## CLONING, CHARACTERIZATION, AND EXPRESSION OF VITELLINE PROTEIN BI AND ITS ENCODING GENE IN THE LIVER FLUKE, FASCIOLA GIGANTICA

Kritsana Homwutthiwong<sup>1</sup>, Ardool Meepool<sup>2</sup>, Rudi Grams<sup>3</sup>, Chaitip Wanichanon<sup>1</sup>, Vithoon Viyanant<sup>3</sup> and Prasert Sobhon<sup>1</sup>

<sup>1</sup>Department of Anatomy, Faculty of Science, Mahidol University, Bangkok; <sup>2</sup>Department of Medical Science, Faculty of Science, Burapha University, Chon Buri; <sup>3</sup>Faculty of Allied Health Sciences, Thammasat University, Prathum Thani, Thailand

**Abstract.** A cDNA containing a 813 bp open reading frame encoding vitelline protein BI (FgVPBI) of *Fasciola gigantica* was cloned. FgVPBI has 96% sequence identity with VPBI of *Fasciola hepatica* and 84% identity with VPBII *F. hepatica*. It is far less similar to eggshell precursor proteins of other trematode species, for example, 29% identity with *C. sinensis*. Northern blot hybridization of total RNA from adult parasites demonstrated a FgVPBI transcript with a size of 1,000 nucleotides. FgVPBI mRNA is localized in the vitelline cells in both vitelline glands and intrauterine eggs. Recombinant FgVPBI was expressed as a 31.5 kDa protein in *Escherichia coli* and used for production of a polyclonal antiserum in rabbits. The FgVPBI antiserum detected immunoblotted rFgVPBI and native eggshell precursor protein at molecular weights of 31.5 kDa and 31 kDa, respectively. Immunolocalization showed strong staining in the cytoplasm of vitelline cells, in eggshell globules and the shells of eggs.

Correspondence: Prof Prasert Sobhon, Department of Anatomy, Faculty of Science, Mahidol University, Bangkok, 10400 Thailand.

Tel: 66 (0) 2201 5406; Fax: 66 (0) 2354 7168

E-mail: scpso@mahidol.ac.th