungi. its three major groups include polygalacturonase (PG), pectin esters (PE) and pectin lysates (PL) [1]. In the Aspergillus niger the Pga gene cluster is responsible for the synthesis of glycoside hydrolase family 28 which secrete polygalacturonase (EC 3.2.1.15) enzymes [2]. Designing and selecting efficient primers with high specificity is the first and most important step in genomic studies and qualitative quantitative assay of the transcription of protein-coding genes [3] In this study sequences related to the PgaA gene encoding end-polygalacturonase enzymes were obtained by using the National Center for Biotechnology Information database, from the species of Aspergillus niger. The Align of sequences and the selection of conserved sequences and eventually the appropriate primer design were performed using the MEGA6 [4], Oligo7 [5], BioEdit bioinformatics software and the web-based 'Primer application. Foeward and revese primers with **PegAF** 'TGCCAAGCCTTTGTTCTG 3' and PgaAR 5 'TCCATCCCACTCCTCGTAC 3' sequence were designed. Their correct performance confirmed using polymerase chain reaction and electrophoresed product analysis on agarose gel.

Keywords: Aspergillus niger; Primer design; PgaA gene; Polygalacturonase; Sequence

References

- [1] Semenova, MV. Grishutin, SG. Gusakov, AV. Okunev, ON and Sinitsyn, AP. Isolation and properties of pectinases from the fungus *Aspergillus japonicus*. Biochemistry (Moscow), 68 (2003) 559-569.
- [2] Cheng, Z. Chen, D. Wang, Q. Xian, L. Lu, B. Wei, Y and Huang. Identification of an acidic endopolygalacturonase from Penicillium oxalicum CZ1028 and its broad use in major tropical and subtropical fruit juices production. Journal of Bioscience and Bioengineering, 123 (2017) 665-672.
- [3] Adamu, O. J and Slater, A Primer Design from Conserved Sequences of the Artemisia Annua Amorpha-4, 11-Diene Synthase and Artemisia Annua Amorpha-4, 11-Diene Mono Oxygenase Synthase Gene. International Journal of Innovative Research and Advanced Studies. 4 (2017) 127-131.
- [4] Tamura, K. Stecher, G. Peterson, D. Filipski A and Kumar S. MEGA6: molecular evolutionary genetics analysis version 6.0. Molecular biology and evolution, 30: (2013) 2725-2729.
- [5] Rychlik W. OLIGO 7 primer analysis software. PCR Primer Design. (2007) 35-59.