



Development of classification criteria for resistance to soybean rust and differences in virulence among Japanese and Brazilian rust populations

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ABSTRACT

In recent years soybean rust, caused by *Phakopsora pachyrhizi* has become one of the most serious threats to soybean production in Brazil. Breeding lines and varieties have been selected for resistance to soybean rust in Asia. However, differences in virulence between Asian and Brazilian rust populations should be considered in order to select and use resistant resources from Asia. Here, we suggest criteria for distinguishing resistant from susceptible types by the analysis of four resistance characters: frequency of lesions having uredinia, number of uredinia per lesion, frequency of open uredinia, and sporulation level, determined by the utilization of 63 genotypes. Under growth chamber conditions, a set of 13 soybean varieties were exposed to three rust populations—one from Japan and two from Brazil—and evaluated for the resistance characters mentioned above. The Japanese and Brazilian populations clearly differed in virulence, as did the two Brazilian populations. Only two resistance genes, *Rpp4* from PI459025 and *Rpp5* from Shiranui, commonly conferred resistance on all three rust populations. The number of resistant varieties or resistance genes useful in both countries appears limited. Therefore, a resistant cultivar that is universally effective against soybean rust should be developed by pyramiding some major resistance genes and by introducing horizontal resistance.

Keywords: *Phakopsora pachyrhizi*, lesion type, pathogenicity, resistant variety.

RESUMO

Desenvolvimento de critério de classificação da resistência à ferrugem asiática da soja e diferenças de virulência entre populações do Japão e do Brasil

Nos últimos anos a ferrugem asiática, causada pelo fungo *Phakopsora pachyrhizi* tornou-se uma das mais sérias ameaças a produção de soja Brasileira. Linhagens melhoradas e variedades têm sido selecionadas para a resistência à ferrugem da soja na Ásia, entretanto para a seleção e utilização dessas fontes de resistência, diferenças de virulência entre populações Asiáticas e Brasileiras desse fungo devem ser consideradas. Neste trabalho sugerimos um critério para se distinguir resistência de susceptibilidade pela análise de quatro caracteres de resistência: frequência de lesões contendo urédias, número de urédias por lesão, frequência de urédias abertas e nível de esporulação determinados pela utilização de 63 genótipos. Sob condições controladas em câmaras de crescimento, treze variedades de soja foram expostas a três populações de fungos — uma população proveniente do Japão e duas populações provenientes do Brasil—e avaliadas quanto aos caracteres de resistência mencionados acima. As populações Brasileiras diferiram entre si claramente quanto a virulência e em relação à população de isolados do Japão. Apenas dois genes de resistência, *Rpp4* presente na variedade PI459025 e *Rpp5* presente na variedade Shiranui conferiram resistência as três populações da ferrugem. O número de variedades ou genes resistentes úteis em ambos os países parece ser limitado. Assim, um cultivar universalmente efetivo contra a ferrugem da soja deveria ser desenvolvido pela piramidação de genes maiores de resistência e pela introdução de resistência horizontal.

Palavras-chave: *Phakopsora pachyrhizi*, tipo de lesão, patogenicidade, variedade resistente.

INTRODUCTION

Soybean rust, caused by *Phakopsora pachyrhizi* Syd. & P. Syd., has been observed in Japan for more than 100

years (Hennings, 1903); it has also been detected in many Asian countries and reached South American countries such as Brazil, Paraguay, Argentina, Bolivia and Colombia in recent years (Rossi, 2003; Yorinori et al., 2005). Since 2001, when soybean rust was discovered in Brazil, it has become one of the most serious threats to soybean (*Glycine max* (L.) Merr.) production in this country because it can potentially

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reduce the soybean yield by more than 75 % (Navarro et al., 2004; Ivancovich, 2005; Yorinori et al., 2005).

The development of resistant varieties is an efficient way to manage soybean rust. Five major resistance genes—*Rpp1*, *Rpp2*, *Rpp3*, *Rpp4*, and *Rpp5*— have been identified in Asian soybean varieties (Hartwig & Bromfield, 1983; Hartwig, 1986; Garcia et al., 2008; Pierozzi et al., 2008). Most of the major resistance genes are considered race-specific (Bromfield & Hartwig, 1980, 1983; Bromfield, 1984; Hartwig, 1986) and the pathogen of soybean rust presents high race variability (Yamaoka et al., 2002; Oliveira et al., 2005). Thus, although many breeding lines and cultivars have been selected for resistance to soybean rust in Asia, they cannot be utilized without knowledge of the differences in virulence between South American and Asian rust populations. In order to use these resistant materials from Asia in the Brazilian breeding program, we need to determine the differences in virulence between Asian and South American rust populations by using a single, comprehensive method of virulence evaluation.

