

عنوان مقاله:

The Effect of β -Adrenergic Agonist Isoproterenol on miR-186-3p and miR-23a Expression in Human Bone Marrow Mesenchymal Stem Cells

محل انتشار:

مجله علمی پژوهشی دانشگاه علوم پزشکی زنجان، دوره 23، شماره 96 (سال: 1393)

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خلاصه مقاله:

Background and Objective: Mobilization of Hematopoietic Stem Cells (HSCs) for transplantation and the importance of β -adrenergic signals in induction of this process have been well investigated. However, little is known about the role of β -adrenergic signals in mobilization of HSCs and factors influenced by these signals. The Chemokine Stromal Derived Factor-1 (SDF-1) which is expressed by human bone marrow-derived mesenchymal stem cells (hMSCs), has a key role in mobilization of HSCs. In addition, miR-186-3p and miR-23a can regulate the expression of SDF-1 in hMSCs. In this study, to investigate the role of miR-186-3p and miR-23a in mobilization process, expression of both miRNAs was evaluated in hMSCs treated by Isoproterenol (a β -adrenergic agonist). **Materials and Methods:** hMSCs were isolated from human bone marrow and cultured. Following flowcytometric analysis, the cells were treated with 100 M isoproterenol. Total RNA was extracted at 12 and 48 hours post treatment, and also from untreated hMSCs as control. Then, miR-186-3p and miR-23a expression levels were quantified by quantitative Reverse Transcriptase PCR. **Results:** The expression level of miR-186-3p increased significantly at 12 and 48 hours post treatment ($P < 0.05$). In addition, the expression level of miR-23a decreased at 12 hours post treatment and increased significantly at 48 hours post treatment ($P < 0.05$). **Conclusion:** Isoproterenol induces miR-186-3p in hMSCs. miR-23a is primarily decreased, and then increased due to treating with isoproterenol. So both miRNAs can contribute to mobilization process. **References** 1- Horowitz MM, Gale RP, Sondel PM, et al. Graft-versus-leukemia reactions after bone marrow transplantation. *Blood*. 1990; 75: 555-62. 2- Sykes M, Nikolic B. Treatment of severe autoimmune disease by stem-cell transplantation. *Nature*. 2005; 435: 620-7. 3- Copelan EA. Hematopoietic stem-cell transplantation. *N Engl J Med*. 2006; 354: 1813-26. 4- Wright DE, Wagers AJ, Gulati AP, Johnson FL, Weissman IL. Physiological migration of hematopoietic stem and progenitor cells. *Science*. 2001; 294: 1933-6. 5- Anderlini P, Körbling M. The use of mobilized peripheral blood stem cells from normal donors for allografting. *Stem cells*. 1997; 15: 9-17. 6- Orkin SH, Zon LI. Hematopoiesis: an evolving paradigm for stem cell biology. *Cell*. 2008; 132: 631-44. 7- Levesque J-P, Winkler IG. Mobilization of hematopoietic stem cells: state of the art. *Curr Opin Organ Transplant*. 2008; 13: 53-8. 8- Motabi IH, DiPersio JF. ... Advances in stem cell mobilization. *Blood Rev*. 2012; 26: 267-78. 9- Siena S, Bregni M, Brando B, R

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