### EVALUATION OF BROWN PLANTHOPPER Nilaparvata lugens (stal) RESISTANCE IN Oryza nivara WILD RICE ACCESSIONS FOUND IN SRI LANKA

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#### Abstract

Brown planthopper (BPH), *Nilaparvata lugens* Stal is one of the most serious rice pests in Sri Lanka. The study evaluated the nature of BPH resistance in seventeen *O. nivara* (WRAC 01, 02, 04, 07, 11, 12, 14, 19, 21, 22, 24, 25, 35, 41, 46, 62, and 9864) accessions collected from different locations in Sri Lanka using standard seed box screening test. Ptb 33(resistant), Bg 379/2, Bg 300(moderately resistant) and Bg 380 (susceptible) were used as check varieties. BPH culture maintained at the Rice Research and Development Institute, Batalagoda on Bg 380 was used for the test. BPH damage was scored (0-to-9 scale) according to the standard evaluation system for rice, developed by International Rice Research Institute. According to the results WRAC 04, 41, 25 and Ptb 33 recorded as resistant (score 0-3), WRAC 46, 35, 24, 22, 21, 14, 7, 2,1, 9864 and Bg 379/2 as resistant to moderately resistant (score 3.1-4.0), WRAC 11, 12 and Bg 300 as moderately resistant (score 4.1-5.0), WRAC 19 and 62 as moderately resistant to moderately susceptible (score 5.1-6.0), while no checked accessions were recorded as susceptible to the BPH indicating potential of using *Oryza nivara* as a source of BPH resistance.

Keywords: Brown planthopper, Oryza nivara, Wild rice, Standard seed box screening test

## **1. INTRODUCTION**

Brown Plant Hopper (BPH) Nilarpavata lugens (stal) (Homoptera: Delphacidae) is one of the most important insect pest in rice. BPH is the most destructive and widespread insect through out the rice growing areas in Asia. In Sri Lanka nearly 5-10% of annual rice production is lost due to BPH damage (Nugaliyadda et al., 2001). At present, farmers mostly depend on chemical pesticides for the control of this pest (Kudagamage and Nugaliyadda, 1995). This dependence leads to many adverse effects. Host plant resistance is identified as the most effective way of BPH management. The varietal resistance is the most economic, least complicated and environmental friendly approach for the control of insect pest damages (Pathak and Kush, 1979). Rice is an annual grass that belongs to genus Oryza which includes 20 wild species. These wild relatives of rice constitute important genes for rice improvement; new sources of resistance to pests and diseases. Identification and characterizing of them may assist for future transfer of genes from wild rice to cultivated rice varieties. O. nivara is the most common and widely distributed wild rice species found in Sri Lanka (Hemachandra et al., 2010). It contains AA genome like O. sativa. Hence identification of BPH resistance in O. nivara is very useful for the rice variety improvement programs in Sri Lanka. Bioassay using standard seed box screening test has been effectively used for screening resistant lines (Heinrichs et al., 1985). Therefore, present study planned to evaluate BPH resistance in selected O. nivara accessions using standard seed box screening test method.

## 2. MATERIALS AND METHODS

#### 2.1. PLANT MATERIALS

Seventeen *O. nivara* accessions collected from different locations in Sri Lanka (Table 1) and maintained by the Rice Research and Development institute (RRDI), Bathalagoda and Ptb 33

(resistant check), Bg 380 (susceptible check), Bg 379/2, Bg 300(moderately resistant check) varieties were used for the study.

Accession No	Collected Location	District
WRAC 01	Bulunawewa	Matale
WRAC 02	Pelbediyawa	Matale
WRAC 04	Otththapahuwa	Anuradhapura
WRAC 07	Ilakkttuwa	Puttlam
WRAC 11	Kabaraya wewa, Poonewa	Anuradhapura
WRAC 12	Sangilikulama wewa	Anuradhapura
WRAC 14	Yakadapotha	Kurunegala
WRAC 19	Ihalawewa	Anuradhapura
WRAC 21	Inamaluwa	Matale
WRAC 22	Rangirigama	Matale
WRAC 24	A-9 road 105Km post	Anuradhapura
WRAC 25	A-9 road 105Km post	Anuradhapura
WRAC 35	Olugaskade tank	Anuradhapura
WRAC 41	Paranagama	Matale
WRAC 46	Mahiyanganaya	Badulla
WRAC 62	Meegahawewa, Weerawewa	Anuradhapura
9864	Matara	Matara