The “lesion type” classification, based on lesion color and the number of sporulating uredinia, has been used to identify the virulence of soybean rust isolates (Bromfield, 1984). However, lesion color can vary a lot and some resistant lesions that do not sporulate can be found in tan-colored lesions. In addition, continuous variation of lesion color among varieties makes it difficult to group all phenotypes into a limited number of lesion types, such as RB (Resistant) and TAN (Susceptible) (Kato & Yorinori, 2008). On the other hand, lesion color is known to be controlled by resistance genes. For example, quantitative trait locus (QTL) analysis of lesion color in 120 F₂ plants revealed that major resistance genes, *Rpp2* from PI230970 and *Rpp4* from PI459025, can genetically contribute to the darkening of lesion color under greenhouse conditions (Yamanaka et al., 2008). Thus, lesion color should also be considered when selecting resistant genotypes. Bonde et al. (2006) suggested that the number and size of uredinia are desirable indexes for detecting resistance derived from major genes and also partial resistance to soybean rust. These indexes are represented by numerical values and are considered more suitable than the index based on lesion color for deciding resistance because of their objectivity.

To choose appropriate traits that would enable us to classify levels of soybean rust resistance in soybean cultivars, we investigated five traits related to resistance in 63 genotypes infected with a Japanese rust population. We also investigated the application of these traits to the determination of differences in virulence among three rust populations collected in Japan and Brazil during 2007 and 2008.

MATERIALS AND METHODS

Japanese and Brazilian rust populations

In this study, we used non-purified isolates derived from single spore isolation and rust populations putatively containing various races. This choice was because our objective was to clarify the regional differences in virulence

of soybean rust by comparing dominant races in each region, rather than some specific races. The Japanese rust population (JRP) was collected from rust-infected soybean in an experimental field of the National Agricultural Research Center (NARO), Tsukuba, Ibaraki, Japan in September 2007. Infected leaflets were collected in plastic bags. Urediniospores on more than 20 leaflets were mixed and put into microtubes. Open microtubes containing urediniospores were placed on silica gel for 1 day and then stored at –80 °C. Urediniospores of JRP were propagated on *Glycine max* cv. Tachinagaha before they were used for inoculation. Brazilian rust population 1 (BRP-1) was obtained from a greenhouse at the Brazilian Agricultural Research Corporation, National Soybean Research Center (Embrapa Soja), Londrina PR, Brazil in January 2008. BRP-1 was originally derived from a commercial field in the state of Mato Grosso and had been maintained on *Glycine max* cv. BRSMS Bacuri in the greenhouse. BRP-1 has been used in previous studies (Yamanaka et al., 2007b; Kato & Yorinori, 2008; Silva et al., 2008; Yamanaka et al., 2008). Brazilian rust population 2 (BRP-2) was also obtained from a greenhouse at Embrapa Soja in August 2008. BRP-2 had the same origin as BRP-1 but had been maintained on various soybean genotypes in a different greenhouse at EMBRAPA Soja. BRP-2 was imported to Japan (Import permit No. 20Y157) and used for the experiments, together with JRP, at the Japan International Research Center for Agricultural Sciences (JIRCAS). The experiment using BRP-1 was performed at EMBRAPA Soja. All experiments were carried out according to the conditions described below. Urediniospores of each rust population were suspended in distilled water with 0.04% (v/v) of polyoxyethylene sorbitan monolaurate (Tween 20, Sigma) to prepare suspensions of about 50,000 spores/mL for inoculation.

Plant materials and growth conditions

Sixty-three soybean genotypes (cultivars/lines), including materials previously reported as having resistance to soybean rust (Table 1), were evaluated for resistance to the JRP. Thirteen varieties (entries 1 to 13 in Table 1), derived and multiplied from a single seed from the Embrapa Soja gene bank, were used to identify virulence differences among the 3 rust populations. Nine soybean cultivars (entries 1 to 9) were known to have major resistance genes and were therefore considered to be resistance standards. Two cultivars, TK5 and Wayne (entries 12 and 13) were used as susceptible materials. The other two PI lines (entries 10 and 11) previously reported as resistant to a mixture of isolates from four regions (Miles et al., 2006), were also included as resistant materials in a set of standard variety. Sixty-three soybean genotypes for infection from JRP were divided into a total of five experimental sets consisting of 12 or 13 genotypes. Each single experiment using thirteen varieties was performed for infection from BRP-1 and

TABLE 1 - Soybean genotypes used in this study. All 63 were used initially to test the Japanese rust population, and entries 1 to 13 were also used to test the Brazilian rust populations

