THE EFFECTS OF TOBACCO TREATED PORPHYROMONAS GINGIVALIS ON HUMAN EPITHELIAL CELLS

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Bacteria and tobacco are risk factors for periodontal diseases. Bacteriahost interactions play a critical role in disease development and progression. The effects of tobacco-treated bacteria such as Porphyromonas gingivalis on epithelial cells have not yet been examined. Therefore, P. gingivalis were treated with different tobacco products (nicotine, cigarette smoke condensate (CSC), and dissolvable smokeless tobacco (DST) strips) to determine the effects that they have on epithelial cells. P. gingivalis were grown with or without the products for 24 hours at 37°C. The cells were separated from the supernatant, washed with 0.9% NaCl and incubated at 60°C to kill the bacteria. Protein assays was performed to determine the protein concentration in the cell pellets and supernatants. Lactate dehydrogenase (LDH) assays are being used to measure the cytotoxicity of the cells and supernatants on epithelial cells in a dose dependent manner. Non-toxic amounts of the cell pellets and supernatants will be used to treat epithelial cells for 72 hours and the media analyzed by cytokine/growth factor protein arrays. The protein assays showed that CSC and nicotine treated P. gingivalis cells had less protein than the others. The total protein in the supernatant for the CSC treated bacteria was less compared to others. The protein data suggests that CSC and nicotine affect protein expression in and by the P. gingivalis cells. Tobacco-treated bacteria are hypothesized to increase the expression of proinflammatory cytokines/growth factors by the epithelial cells, thereby contributing to the inflammation seen in periodontal diseases.

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