

# FMSA 2018

CROSSTALK BETWEEN INNATE AND ADAPTIVE IMMUNITY  
IN HEALTH AND DISEASE

Hosted by



**Name** Leow Chiuann Yee

**Current Position** Senior Lecturer

**Email** yee.leow@usm.my

**Office Number** 106

## **Qualifications**

PhD, University of Queensland, Australia, 2014

PgCert, Industrial Bioinformatics, Bioinformatics Institute of India, 2014

MSc, (Pharmacy) Universiti Sains Malaysia, 2006

B.App.Sc. (Hons.) (Biotechnology), Universiti Sains Malaysia, 2002

## **Affiliations**

Young Scientists Network - Academy of Sciences Malaysia (YSN-ASM)

Malaysian Society of Parasitology and Tropical Medicines

Australian Society for Parasitology

## **Research Interests**

Antigen discovery, immunoinformatics, molecular immunology, emerging infectious diseases (diagnosis, pathogenesis, and vaccine development)

## **Research Overview 1**

### **Molecular Immunological Approach to Shigellosis Vaccine Development**

In Malaysia, *Shigella* spp. is the third most common bacterial agent responsible for childhood diarrhoea. Like many other tropical diseases, shigellosis cannot be sustainably controlled by mass drug administration due to the possibility of antibiotic resistance. Therefore, a vaccine is desperately needed to combat this microbial pathogen. Prior to achieve this, the comprehensive understanding of immune response associated with shigellosis is critically important. *Shigella* infection represents an interesting paradigm of imbalance of the host immune mechanisms that regulate inflammation, and of bacterial strategies developed to escape killing by host immune cells that are equipped with a large repertoire of anti-microbial weapons to control microbial infection. Previous study showed that in vivo activation of T

cells in the blood in patients during shigellosis and a correlation between the degree of T-cell activation and disease severity, confirming the importance role of cell-mediated immune response during *Shigella* infection. To understand the role of cellular T cells in the pathogenesis of and protection against *Shigella* infection and to further the development of effective vaccines, knowledge about *Shigella* derived molecules or epitopes that may induce T-cell-specific responses is required.

## **Research Overview 2**

### **Identification of immune epitopes for the development of diagnostic assay and vaccines against pathogens**

Brucellosis is endemic in many developing countries and is caused by *Brucella* species that affect human, domestic and some wild animals. In livestock, *Brucella* causes abortion, still birth, reduced milk production and infertility, thus directly affecting livestock related food production. This zoonosis disease has made a significant economic impact in livestock that is widely distributed among cattle and related wildlife species. Human brucellosis has been reported in Malaysia since 2000. More importantly, food borne brucellosis in human was first reported in 2010 involving a seven-year boy after after consumed a *Brucella* infected raw milk. Recently, several outbreaks of brucellosis have been reported among humans in Malaysia which were associated with the consumption of raw cow's milk contaminated primarily with *B. abortus* and occasionally with *B. melitensis* and *B. suis*. Owing to the cross-reactivity of *Brucella* LPS with other Gram negative bacteria, and the lack of available vaccine, our primary research interest is to understand pathogenesis and immunology of brucellosis and hence contributing towards the diagnostic assays and vaccines development against this intracellular pathogen.

### **Selected Publications**

1. Karunaratne, D.S., Horne-Debets, J.M., Faleiro, R., Leow, C.Y., et. al., (2016). Programmed death-1 ligand 2 (PD-L2) is crucial for establishing enduring CD4+ Th1 immunity against malarial infections. *Immunity*. 45(2):333-45.
2. Leow, C.Y., Willis, C., Hofmann, A., and Jones, M.K. (2015). Structure-function analysis of apical membrane-linked molecules for treatment and control of schistosome parasites of humans: insights from studies into annexins. *British Journal of Pharmacology*. 172: 1653.
3. Wykes, M.N., Horne-Debets, J.M., Leow, C.Y. and Karunaratne, D. (2014). Malaria drives T cells to exhaustion. *Front. Microbiol.* 5:249.
4. Leow, C.Y., Willis, C., et. al. (2014). Crystal structure and immunological properties of the first annexin from *Schistosoma mansoni* - Insights into the structural integrity of the schistosomal tegument. *FEBS Journal*. 281(4): 1209-1225.
5. Cantacessi, C., Seddon, J.M., Miller, T.L., Leow, C.Y., et. al., (2013). A genome-wide analysis of annexins from parasitic organisms and their vectors. *Scientific Reports*. 3: 2893.
6. Hofmann, A., Osman, A., Leow, C.Y., Driguez, P., McManus, D.P., and Jones, MK. (2010). Parasite annexins - new molecules with potential for drug and vaccine development. *BioEssays*, 32: 967-976.