## عنوان مقاله:

Fluorouracil-Loaded PLGA Declined Expression of Pro-Inflammatory Genes IL-9, IL-1VA, IL-1V and IFN- V in the HT-Y9 Colon Cancer Cell Line-0

## محل انتشار:

مجله گزارش های بیوشیمی و زیست شناسی مولکولی, دوره 12, شماره 4 (سال: 1402)

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#### خلاصه مقاله:

Background: Pro-inflammatory cytokines play critical roles in cancer pathobiology and have been considered potential targets for cancer management and therapy. Understanding the impact of cancer therapeutics such as  $\Delta$ -fluorouracil ( $\Delta$ -FU) on their expression might shed light on development of novel combinational therapies. This study aimed to encapsulate  $\Delta$ -FU into PLGA and evaluate their effects on the expression of pro-inflammatory genes IL- $^4$ , IL- $^4$ YF, and IFN- $^4$ Y in the HT- $^4$ P cells. Methods: PLGA- $^4$ FU NPs were constructed and characterized by Dynamic Light Scattering (DLS) and Atomic Force Microscopy (AFM). The cytotoxicity was evaluated by MTT test and, the IC $^4$ P was identified. HT- $^4$ P cells were treated with different concentrations of the PLGA- $^4$ PU NPs for  $^4$ P hours and, gene expression levels were analyzed by qRT-PCR. Results: DLS and AFM analysis revealed that the prepared PLGA- $^4$ PU NPs were negatively charged spherical-shaped particles with a mean size of  $^4$ PCR. Results: DLS and AFM analysis revealed the viability of HT- $^4$ P cells in a dose- and time-dependent manner. The qRT-PCR results revealed a dose-dependent decrease in the expression of IL- $^4$ PCPVA, IL- $^4$ PCR and IFN- $^4$ P genes, and their expressions were significantly different in both  $^4$ PCR and  $^4$ PCR treated groups compared to the control. However, although the treatment of HT- $^4$ PC cells with  $^4$ PCR resulted in decreased expression of the studied genes, the differences were not statistically significant compared to the control group. Conclusions: PLGA- $^4$ PCR into PLGA improved considerably impact of the  $^4$ PCU on the HT- $^4$ PC cells

# كلمات كليدي:

.Cancer therapy, Colorectal cancer, Fluorouracil, Polylactic Acid-Polyglycolic Acid Copolymer (PLGA), Pro-inflammatory cytokine

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