Entry	Genotype	Characteristics	Source ⁴	Reference
1	PI200492 (Komata)	Standard variety (SV) having <i>Rpp1</i>	EMBRAPA	Hartwing and Bromfield, 1983
2	Tainung 4	SV having <i>Rpp1</i>	EMBRAPA	Yamaoka et al., 2002
3	PI587880A (Huang Dou)	SV having <i>Rpp1</i>	EMBRAPA	Ray et al., 2009
4	PI587886 (Bai Dou)	SV having <i>Rpp1</i>	EMBRAPA	Ray et al., 2009
5	PI230970 (No.3)	SV having <i>Rpp2</i>	EMBRAPA	Hartwing and Bromfield, 1983
6	PI417125 (Kyushu 31)	SV having <i>Rpp2</i> ¹	EMBRAPA	Nogueira et al., 2008
7	PI462312 (Ankur)	SV having <i>Rpp3</i>	EMBRAPA	Hartwing and Bromfield, 1983
8	PI459025 (Bing Nan)	SV having <i>Rpp4</i>	EMBRAPA	Hartwing, 1986
9	Shiranui	SV having <i>Rpp5</i> ²	EMBRAPA	Garcia et al., 2008
10	PI416764 (Akasaya)	SV	EMBRAPA	Arias et al., 2008
11	PI587905 (Xiao Huang Dou)	SV	EMBRAPA	Miles et al., 2006
12	TK5	SV of susceptible control	EMBRAPA	Yamaoka et al., 2002
13	Wayne	SV of susceptible control	EMBRAPA	Yamaoka et al., 2002
14	PI459025A	Resistant variety putatively having <i>Rpp4</i>	EMBRAPA	Pierozzi et al., 2008
15	PI594767A	Resistant variety	EMBRAPA	Miles et al., 2006
16	GC00002-100	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1989
17	GC00138-29	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1989
18	GC60020-8-7-7-18	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1989
19	GC84040-16-1	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
20	GC84051-9-1	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
21	GC84058-21-4	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
22	GC85037-2-3-5-1	Resistant variety developed by AVRDC	AVRDC	Yorinori, 2008
23	GC85039-1-2-1-1	Resistant variety developed by AVRDC	AVRDC	-
24	GC86004-9	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
25	SS86045-23-2	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
26	GC87012-10-B-5	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
27	GC87016-11-B-2	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1992
28	SRE-B-15C	Resistant variety developed by AVRDC	AVRDC	AVRDC, 1989
29	SRE-D-11C	Resistant variety developed by AVRDC	AVRDC	Oloka et al., 2008
30	Xiao Jing Huang	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
31	Niu Mao Huang	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
32	Qin Dou	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
33	Da Bai Qi	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
34	6611	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
35	Himedaizu	Resistant variety screened from genetic resources	JIRCAS	Yamanaka et al., 2007a, 2008
36	Lu Pi Dou	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
37	Hei Dou	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
38	Da Li Zi	Resistant variety screened from genetic resources	JAAS	Yamanaka et al., 2007a, 2008
39	Hougyoku	Resistant variety	NICS	-
40	Sachiyutaka	Resistant variety	NICS	-
41	FT2	Resistant variety putatively having <i>Rpp3</i> ³	EMBRAPA	Yamanaka et al., 2007b
42	Davis	Susceptible variety	EMBRAPA	Yamanaka et al., 2007b
43	Kinoshita	Resistant variety having <i>Rpp5</i> ⁴	EMBRAPA	Garcia et al., 2008
44	Abura	Resistant variety	EMBRAPA	Laperuta et al., 2008
45	RI75	Susceptible variety	EMBRAPA	-
46	BR01-17996	Breeding line resistant to soybean rust	EMBRAPA	-
47	BR01-18437	Resistant line having single recessive major gene	EMBRAPA	Pierozzi et al., 2008
48	BRSMS-Bacri	Resistant variety putatively having <i>Rpp3</i> ³	EMBRAPA	Kato and Yorinori, 2008
49	EMBRAPA48	Susceptible variety	EMBRAPA	Ribeiro et al., 2008
50	Misuzudaizu	Susceptible variety	Chiba Univ.	Yamanaka et al., 2001
51	Moshidou Gong 503	Susceptible variety	Chiba Univ.	Yamanaka et al., 2001
52	BRS184	Susceptible variety	EMBRAPA	Yamanaka et al., 2008
53	BRS231	Tolerant variety	EMBRAPA	Yamanaka et al., 2008
54	BB17	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
55	BB18	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
56	BB19	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
57	BB20	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
58	BB21	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
59	BB22	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
60	BB25	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
61	BB26	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
62	BB28	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008
63	BB29	Recombinant Inbred Line from BRS184 X BRS231	EMBRAPA	Yamanaka et al., 2008

¹PI417125 has been confirmed to have *Rpp2* by allelism testing with PI230970 (Nogueira et al. 2008); ²Kinoshita and Shiranui have a single resistance gene, which is mapped as *Rpp5* (Garcia et al. 2008); ³FT2 and BRSMS-Bacri putatively have the same resistance gene, *Rpp3*, located in linkage group C2 (Monteros et al. 2006; Arias et al. 2008); ⁴EMBRAPA: Brazilian Agricultural Research Corporation, AVRDC: Asian Vegetable Research and Development Center, JAAS: Jilin Academy of Agricultural Sciences, JIRCAS: Japan International Research center for Agricultural Sciences, NICS: National Institute of Crop Science.

BRP-2, respectively. All sets of experiments were carried out using the following growth conditions.

