Short Communication

High Heritability in a Resistant Barley Genetic Source to Spot Blotch (Cochliobolus sativus)

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Abstract. The objective of the present research was to assess and understand the heritability of the resistant barley genetic source cv. Banteng to spot blotch SB disease caused by *Cochliobolus sativus*. A cross was made between this resistant cultivar and the universally susceptible cv. WI 2291. Analysis of variance for the studied trait indicated highly significant differences among cultivars. High broad sense heritability was found ($H \approx 88 \%$). However, in all cases, the results obtained for the F₂ plants demonstrated that the observed segregation pattern fitted 1:15 ratios.

Keywords: Hordeum vulgare, Cochliobolus sativus, fungus resistance, inheritance

Spot blotch (SB) caused by *Cochliobolus sativus* (Ito & Kurib.) Drechsler ex Dastur [anamorph: *Bipolaris sorokiniana* (Sacc.) Shoem.] is economically one of the most important fungal diseases of barley (*Hordeum vulgare* L.) throughout the world (Mathre *et al.*, 2003).

The economic damage caused by SB and planting of resistant genotypes has been studied extensively by many researchers (Zhou and Steffenson, 2013; Ghazvini and Tekauz, 2008; Joshi *et al.*, 2007a; Bilgic *et al.*, 2006; 2005; Arabi, 2005a; Arabi and Jawhar, 2004; 2003; Steffenson *et al.*, 1996).

The present study therefore, was initiated to investigate the inheritance pattern of SB resistance in the German barley Banteng cultivar to design suitable strategies to enhance resistance of barley cultivars.

Single plant selections of resistant (Banteng) and susceptible (WI2291) cultivars were multiplied and used in the cross. Resistant parent Banteng (a germplasm cultivar introduced from Germany) was crossed with the universal susceptible WI2291 (originated from the Waite Institute, Glen Osmond, Australia) which is otherwise higher yielding with good agronomic performance.

Parents and F_1 progenies (29 plants) were evaluated for resistance to SB under an induced epiphytotic created in the field at station, west of Damascus , Syria under rainfed conditions (500mm rainfall). Seeds were planted in a randomised complete block design, with three replicates. Plots of the F_1 generation consisted of two 2-m rows seeded 25 cm apart with 30 cm between plots. The susceptible barley cultivar WI 2291 was planted in the alleys and borders, two weeks before sowing the experiment to enhance the spread of disease. Soil fertilizers were drilled before sowing at a rate of 50 kg/ha urea (46% N) and 27 kg/ha superphosphate (33% P). The progenies of the cross were advanced to the F_2 generation (457 plants) following the method described by Joshi *et al.* (2004) where a random plant in each generation from each line was harvested for advancing the generation.

A mixture of equal ratio of pure aggressive isolates of *C. sativus* (Arabi and Jawhar, 2004) was used to inoculate the parents as well as plants of the F_1 and F_2 generation. A spore suspension (approximately 2 x 10⁴ spores/mL) containing the surfactant Tween 20, was uniformly sprayed onto plants during the evening hours by using a hand-held atomizer, then plants were covered with polyethylene for 3 days to maintain humidity for infection and subsequent disease development (Joshi *et al.*, 2007a; 2007b).

Percentages were transformed into a 1-4 scale. Cultivars that scored less than 26% were considered resistant, between 26 and 50% as moderately susceptible; between 60 and 70% as susceptible, and those having higher than 70 as highly susceptible (Joshi *et al.*, 2007a; 2007b).

Statistical analysis. For every line, disease scores of all the plants including the most susceptible and most resistant ones were recorded. Broad-sense heritability

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 (h^2b) was used to eliminate the influence of environment on the expression of disease severity (Mulitze and Baker, 1995) and computed as follows:

$$\begin{split} h^{2}{}_{b} &= \sigma^{2}{}_{G} \; / \; \sigma^{2}{}_{P} \; x \; 100, \; \; \sigma^{2}{}_{G} &= \sigma^{2}{}_{P} - \sigma^{2}{}_{E}, \; \; \sigma^{2}{}_{E} &= \sigma^{2}{}_{P_{1}} + \\ \sigma^{2}{}_{P_{2}} + \sigma^{2}{}_{F_{1}} / 3, \; \; \sigma^{2}{}_{P} &= \sigma^{2}{}_{G} - \sigma^{2}{}_{E} \end{split}$$

where:

 σ_{P}^{2} = phenotypic; σ_{E}^{2} = environmental; σ_{G}^{2} = genetic; $\sigma_{P_{1}}^{2}$ = Banteng; $\sigma_{P_{2}}^{2}$ = WI 2291 variance.

Significant differences (p = 0.05) in mean severity values were detected between the two barley cultivars. The cv. Banteng had a mean disease severity of 13.45 %, whereas the susceptible cv. WI 2291 had a mean disease severity 73.44% (Table 1). The general combining ability (GCA) mean square was significant at p = 0.05, which shows the variability of (GCA) of the parent. Estimates of GCA effects of each parental genotype are presented in Table 1. Compared to the parents, the SB severity of the 29 F₁ plants appeared to be intermediate (Fig. 1), indicating the absence of dominance for the genes governing resistance.

Moreover, F_2 progeny distributions in the cross (Fig. 1) indicated that resistance genes interacted in an additive manner. F_2 progeny exhibited a wide range of SB severities from 2 to 88 % (Fig. 1). As discrete classes were observed in the distribution of SB reaction in the F_2 , the plants were classified within parental classes using phenotypic values observed for the parents grown in the same environment. This was only applicable to the class of the resistance of parent (≤ 20 %), as the range of variation of the standard resistance parent was similar to that of the F_1 . However, in all cases, the results obtained for the F_2 plants demonstrated that the observed segregation pattern fitted 1:15 ($x^2 = 7.11$; $\alpha = 0.10$) ratios.

Table 1. Range and mean spot blotch (SB) severity (%) of barley parents during two years of testing under different field conditions, and the general combining ability (GCA) estimates in the F_1 generation^y

Genotype	Origin	SB reaction		GCA	
		Range	Mean	Seedling	Adult plants
Banteng	Germany	1.14-22	13.45	-0.90*	-0.92*
W1229	Australia	54-90	73.44	0.75*	0.72*
				0.18	0.15
F ₁		35-50	42.00		
F_2		6-80	41.64		

*Significant at p = 0.05, ^y =Arabi, 2005a



Fig. 1. Histogram of % disease severity in cv. Banteng (a), cv. WI 2291 (b), F_1 (c) and F_2 (d).

Chromosomes 1S and 5S have been identified in barley as harboring loci for SB resistance at the adult stage. Information regarding the genes controlling durable resistance is of a paramount importance to breeders. Some of the previous studies concerning the inheritance of SB resistance in barley also indicated the control by many genes (Kuldeep *et al.*, 2008), who reported that the heritabilities of SB resistance were moderately high and ranged from 0.77 to 0.83 across four environments.

The results of the present study indicate that the heritability of resistance in cv. Banteng was high at 88%. Thus, this cultivar should be considered as a possible donor in future breeding efforts. Since heritability was high, effective selection could be applied in early generations. Furthermore, the cv. Banteng was resistant to net blotch, barley stripe (Arabi, 2005b) and powdery mildew (Arabi and Jawhar, 2012), which could give it special interest in barley breeding programmes.

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