

Accepted Manuscript

Title: Neurokinin-1 receptor (NK1R) inhibition sensitizes APL cells to anti-tumor effect of arsenic trioxide via restriction of NF- κ B axis: Shedding new light on resistance to Aprepitant

Authors: Davood Bashash, Ava Safaroghli-Azar, Samaneh Bayati, Elham Razani, Atieh Pournbagheri-Sigaroodi, Arshia Gharehbaghian, Majid Momeny, Maryam Sanjadi, Mostafa Rezaie-Tavirani, Seyed H. Ghaffari



PII: S1357-2725(18)30180-8
DOI: <https://doi.org/10.1016/j.biocel.2018.08.010>
Reference: BC 5403

To appear in: *The International Journal of Biochemistry & Cell Biology*

Received date: 19-5-2018
Revised date: 16-7-2018
Accepted date: 17-8-2018

Please cite this article as: Bashash D, Safaroghli-Azar A, Bayati S, Razani E, Pournbagheri-Sigaroodi A, Gharehbaghian A, Momeny M, Sanjadi M, Rezaie-Tavirani M, Ghaffari SH, Neurokinin-1 receptor (NK1R) inhibition sensitizes APL cells to anti-tumor effect of arsenic trioxide via restriction of NF- κ B axis: Shedding new light on resistance to Aprepitant, *International Journal of Biochemistry and Cell Biology* (2018), <https://doi.org/10.1016/j.biocel.2018.08.010>

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

Neurokinin-1 receptor (NK1R) inhibition sensitizes APL cells to anti-tumor effect of arsenic trioxide via restriction of NF- κ B axis: Shedding new light on resistance to Aprepitant

Davood Bashash^{1*}, Ava Safaroghli-Azar², Samaneh Bayati³, Elham Razani², Atieh Pourbagheri-Sigaroodi⁴, Arshia Gharehbaghian⁵, Majid Momeny⁶, Maryam Sanjadi⁷, Mostafa Rezaie-Tavirani⁸, Seyed H. Ghaffari³

¹*Department of Hematology and Blood Banking, School of Allied Medical Sciences, Shahid Beheshti University of Medical Sciences, Tehran, Iran*

²*Student Research Committee, Department of Hematology and Blood Banking, School of Allied Medical Sciences, Shahid Beheshti University of Medical Sciences, Tehran, Iran*

³*Hematology, Oncology and Stem Cell Transplantation Research Center, Tehran University of Medical Sciences, Tehran, Iran*

⁴*Department of Biotechnology, faculty of Advanced Sciences and Technology, Pharmaceutical sciences branch, Islamic Azad University (IAUPS), Tehran, Iran*

⁵*School of Biology University of Tehran, Tehran, Iran*

⁶*Cancer Cell Signaling, Turku Center for Biotechnology, University of Turku and Åbo Akademi University, Turku, Finland*

⁷*Falavarjan branch Islamic Azad University, Isfahan, Iran*

⁸*Proteomics Research Center, Faculty of Paramedical Sciences, Shahid Beheshti University of Medical Sciences, Tehran, Iran*

*Corresponding author: Davood Bashash, Ph.D

Department of Hematology and Blood banking, School of Allied Medical Sciences, Shahid Beheshti University of Medical Sciences, Tehran, Iran

Tel: +98-21-22717504

Fax: +98-21-22721150

E-mail: d.bashash@sbmu.ac.ir

Abstract

While a batch of efforts are fastened on synthesizing the novel targeted anti-cancer agents, recent investigations have achieved a breakthrough in identifying a favorable anti-tumor activity for some supportive drugs, which their safety have been confirmed thus far. The results of the present study highlighted the efficacy of Aprepitant, an oral antagonist of the neurokinin-1 receptor (NK1R), against both APL (NB4) and pre-B ALL (Nalm-6) cell lines; however, a differential sensitivity pattern was found in these cells. To the best of our knowledge, this is the first time that the molecular mechanisms of resistance to Aprepitant have been investigated and, herein, we proposed that the

Download English Version:

<https://daneshyari.com/en/article/9954158>

Download Persian Version:

<https://daneshyari.com/article/9954158>

[Daneshyari.com](https://daneshyari.com)