

## عنوان مقاله:

Adenosine A<sub>1</sub> Receptor Antagonist Up-regulates Casp<sup>3</sup> and Stimulates Apoptosis Rate in Breast Cancer Cell Line T47D

## محل انتشار:

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## خلاصه مقاله:

Background: Adenosine receptor family, especially A<sub>1</sub> type is-overexpressed in breast-derived tumor cells and the P<sup>53</sup> gene is mutant in some of these cells while the casps gene is of wild type as well. The aim of this study was to evaluate the effect of the A<sub>1</sub> receptor function on cell programmed death or proliferation, as well as the relationship between this receptor stimulation/inhibition and caspase 3 (casp<sup>3</sup>) expression in T47D cell line that has a mutant and non-functional P<sup>53</sup> gene. Materials and Methods: The expression of casps<sup>3</sup> was measured by real-time polymerase chain reaction and then flow cytometry and MTT assay were used to assess the apoptotic and proliferation cell rate after the treatment of T47D cells with specific agonist N<sup>6</sup>-cyclopentyladenosine (CPA) and antagonist 1,3-dipropyl-8-cyclopentylxanthine (DPCPX) of this receptor 24, 48, and 72 hours after treatment. Result: Our results indicated that DPCPX significantly induces apoptosis in T47D cells and the rate of survival cell after the reduction of this treatment, especially 72 hours after treatment. Finally, the expression of casp<sup>3</sup> was up-regulated by DPCPX treatment, especially in 72 hours while CPA treatment had opposite results ( $P > 0.05$ ). Conclusion: In general, DPCPX could up-regulate casp<sup>3</sup> gene expression and subsequently increase the apoptosis rate in T47D cells with casp<sup>3</sup> expression without the

.P53 gene interference. Therefore, adenosine A1 receptor antagonists may be introduced as anti-cancer agents

## کلمات کلیدی:

Receptor, Adenosine A1, Apoptosis, Genes, Casp3, T47D Cells

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