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Gene increases effectiveness of drugs used to fight cancer and allows reduction in dosage

by [editor](#) on November 28, 2009

in [Indian Press Release](#), [Web News Wire](#)

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This research was conducted by Ana Rosa Rama Ballesteros, from the Department of Anatomy and Human Embryology at the [University](#)  of Granada, and directed by professors Antonia Aránega Jimenez, José Carlos Prados Salazar and Consolación Melguizo Alonso. Its aim was to study the possibility of reducing the dosage of drugs currently administered to cancer patients using combination therapy with suicide gene E.

Scientists from the UGR have shown that the bacteriophage phiX174 killer gene called E, can be used to induce death in tumour cells. So far, attempts to use many chemotherapeutic (cytotoxic) agents similar to the E gene have shown severe limitations resulting from their toxicity and their poor affinity with the tumour.

Advantages of gene therapy

As Ana Rosa Rama explains, chemotherapy, radiotherapy and surgery [show](#)  at present limited results in advanced stages of cancer. That is why it is urgent to find new therapies, and gene therapy has emerged as a potentially powerful therapeutic platform. Her [work](#)  has shown that it is possible to use gene therapy as an aid to chemotherapy, improving its results when it comes to attacking cancer, thus allowing the dosage of agents to be reduced and contributing to a reduction in side effects for the patient.

In order to understand how the E gene works, the researchers conducted studies using various techniques. The results indicate that the E genes mechanism of action is to induce apoptosis (cell death), possibly through mitochondrial injury.

Therefore, they stress that this new E gene appears as an ideal candidate to be transfected into tumour cells in order to induce apoptosis, possibly through mitochondrial activation, and to increase the sensitivity of these cells to the action of the drug developed specifically to act on them.

The results of this research suggest the possibility of reducing the concentration of chemotherapeutic agents in current use with cancer patients. Thus, in lung cancer cell line A-549, scientists from the UGR achieved a 14% inhibition of tumour growth and reduced by 100 times the dose of Paclitaxel agent when it was combined with gene E. In the case of colon cancer, the results were similar. However, the most relevant fact was found in the breast cancer cell line MCF-7, in which the dose of the chemotherapeutic agent, doxorubicin, was reduced by 100 times, reaching up to a 21% greater inhibition of tumour proliferation when combined with gene E. Currently, researchers from the UGR are in the process of obtaining a patent for gene E.

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Reference:

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