Antimicrobial effects of lactoferrin and cannabidiol on *Borrelia burgdorferi* Dylan Haenel Department of Biology and Environmental Science/ Biotechnology Eva Sapi Ph.D.

Abstract

Robust antimicrobial resistance in pathogenic bacteria such as *Borrelia burgdorferi*, the Lyme disease bacteria, has proven difficult to alleviate in affected patients. Due to the bacterium's ability to form biofilms in unfavorable conditions, standard means of eliminating infection has shown to be ineffective and has lead the need to find a novel biologically active compound to combat these infectious agent. Lactoferrin previously has been shown to have antimicrobial effects on biofilms formed by *Pseudomonas aeruginosa* in cystic fibrosis patients. In this project, the effect of lactoferrin was tested against the biofilm form of *Borrelia burgdorferi*. The results show that the amount of biofil

Materials and Methods:

Low passage isolates of B31 strain of *B. burgdorferi* sensu stricto were cultured in Barbour-Stoner-Kelly H (BSK-H) complete medium supplemented with 6%

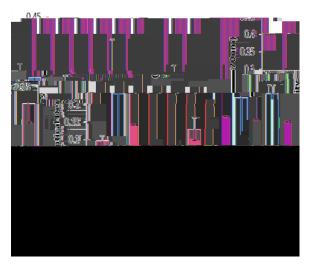


Fig 2. Quantitative analysis of Borrelia burgdorferi biofilm treated with different concentration of allicin using MTT viability assay. Negative control was a sample treated with vehicle (PBS buffer) and positive control had doxycycline (doxy) and Stevia, which was shown by previous research to have a significant effect on B. burgdorferi biofilm.

Conclusions and Future Work:

Data from this study showed that lactoferrin could have a significant effect on the viability of B. burgdorferi biofilm, while allicin did not show any potential effect. The mechanism of the antimicrobial effect of lactoferrin has been previously suggested to work by absorbing and therefore sequestering iron and manganese, so bacteria cannot utilize it. In a recent Nature publication, lactoferrin was demonsto7(e)D [(t) -1.149.59 1to-7()]12(s)-(i)g-7()5(s)-12(n)7(ei)-3(a)-3(n)57(t 1-12(y)25 c)-13onou7(e)Dnoi-12(s)2(

Acknowledgements:

I would like to thank Dr. Eva Sapi and the graduate students of the UNH Lyme Disease Research group for their continuous support and guidance. I would particularly like to thank Priyanka Theophilus MS and Jacintha Victoria MS.

Also, I would like to thank the University of New Haven for their support and allowing me an opportunity to present this project. This project was exclusively supported by the University of New Haven SURF Program.

Biography:

Dylan Haenel is a senior at the University of New Haven majoring in Biotechnology. He works as an Academic Peer Mentor for Bethel HTc n-1.1596nR7(yD1596nR7-7(a(a)-0)-4()](B))0(ks)-3(ae5bTw T* [o)-4(rn)-3(a)-njrn(hno)-12(l)-