Three plants per genotype were grown in pots filled with soil from the field, in a growth chamber at 24°C and under a 14-h light photoperiod. Luminance at the soil surface of the chamber was 19,000 to 23,000 lux, provided by fluorescent lamps. Plants were inoculated with spore suspension when they had reached the V3 to V4 growth stage (Fehr et al., 1971). The lower surface of first trifoliolate leaves was scrubbed by the fingers with distilled water for about 10 seconds per leaflet in order to receive the spore suspension well, and then the spore suspension (about 0.1mL per one leaf) was spread homogeneously on the lower surface of the leaves with a paintbrush. Inoculated plants were covered with plastic bags to maintain humidity and the temperature was kept at 24°C for 24 h. Then the plastic bags were removed and the inoculated plants were transferred into the growth chamber under the same temperature and luminance conditions as described above for another 13 days until evaluation. Each experiment respectively for the infections from JRP, BRP-1, and BRP-12 was performed in the present study.

Evaluation

Lesion color (LC) and number of uredinia per lesion (NoU), which have been used to determine resistance to soybean rust (Bromfield 1984.), were scored two weeks after inoculation. In addition, frequency of lesions that had uredinia (%LU), frequency of open uredinia (%OU), and sporulation level (SL) were also evaluated. Thus a total of five resistance characters were evaluated under a stereomicroscope. LC of each lesion was classified into six categories in accordance with the method of Kato & Yorinori (2008): 1 (very dark) to 6 (very light). SL of each lesion was classified into 0 (no spores) to 3 (abundant spores) (Figure

1). Phenotypic data for %LU and %OU were obtained from at least 30 lesions per genotype and phenotypic data for LC, NoU and SL were decided by the average values of at least 30 lesions from three plants. Pearson’s correlations between these five characters and their significance were also calculated.

Phenotypic values in these resistance characters were respectively classified into resistant or susceptible types. Then five resistance categories of soybean genotypes: Immunity, Highly resistant, Resistant, Slightly resistant, or Susceptible, were determined based on the types in the resistance characters. We judged the differences in virulence among rust populations by these five categories of resistance in soybean genotypes.

RESULTS

Phenotypes of resistance against JRP

Phenotypic values of the five resistance characters in the 63 soybean genotypes, which had been infected with JRP, were arranged on the basis of their SLs (Table 2). Then frequency distributions of the five resistance characters in the 63 genotypes (Figure 2) were created. All the frequency distributions except that for LC were bimodal, and the genotypes could mostly be clearly classified into two classes: resistant and susceptible, although some showed intermediate values. On the other hand, continuous variation in LC was observed. In addition, in three genotypes, SRE-D-11C, PI587905, and Shiranui showed imperfect formation of lesions that might have been caused by very strong resistance (e.g. SRE-D-11C in Figure 3). Therefore, various types of lesions in terms of LC and SL were also observed (Figure 3).

All correlations among the five characters were significant at the level of $P < 0.001$ by t-test. Correlations among the four characters excluding LC were very high, with values of 0.868 to 0.953; LC had lower correlations

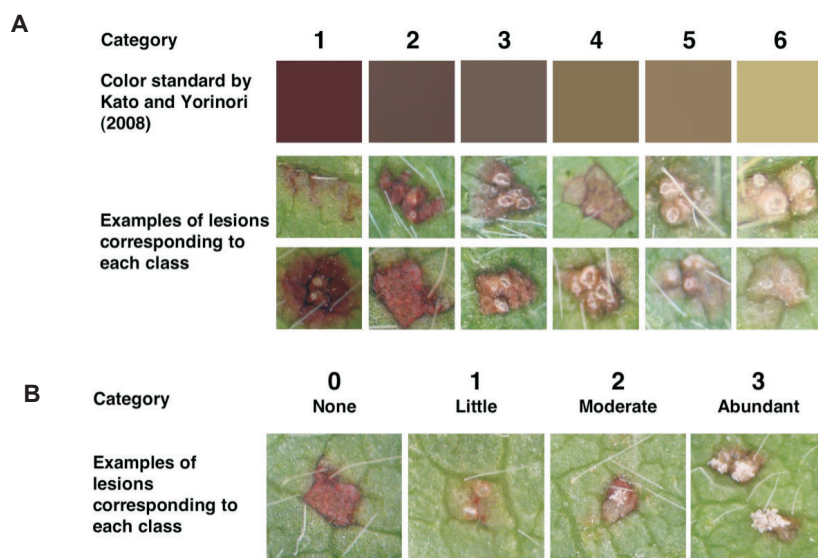


FIGURE 1 - Standards for lesion color **A.** and sporulation level **B.** Pictures of lesions for the color standard were taken after removal of the urediniospores to clearly reveal the color. Imperfect formation of lesions might have been the result of a hypersensitive response; lesions with this type of pigmentation were classified into lesion color class 1.

