## TOLL IMMUNE PATHWAY IN THE MOSQUITO AEDES AEGYPTI

## Sang Woon Shin<sup>1</sup>, Guowu Bian<sup>1</sup>, Zhiyong Xi<sup>2</sup>, George Dimopolous<sup>2</sup>, and <u>Alexander S. Raikhel<sup>1</sup></u>

<sup>1</sup>Department of Entomology and the Institute for Integrative Genome Biology, University of California, Riverside California 92521, USA and <sup>2</sup>Malaria Research Institute, Johns Hopkins Bloomberg School of Public Health, 615 North Wolfe Street, Suite W4609, Baltimore, MD 21205

The availability of the A. aegypti genome enables researchers to perform comparative and functional analyses of the A. aegypti immune system. Transgenic approach, in combination with the RNAi technique is being used to elucidate the regulation of Toll immune pathway in the mosquito Aedes aegypti. By creating gain-of-function and loss-of-function transgenic *Aedes* strain we have shown that AaREL1, a homolog of *Drosophila* Dorsal, is a key regulator of Toll immune pathway in the adult mosquito A. aegypti. The Toll receptor and its ligand, Spätzle (Spz), links extracellular immune signals to the Toll intracellular transduction pathway. Genome analysis has shown that in the mosquito Ae. aegypti, there are five homologues of the Drosophila Toll (Toll1) receptor and three homologues of the Drosophila cytokine Spätzle (Spz). The transgenic approach, in combination with the RNAi technique, has shown that AeToll5 and Spz1C function as cytokine-receptor systems specific to the Toll receptormediated immune response following fungal challenge in the mosquito fat body. High throughput microarray analyses that focus on the determination of the REL1 regulated gene repertoire are being performed using gain-of-function REL1 transgenic mosquitoes.

Key words: *Aedes aegypti*, Toll pathway, microarray