

The Chemical Biology of DNA Damage, 2010, 488 pages, Nicholas E. Geacintov, Suse Broyde, 3527630120, 9783527630127, John Wiley & Sons, 2010

Bringing the power of biochemical analysis to toxicology, this modern reference explains genotoxicity at the molecular level, showing the links between a DNA lesion and the resulting cellular or organismic response. Clearly divided into two main sections, Part 1 focuses on selected examples of important DNA lesions and their biological impact, while the second part covers current advances in assessing and predicting the genotoxic effects of chemicals, taking into account the biological responses mediated by the DNA repair, replication and transcription machineries. A ready reference for biochemists, toxicologists, molecular and cell biologists, and geneticists seeking a better understanding of the impact of chemicals on human health.

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The Role of Human Rev7, the Accessory Subunit of Human DNA Polymerase Zeta, in Cell Survival and DNA Damage Induced Mutagenesis , Jessica A. Neal, 2008, , 212 pages. The presence of DNA damage can pose a considerable threat to cell survival because the high fidelity polymerases that are charged with the task of replicating DNA are often.

Long-Range Charge Transfer in DNA I, Volume 237 , Gary B. Schuster, Mar 24, 2004, Science, 219 pages. .

Mechanisms of DNA damage and repair implications for carcinogenesis and risk assessment, Michael G. Simic, United States. National Bureau of Standards, 1986, Medical, 578 pages. .

The Effect of Cytosine Methylation on the Structure and Repair of Carcinogenic DNA Adducts , Jessica B. Back, 2006, , 172 pages. The E. coli nucleotide excision repair complex, UvrABC, is able to repair a variety of DNA damage products. The repair efficiency of this complex appears to be influenced by.

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Chemical Carcinogens and DNA. Volume I, Philip L. Grover, 1979, Medical, 256 pages. .

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Oxidized Purine Lessons: Influences on DNA Ligation, Base Excision Repair, and DNTP Insertion , Xiaobei Zhao, 2008, , 161 pages. The biochemical experiments carried out in the dissertation indicated that Gh and Sp lesions are highly mutagenic in biological processes..

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