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**“Evaluating cell survival and DNA damage of cells that are deficient in a DNA repair gene
exposed to disinfectant chlorine dioxide”**

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Abstract:

Chlorine dioxide (ClO₂) is an effective biocidal product used to decontaminate medical utensils, swimming pools, and for the treatment of drinking water. ClO₂ is thought to be an oxidizing agent however the mechanism of action is not well known. The purpose of this research was to compare the cell survival rates and DNA damage of mouse embryonic fibroblasts (MEF₁) cells that are normal (MEF WT) versus cells that lack the essential DNA repair polymerase beta protein (MEF Pol β). An MTT which is a colorimetric assay for assessing cell survival and metabolic activity, was performed on the two cell lines with ClO₂ concentrations ranging from 2.5mM to 250mM. The ClO₂ concentration of 2.5mM showed a large gap between the MEF WT and MEF Pol β cells indicating increased cell death in the MEF Pol β cells. Flow cytometry data obtained from MEF WT cells stained with the anti-phosphor H2AX antibody showed that cells treated with ClO₂ experienced more DNA double stranded breaks than untreated cells. The results suggest that ClO₂ can be used as a possible treatment for cancer cells which are usually deficient in DNA repair genes because the MEF Pol β cells experienced more cell death.

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Autobiography:

My name is Syria McCullough and I am a junior at the University of New Haven. Next, I plan to attend graduate school and ultimately work in a lab pertaining to one of my majors, Forensic Science and Biology.