Table 1: The accession numbers and collected locations of O. nivara

#### 2.2. BPH BIOASSY

Experiment was conducted in the screen house at RRDI, Bathalagoda. BPH culture maintained at RRDI, Bathalagoda on susceptible rice variety, Bg 380 was used for the test. Bioassay was done following the standard seed box screening test method. Seeds of tested *O. nivara* accessions and check varieties were sown in a galvanized tray (60\*40\*10) filled with sterilized soil 5cm deep. The seeds were sown in 40cm long-rows and spaced 5cm among rows. When the seedlings were 7 days old (at 3 leaf stage), they were infested with 2<sup>nd</sup> instars nymphs at the rate of 3nymps/seedling. Complete Randomized design (CRD) with four replicates was applied for the test. The standard evaluation system for rice, developed by International Rice Research Institute (IRRI), in 1980 was used to score the BPH damage. BPH damage was rated by a 0-to-9 scale (Table 2).

**Table 2:** Standard evaluation system of BPH resistance in rice

Scale value	Type of resistance	
0.0-3.0	Resistant(R)	
3.1-4.0	Resistant/Moderately Resistant(R/MR)	
4.1-5.0	Moderately Resistant(MR)	
5.1-6.0	Moderately Resistant/Moderately Susceptible(MR/MS)	
6.1-7.0	Moderately Susceptible(MS)	
7.0<	Susceptible(S)	

### **3. RESULTS AND DISCUSSION**

BPH resistance has been identified in a wide range of traditional rice varieties and wild rice species (Pathak & Khush 1979 and Heinrichs *et al.* 1985). Scientists have already characterized and exploited some resistant species and upgraded cultivated rice varieties. To

date these varieties play a significant role in controlling BPH outbreaks in rice cultivations. However development of virulent BPH populations is becoming a serious threat for rice cultivation at present. Thus, constant search for resistant species is very important. Several reports have highlighted wild rice species of *O. latifolia*, *O. nivara*, *O. punctata* and *O. minuta* as important sources of BPH resistance (Wu *et al.* 1986).

Results of the standard seed box screening test showed that tested *O. nivara* accessions were almost resistant to the Sri Lankan BPH. Lowest mean damage scores were recoded in Ptb 33, WRAC 04, WRAC 25, WRAC 41, WRAC 01, WRAC 02, WRAC 07, WRAC 14, WRAC 21, WRAC 22, WRAC 24, WRAC 35, WRAC 46, 9864 and Bg 379/2 accessions and varieties indicating their resistant nature to the BPH. Highest mean damage score was recorded in variety Bg 380(Fig.1).

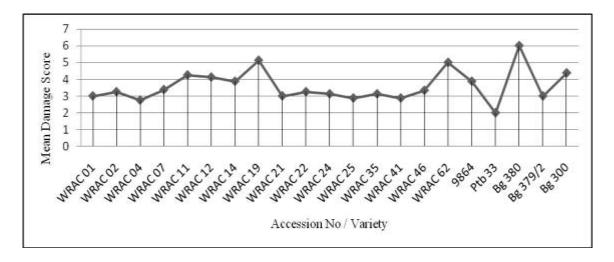


Figure 1: Variation of mean damage score in tested accessions and varieties

According to the scale of the standard evaluation system of BPH resistance, Ptb 33, WRAC 04, WRAC 25 and WRAC 41 recoded as resistant to BPH as all the other *O. nivara* accessions were categorized under the resistant to moderately resistant, moderately resistant and moderately resistant to moderately susceptible groups (Table 3).

Type of resistance	Accession No/Variety
Resistant(R)	WRAC 04, WRAC 25, WRAC 41, Ptb 33
Resistant/Moderately	WRAC 01, WRAC 02, WRAC 07, WRAC 14, WRAC 21,
Resistant(R/MR)	WRAC 22, WRAC 24, WRAC 35, WRAC 46, 9864, Bg 379/2
Moderately Resistant(MR)	WRAC 11, WRAC 12, Bg 300
Moderately Resistant/Moderately	
Susceptible(MR/MS)	WRAC 19, WRAC 62
Susceptible(S)	Bg 380

Table 3: Type of resistance recorded in tested O. nivara accessions and check varieties

According to the results no checked *O. nivara* accession was recorded as susceptible to the BPH indicating potential of using *O. nivara* as donors of BPH resistance for the future rice variety improvement program in Sri Lanka.

### 4. CONCLUSION

Results of the BPH bioassay revealed that most of the tested *O. nivara* accessions were resistant to the Sri Lankan BPH. Resistance observed in these accessions may be due to already identified BPH resistant genes or due to new genes. So further experiments should be carried out with markers for already identified BPH resistant genes to detect whether that the resistance observed in *O. nivara* accessions is due to existing resistant genes or new genes to avoid genetic uniformity of BPH resistance in cultivated rice in future.

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