

Evaluating Genetic Variations on Brown Plant Hopper (*Nilaparvata lugens* (Stål)) Resistance Gene *Os06g0125132* of Rice for Molecular Marker Development

R.M.S.D.L. Abeyrathne¹, H.M.N.N. Herath¹, E.Y. Fernando²,
N.H.L.D.L.D. Nanayakkara¹ and D.V. Jayatilake^{1*}

¹*Department of Agricultural Biology, Faculty of Agriculture,
University of Peradeniya, Peradeniya 20400, Sri Lanka*

²*Department of Biological Sciences, Faculty of Applied Sciences,
Rajarata University of Sri Lanka, Mihinthale 50300, Sri Lanka*

**djayatilake@agri.pdn.ac.lk*

Brown plant hopper (BPH) is the most devastating insect pest of rice (*Oryza sativa* L.). Hence, cultivation of rice varieties carrying genetic resistance to BPH is the most promising approach for mitigating its damage. BPH resistance is controlled by many genes. The expression of *Os06g0125132* was found to be significantly upregulated under BPH infection and hence, a KASP marker for marker-assisted selection (MAS) was developed targeting a SNP in the intron region. Non-synonymous sequence variations in the exons that could alter the protein structure and/or its domains could be potential target sites for the development of alternative molecular markers compatible with technology-limited setups. In the current study, to identify such genetic variations, a haplotype analysis was conducted using the coding sequence of *Os06g0125132* (177bp), retrieved from 2,804 accessions sequenced in the 3K Rice Genomes Project. The extracted sequences were aligned using UGENE and four SNPs with $\geq 5\%$ occurrence frequency in the rice panel were identified (three SNPs at exon 1 and one SNP at exon 2) and three confirmed haplotypes (with $\geq 1\%$ occurrence frequency) were defined using DnaSP. Given all three haplotype-defining SNPs were non-synonymous, the respective putative amino acid sequences were modeled using the *ab-initio* algorithm *TrRefineRosetta* in Robetta. The modeled output suggested that all three putative proteins would fold into alpha helical structures (confidence of ≥ 0.8) and with no detectable 3-D structural difference. InterPro scan of the amino acid sequences detected a TMhelix, Transmembrane, Cytoplasmic and Non cytoplasmic domain each for all three putative proteins, indicating that the three identified non-synonymous SNPs had no impact on the domains. However, given the significance of *Os06g0125132* for BPH resistance, and its potential involvement in carbohydrate metabolism via the tricarboxylic acid cycle, further investigation is warranted to identify alternative sites for marker development targeting users in technology limited setups.

Keywords: Resistance to brown plant hopper, *Os06g0125132*, Haplotyping, Protein domains, Protein modeling

Funding from OWSD (Agreement No:4500384848) is acknowledged.