TABLE 2 - Phenotypes of the five resistance characters against the Japanese rust population in 63 soybean genotypes. List arrangement is determined by the values of the 5 resistance characters in the following order: first: sporulation level (SL); second: number of uredinia (NoU); third: frequency of lesions with uredinia (%LU); fourth: frequency of open uredinia (%OU); fifth: lesion color (LC). All values are rounded off to the first decimal place

Entry	Genotype	LC	%LU ¹	NoU ²	%OU ³	SL ⁴	Resistance classification ⁵
3	PI587880A	—	—	—	—	—	Immune
29	SRE-D-11C	1.0	0.0%	0.0	0.0%	0.0	Highly resistant (almost immune)
11	PI587905	2.3	0.0%	0.0	0.0%	0.0	Highly resistant (almost immune)
9	Shiranui	2.6	0.0%	0.0	0.0%	0.0	Highly resistant (almost immune)
15	PI594767A	1.3	0.0%	0.0	0.0%	0.0	Highly resistant
39	Hougyoku	1.5	0.0%	0.0	0.0%	0.0	Highly resistant
43	Kinoshita	1.5	0.0%	0.0	0.0%	0.0	Highly resistant
22	GC85037-2-3-5-1	1.9	0.0%	0.0	0.0%	0.0	Highly resistant
10	PI416764	1.9	0.0%	0.0	0.0%	0.0	Highly resistant
2	Tainung 4	2.1	0.0%	0.0	0.0%	0.0	Highly resistant
1	PI200492	2.8	0.0%	0.0	0.0%	0.0	Highly resistant
17	GC00138-29	1.9	1.7%	0.0	0.0%	0.0	Resistant
16	GC00002-100	1.9	1.9%	0.1	0.9%	0.0	Resistant
44	Abura	1.2	7.1%	0.1	0.0%	0.0	Resistant
21	GC84058-21-4	2.7	13.9%	0.1	0.0%	0.0	Resistant
48	BRSMS-Bacri	1.1	12.5%	0.8	0.0%	0.0	Resistant
18	GC60020-8-7-7-18	1.4	13.2%	0.2	7.1%	0.0	Resistant
46	BR01-17996	1.0	4.3%	0.0	0.0%	0.0	Resistant
41	FT2	1.3	37.8%	0.4	28.9%	0.1	Resistant
8	PI459025	4.5	36.7%	0.6	0.0%	0.3	Resistant
14	PI459025A	1.9	40.0%	0.6	0.0%	0.4	Resistant
7	PI462312	2.0	63.3%	0.9	57.1%	0.7	Resistant
30	Xiao Jing Huang	3.5	100.0%	1.9	36.7%	1.0	Slightly resistant
6	PI417125	1.2	100.0%	2.1	67.7%	1.6	Slightly resistant
57	BB20	4.0	96.0%	1.4	97.6%	1.8	Slightly resistant
4	PI587886	1.5	100.0%	1.9	69.0%	1.9	Slightly resistant
38	Da Li Zi	3.3	100.0%	2.0	48.3%	2.0	Slightly resistant
56	BB19	3.6	90.0%	2.1	95.2%	2.2	Susceptible
54	BB17	3.5	100.0%	1.5	100.0%	2.3	Slightly resistant
63	BB29	3.7	93.3%	1.6	81.6%	2.3	Slightly resistant
36	Lu Pi Dou	2.0	82.4%	2.9	81.5%	2.3	Susceptible
58	BB21	3.9	93.3%	2.3	91.4%	2.4	Susceptible
13	Wayne	2.9	100.0%	2.5	84.2%	2.4	Susceptible
62	BB28	3.8	100.0%	3.0	84.9%	2.4	Susceptible
20	GC84051-9-1	2.8	98.3%	2.3	90.7%	2.5	Susceptible
61	BB26	4.2	100.0%	2.4	83.6%	2.5	Susceptible
26	GC87012-10-B-5	3.4	100.0%	3.0	84.0%	2.5	Susceptible
59	BB22	4.1	100.0%	3.5	91.3%	2.6	Susceptible
60	BB25	4.0	96.7%	2.6	91.0%	2.6	Susceptible
40	Sachiyutaka	1.7	88.2%	2.9	95.9%	2.6	Susceptible
23	GC85039-1-2-1-1	2.5	96.7%	2.5	93.3%	2.7	Susceptible
47	BR01-18437	1.1	100.0%	2.4	85.6%	2.7	Susceptible
27	GC87016-11-B-2	2.1	100.0%	3.2	90.1%	2.7	Susceptible
24	GC86004-9	3.1	100.0%	2.1	75.0%	2.7	Susceptible
19	GC84040-16-1	2.5	94.8%	2.5	66.4%	2.7	Slightly resistant
28	SRE-B-15C	2.0	100.0%	2.9	91.1%	2.8	Susceptible
50	Misuzudaizu	2.9	100.0%	3.2	85.1%	2.8	Susceptible
53	BRS231	2.4	100.0%	4.2	84.9%	3.0	Susceptible
25	SS86045-23-2	3.2	100.0%	4.4	86.4%	3.0	Susceptible
37	Hei Dou	2.7	98.3%	3.9	80.8%	3.0	Susceptible
42	Davis	2.9	100.0%	4.0	90.2%	3.0	Susceptible
45	RI75	3.8	100.0%	3.9	92.1%	3.0	Susceptible
35	Himedaizu	3.9	93.3%	2.1	70.6%	3.0	Susceptible
5	PI230970	1.3	100.0%	2.2	84.8%	3.0	Susceptible
33	Da Bai Qi	4.7	100.0%	2.5	89.0%	3.0	Susceptible
32	Qin Dou	4.1	100.0%	3.2	91.6%	3.0	Susceptible
55	BB18	4.7	100.0%	3.2	90.7%	3.0	Susceptible
34	6611	4.5	100.0%	3.3	97.9%	3.0	Susceptible
31	Niu Mao Huang	3.2	100.0%	3.5	74.2%	3.0	Susceptible
49	EMBRAPA48	2.2	100.0%	3.7	90.3%	3.0	Susceptible
12	TK5	1.6	100.0%	4.0	78.3%	3.0	Susceptible
52	BRS184	2.4	100.0%	4.8	97.8%	3.0	Susceptible
51	Moshidou Gong 503	2.9	100.0%	5.1	87.0%	3.0	Susceptible

