

# Toxicity of DNA Intercalating Agents in the Human Body

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## INTRODUCTION:

Throughout this Poster I will be showing the research and knowledge I have gathered not only on the importance of this topic but on how Intercalating agents could possibly be used in the future as a possible very successful cancer treatment. Discussed will be the mechanism and concepts of the process that these drugs perform inside the body.

Chemical Mechanisms of Intercalation occurring

## Agents, Mechanisms and Toxicity:

- There are multiple intercalating agents under research, the three main ones being, Acridine, Actinomycin, Ethidium Bromide
- These agents bind to DNA which in turn inhibits a key replication enzyme called TOPoisomerase II
- Due to this inhibition and space between base pairing, mutations can arise in DNA replication
- Not only is replication inhibited but all DNA repair and translation becomes effected leading to cell death



## Acridine and Measurement of Binding Affinity:

- Due to the property of fluorescence Acridine has it makes it very easy to measure and study in the body
- Acridine has a very high binding affinity leading to high success rates in DNA binding
- The permanent inhibition this agent has on DNA replication makes it very useful to study intercellular complex mechanisms with.

Statistical Measurements of successful binding affinity to DNA

## Possibilities for the Future:

With cancer being uncontrollable cell replication this research and technology with extreme studies performed could lead to a breakthrough in successful cancer research. If a way to attach a signal ligand to tumor cells and avoid healthy cells was discovered, this would lead to an extreme increase in cell selectivity and binding affinity. It is impossible for treatments such as this to be successful.

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC17493749/

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