A COMPARATIVE STATEMENT ON MOLECULAR APPROACH OF LARGE HSPS FROM MACROBRACHIUM ROSENBERGII

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ABSTRACT

This study reported a comparative account of large heat shock proteins (HSPs) namely HSP60, 70 and 90 from freshwater prawn Marcobrachium rosenbergii (Mr) at molecular level. The MrHSP60, 70 and 90 was 2158, 1998 and 2220 base pairs (bp) long that contain an open reading frame (ORF) of 1722, 1995 and 2157 bp, respectively. These MrHSP60, 70 and 90 ORFs encoded a polypeptide of 574, 665 and 719 amino acids and their respective molecular mass was 60.75, 71.40 and 82.65 kDa. MrHSP60 and 70 are mitochondrial, whereas MrHSP90 is cytoplasmic in nature. The bioinformatics analysis showed that MrHSP60 possess a chaperonin (cpn) 60 domain at 46-547 and a cpn 60 signature motif between 427 and 438. Similarly, MrHSP70 possessed a HSP70 domain at 21-624, whereas MrHSP90 contained a HSP90 domain at 188-719 along with a HSP90 family signature between 30 and 39. MrHSP60 and 70 are rich in hydrophobic (glycine) residues at C-terminal ends, whereas 90 is rich in hydrophilic (aspartate) residues. Homology and evolutionary analysis revealed that these three MrHSP60, 70 and 90 are very close to Macrobrachium nipponense, Cherax cainii and Palaemon carinicauda, respectively. The structural analysis showed that MrHSP60 carried maximum amino acids (52%) in helices and it carries cpn 60 signature in helical and coil regions. In MrHSP70, the nucleotide binding domain was distributed throughout the helices, sheets and coils, whereas in MrHSP90, the HSP90 family signature is present in the helical region. Tissue distribution results revealed that significantly (P<0.05) highest expression of MrHSP60, 70 and 90 was observed in haemocyte, gill and hapatopancreas, respectively. Moreover, their mRNA regulation upon exposure to microbial pathogens significantly (P<0.05) increased their expression. Overall, the study showed the potential involvement of large HSPs against microbial stress in immune function of M. rosenbergii.

KEYWORDS

M. rosenbergii; HSP; Tissue distribution; Pathogens; Gene expression

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