¹The values $0.0 \leq x < 70.0$ and $70.0 \leq x \leq 100.0$ are respectively classified as indicating resistance (shading) and susceptibility for %LU; ² The values $0.0 \leq x < 2.0$ and $2.0 \leq x$ are respectively classified as indicating resistance (shading) and susceptibility for NoU; ³ The values $0.0 \leq x < 70.0$ and $70.0 \leq x \leq 100.0$ are respectively classified as indicating resistance (shading) and susceptibility for %OU; ⁴ The values $0.0 \leq x < 2.0$ and $2.0 \leq x \leq 3.0$ are respectively classified as indicating resistance (shading) and susceptibility for SL; ⁵ Final classification of resistance was determined by the following criteria: "Immune": having no lesions; "Highly resistant": having lesions showing the resistant phenotype in four characters and with no uredinia; "Resistant": having lesions showing resistant phenotype in four characters and with uredinia; "Slightly resistant": having lesions showing resistant phenotype in any of four resistance characters; "Susceptible": having lesions showing susceptible phenotype in all four resistance characters. "Almost immune" means that the genotypes showed imperfect formation of lesions that might have been caused by very strong resistance (shown in Figure. 3).

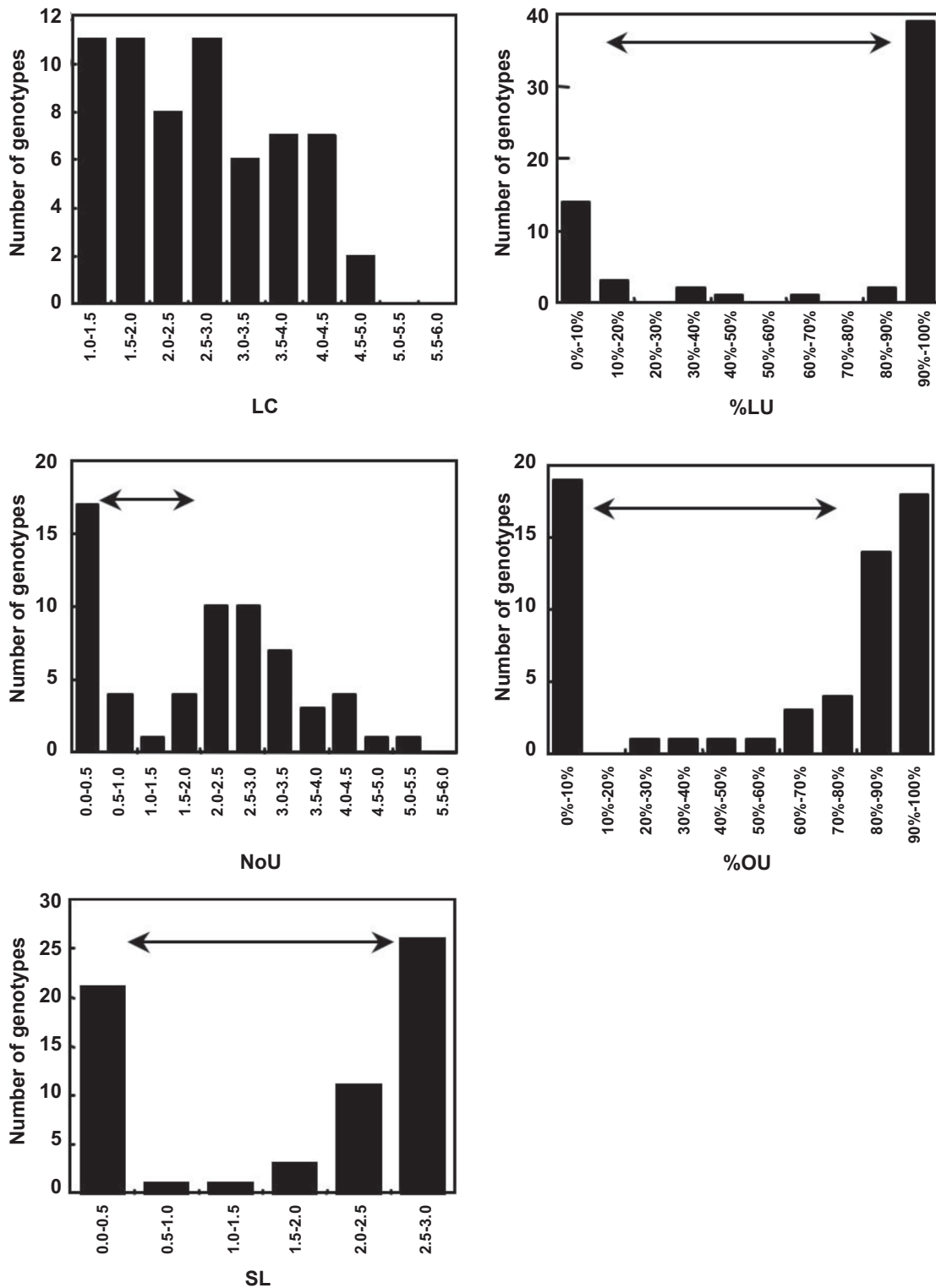


FIGURE 2 - Frequency distribution of the 5 characters for resistance against the Japanese rust population in the 63 genotypes. LC, %LU, NoU, %OU, and SL mean lesion color, frequency of lesion having uredinia, number of uredinia per lesion frequency of open uredinia, and sporulation level, respectively. Estimated boundary values between resistant and susceptible phenotypes for each character are shown by arrows.

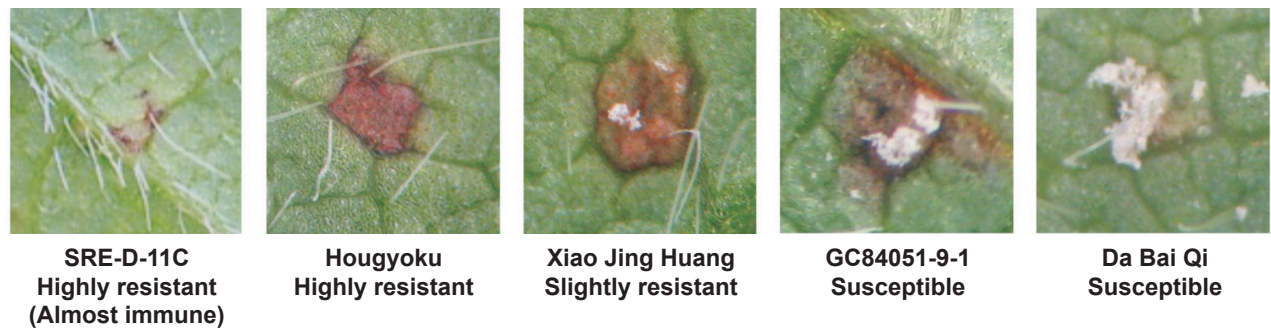


FIGURE 3 - Examples of the different pigmentation and sporulation of lesions observed in the 63 genotypes in response to inoculation with the Japanese rust population.

