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***In Vitro* Effects of Blue Laser Light as an Antimicrobial Agent on Microbial-Infected Fibroblast Cells**

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In Vitro Effects of Blue Laser Light as an Antimicrobial Agent on Microbial-Infected Fibroblast Cells

Francis Obeng Brenya¹ and Nicolette Nadene Houreld¹

¹Laser Research Centre, Faculty of Health Science, University of Johannesburg, Johannesburg, South Africa

Pseudomonas aeruginosa, *Staphylococcus aureus*, and *Streptococcus pyogenes* are key pathogens that delay healing and pose challenges due to their antibiotic resistance. Antimicrobial photobiomodulation (aPBM) using blue light (400-470 nm) has been shown to have antibacterial properties; however, its effects on mammalian cells are not well understood. We investigated the effect of blue laser light (470 nm, 82.7 mW/cm², 10 J/cm², 2 min) on bacteria-infected BJ-5ta fibroblast cells. BJ-5ta cells were co-cultured for 24 h with each of the three bacterial strains (1.50 x 10³ CFU/mL) and then exposed to blue light. Fibroblast cell viability and bacterial colony counts were assessed 24 h post-aPBM. Control cells (0 J/cm²) infected with *S. aureus* exhibited 95% fibroblast cell viability and increased bacterial counts (1.80 x 10⁵ CFU/mL). Control cells (0 J/cm²) infected with *S. pyogenes* and *P. aeruginosa* showed 89.5% fibroblast cell viability, with bacterial counts increasing to 3.00 x 10⁵ and 2.36 x 10⁵ CFU/mL, respectively. In irradiated (10 J/cm²) BJ-5ta cells infected with *S. aureus*, *P. aeruginosa*, and *S. pyogenes*, fibroblast cell viability was 89.2%, 94.6%, and 77.6%, respectively. As compared to the controls, bacterial counts decreased to 1.30 x 10⁵ CFU/mL, 1.35 x 10⁵ CFU/mL, and 1.20 x 10⁵ CFU/mL, respectively. Blue light (470 nm, 82.7 mW/cm², 10 J/cm²) induced bacterial death while preserving fibroblast cell viability after a single exposure. aPBM has the potential to address the medical challenges associated with infected wounds and open new avenues for future research.

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Primary author: Mr OBENG BRENYA, Francis (Laser Research Centre, Faculty of Health Science, University of Johannesburg, Johannesburg, South Africa)

Co-author: Prof. NADENE HOURELD, Nicolette (Laser Research Centre, Faculty of Health Science, University of Johannesburg, Johannesburg, South Africa)

Presenter: Mr OBENG BRENYA, Francis (Laser Research Centre, Faculty of Health Science, University of Johannesburg, Johannesburg, South Africa)

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