

Public Abstract

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Title:FIRST CONTACT: POTENTIAL CONSEQUENCES OF EXPOSURE OF MACROPHAGE-LIKE CELLS TO DIVERGENT BACTERIA

The innate immune system must react to a wide variety of foreign stimuli, only a relatively small proportion of which may be classified as harmful. The goal of this thesis was to investigate interactions between macrophages and divergent bacteria with a broad range of adaptation to these mammalian cells that serve a central role at the interface of the innate and adaptive immune responses. The first study aimed to model innate macrophage responses to inactivated *Bacillus* spores, a proposed vaccine platform. This was done through cell culture of the macrophage-like cell line, J774, exposed to UV-irradiated *Bacillus thuringiensis* spores and molecules with known pathogen associated molecular patterns (PAMPs). Results indicated that *B. thuringiensis* spores activated J774 cells. J774 cells secreted cytokines TNF- α , IL-6 and IL-10 in response to these spores. IL-1 β , a pro-inflammatory cytokine, was only detected in J774 culture supernatant after exposure to viable *Bacillus* spores, or in the presence of UV-irradiated spores and an adjunct PAMP-containing molecule. The second study in this thesis concerned the interaction of parasitic bacteria with different mammalian host cell lines. *Ehrlichia canis* and *E. chaffeensis*, which are considered pathogenic and nonpathogenic to dogs, respectively, were chosen to begin work to test the possible role of this host-pathogen interface in the development of severe acute ehrlichiosis. The pathogenic pairing used was *E. canis* infecting the canine macrophage-like cell line, DH82. The non-pathogenic pairings were *E. canis* grown in the murine cell line, J774, and *E. chaffeensis* in the canine DH82 cell line. In both cases of comparison, the *Ehrlichia* grew more robustly when part of a pathogenic pairing.