(0.412 to 0.515) with the other four resistance characters (Table 3). That is, most varieties had the same phenotypes—resistance or susceptible—in terms of the four resistance characters, with the exception of LC, and the darkness of LC was not always correlated with the production of uredinia and spores. For example, PI459025 (*Rpp4*) produced lesions that had a low NoU and SL but a light LC (with score of 4.5, Table 2). On the other hand, PI230970 (*Rpp2*) and TK5, which had abundant spore production, had dark LC (with scores of 1.3 and 1.6, respectively).

Classification criteria for resistance to soybean rust

The frequency distributions of four characters (excluding LC) in the 63 genotypes were bimodal (Figure 2), and two peaks that may have been associated with resistant and susceptible genotypes were observed. The thresholds between resistant and susceptible were determined between these two peaks (Figure 2) for the four resistance characters (excluding LC), as follows. The value of 2.0 for SL was primarily determined as the boundary value for distinguishing resistant (R) and susceptible (S). In this study, SL is considered to be the most important character for soybean rust resistance among the five because the amount of spores produced directly influences multiplication of soybean rust. An SL of less than 2.0 was clearly different from a larger (susceptible) SL. The boundary values between R and S for the other three characters were then determined to minimize the number of genotypes showing both R and S phenotypes in the four resistance characters. That is, the values $0.0 \leq x < 2.0$ and $2.0 \leq x$ were respectively classified as resistant and susceptible for NoU. The values $0.0 \leq x < 70.0$ and $70.0 \leq x \leq 100.0$ were respectively classified as resistant and susceptible for both %OU and %LU.

