



**Project Details:**

<b>Host Institution:</b>	University College Dublin (UCD)
<b>Location:</b>	Belfield, Dublin 4, Ireland
<b>College/Company:</b>	College of Science
<b>School/Unit:</b>	School of Biomolecular and Biomedical Science
<b>Website:</b>	<a href="http://www.ucd.ie/sbbs/">http://www.ucd.ie/sbbs/</a>

**Project Lead:**

<b>Name:</b>	Dr Siobhán McClean
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<b>Telephone Contact</b>	

**Project Title:**

Determining the host response to novel vaccine antigens

**Brief Project Description:**

Antimicrobial resistance is a massive growing problem in the fight against bacterial infections. The number of antibiotics that are effective at treating many bacterial infections is shrinking. Vaccines represent one of the best ways to prevent bacterial infections and have also been shown to reduce antimicrobial resistance (Mishra, Oviedo-Orta et al. 2012). In our lab we aim to develop anti-bacterial vaccines in order to prevent these difficult and challenging infections. We use a proteomic approach to identify highly effective vaccine antigens which prevent infections in mouse models. We have number of vaccine projects ongoing in our laboratory against antibiotic resistant infections such as respiratory infections that impact the lives of people with cystic fibrosis (McClean, Healy et al. 2016); the tropical infection, melioidosis (Casey and McClean 2015, Casey, Spink et al. 2016); the potentially lethal hospital acquired infection, *Klebsiella pneumoniae* and verotoxin resistant *E. coli*. Some of these are progressing towards human trials.

We have tested our vaccine antigens in mice and are currently examining the protective immunological responses, including antibody responses and cytokine responses in serum or immune cells. This project will involve using ELISA to determine the levels of antigen specific IgGs in immunised mice. In addition the host response will be further examined by exposing immune cells to antigen and evaluating the profile of cytokines produced using flow cytometry and/ or ELISA. Understanding how the antigens protect against infection is an important stage in progressing the vaccines towards human trials. The project would suit someone with an interest in immunology, microbiology or biochemistry.

**References:**

- Casey, W. T. and S. McClean (2015). "Exploiting molecular virulence determinants in *burkholderia* to develop vaccine antigens." *Curr Med Chem* **22**(14): 1719-1733.
- Casey, W. T., N. Spink, F. Cia, C. Collins, M. Romano, R. Berisio, G. J. Bancroft and S. McClean (2016). "Identification of an OmpW homologue in *Burkholderia pseudomallei*, a protective vaccine antigen against melioidosis." *Vaccine* **34**(23): 2616-2621.
- McClean, S., M. E. Healy, C. Collins, S. Carberry, L. O'Shaughnessy, R. Dennehy, A. Adams, H. Kennelly, J. M. Corbett, F. Carty, L. A. Cahill, M. Callaghan, K. English, B. P. Mahon, S. Doyle and M. Shinoy (2016). "Linocin and OmpW Are Involved in Attachment

of the Cystic Fibrosis-Associated Pathogen *Burkholderia cepacia* Complex to Lung Epithelial Cells and Protect Mice against Infection." *Infect Immun* **84**(5): 1424-1437. Mishra, R. P., E. Oviedo-Orta, P. Prachi, R. Rappuoli and F. Bagnoli (2012). "Vaccines and antibiotic resistance." *Curr Opin Microbiol* **15**(5): 596-602.

**Project Dates:**

From the end of May to August (specific dates and weekly hours can be agreed between the PI and the student directly over a 10-week period).

**Candidate Requirements:**

The project would suit someone with an interest in immunology, microbiology or biochemistry - a strong academic background required.