## INTRAMOLECULAR INTEGRATION ASSAY VALIDATES INTEGRASE PHI C31 AND R4 POTENTIAL IN A VARIETY OF INSECT CELLS

Jakkrawarn Chompoosri<sup>1,2,4</sup>, Tresa Fraser<sup>1</sup>, Yupha Rongsriyam<sup>2</sup>, Narumon Komalamisra<sup>2</sup>, Padet Siriyasatien<sup>3</sup>, Usavadee Thavara<sup>4</sup>, Apiwat Tawatsin<sup>4</sup> and Malcolm J Fraser Jr<sup>1</sup>

<sup>1</sup>Department of Biological Sciences, Eck Institute of Global Health, University of Notre Dame, Notre Dame, IN, USA; <sup>2</sup>Department of Medical Entomology, Faculty of Tropical Medicine, Mahidol University, Bangkok; <sup>3</sup>Department of Parasitology, Faculty of Medicine, Chulalongkorn University, Bangkok; <sup>4</sup>National Institute of Health, Department of Medical Sciences, Nonthaburi, Thailand

Abstract. Phage  $\phi$ C31 and R4 integrases are site-specific and unidirectional serine recombinases. We have analyzed the ability of these integrases to mediate intramolecular integration between their *attB* and *attP* sites in 7 important insect cell lines as a means of predicting their relative mobility in the corresponding insect species. Both integrases exhibit significantly higher frequencies in *Drosophila* S2 cells than in the other insect cell lines examined, but do work well in all of the species tested. Our results, coupled with previous results of the activity of  $\phi$ C31 integrases in *D. melanogaster* and *Aedes aegypti*, suggest the family of serine catalyzed integrases will be useful site-specific integration tools for functional genome analysis and genetic engineering in a wide range of insect species.

Tel: 574-631-6209; Fax: 574-631-7413 E-mail: fraser.1@nd.edu

Correspondence: Malcolm J Fraser Jr, Eck Institute of Global Health, Department of Biological Sciences, University of Notre Dame, Indiana, 46556, USA.