Finally, all 63 genotypes were classified into five kinds of resistance classification, “Immune”, “Highly resistant”, “Resistant”, “Slightly resistant”, and “Susceptible”, according to the classification criteria of “Immune” = having no lesions; “Highly resistant” = having lesions showing the resistant phenotype in all four characters and with no uredinia; “Resistant” = having lesions showing the resistant

phenotype in all four characters and possessing uredinia; “Slightly resistant” = having lesions that show resistant phenotype in any of four characters; and “Susceptible” = having lesions with susceptible phenotypes in all four resistance characters. These five resistance categories of genotypes were also applied to the other two Brazilian rust populations in order to identify the difference in the virulence among three populations.

Pathogenic differences among the 3 rust populations

We examined the pathogenic differences among JRP, BRP-1, and BRP-2 in the 13 standard varieties based on the criteria we determined (Figure 4). With BRP-1 infection, a few whitish lesions in 100 dark brown lesions were observed in the genotype, PI416764. With BRP-2 infection, two clearly different types of lesions with similar frequencies were also observed on PI587905. Such mixed lesions were not observed in JRP.

Some pathogenic differences were observed between BRP-1 and BRP-2. Major differences in virulence were observed in PI230970 (*Rpp2*), PI417125 (*Rpp2*), PI462312 (*Rpp3*), and PI416764 (Figure 4). Until 2008, it had not been reported that Brazilian rust populations show virulence against genotypes containing the *Rpp2* gene. In addition, stronger

TABLE 3 - Pearson correlations among the five characters for resistance against the Japanese rust population in the 63 genotypes. All correlations are significant ($P < 0.001$) by t-test. For key to table headings see caption to Table 2

	LC	%LU	NoU	%OU	SL
LC	–	0.515	0.412	0.502	0.499
%LU	–	–	0.873	0.951	0.944
NoU	–	–	–	0.868	0.926
%OU	–	–	–	–	0.953
SL	–	–	–	–	–

Standard varieties	<i>Rpp1</i>				<i>Rpp2</i>		<i>Rpp3</i>	<i>Rpp4</i>	<i>Rpp5</i>	(unknown)			
	1. PI200492	2. Tainung 4	3. PI587880A	4. PI587886	5. PI230970	6. PI417125	7. PI462312	8. PI459025	9. Shiranui	10. PI416764	11. PI587905	12. TK5	13. Wayne
Japanese rust population (JRP)	HR	HR	I	SR	S	SR	HR	R	HR	HR	HR	S	S
Brazilian rust Population 1 (BRP-1)	S	S	S	S	R	R	S	R	R	(Mix)	S	S	S
Brazilian rust Population 2 (BRP-2)	S	S	S	S	S	S	SR	SR	SR	SR	(Mix)	S	S

I :Immunity **HR** :Highly resistant **R** :Resistant **SR** :Slightly resistant **S** :Susceptible **(Mix)** :2 types of lesions mixed

FIGURE 4 - Phenotypes of resistance against Japanese and Brazilian rust populations in 13 standard varieties. 1-9: Varieties having one of known resistance genes, 10 and 11: Varieties having identified their resistance but not identified their genes yet, 12 and 13: Susceptible standard varieties.

virulence of BRP-2 than BRP-1 was observed in PI459025 with the *Rpp4* gene. Thus, the resistance conferred by *Rpp4* from PI459025 against Brazilian populations might have started to break down. Clear differences in virulence between the Japanese and the two Brazilian populations were observed in all four varieties that have the *Rpp1* gene: PI200492, Tainung 4, PI587880A, and PI587886. All the differences in these varieties involved the Brazilian populations causing stronger disease than the Japanese one.

DISCUSSION

Various types of lesions obtained from JRP infection

We considered that following factors were likely to have been associated with various types of lesions observed in JRP infection. First, not only major resistance genes but also minor resistance genes may have influenced these resistance characters. A continuous degree of resistance has previously been observed among soybean lines that have identical compositions of the major resistance genes but have different genetic backgrounds (Nogueira et al., 2008; Yamanaka et al., 2008). In this study, PI417125, which has the major resistance gene, *Rpp2*, did not show as clearly resistant but were seen to be slightly resistant (Table 2). This is also an example of the major resistance genes showing their resistance quantitatively or soybean varieties having quantitative trait loci for resistance characters except for major genes. Second, the mixture of races in JRP might have produced continuous variation in the average values of the characters in the 30 lesions sampled per genotype. Mixed types of lesions, which can be clearly distinguished from each other, were not observed in JRP infection. However, it is difficult to differentiate types of pathogenic races without clear differences in their LCs or SLs. Thus, the average values of the resistance characters could have been derived

from different pathogenic races. Consequently, the use of both lines that are isogenic for each major resistance gene and single-lesion isolates taken from rust populations will be necessary to determine the boundaries between resistant and susceptible types more exactly.

Classification criteria for resistance to soybean rust

The resistance genes *Rpp2* and *Rpp4* contribute genetically to the darkness of LC against BRP-1 under greenhouse conditions (Yamanaka et al., 2008). PI459025 (*Rpp4*) had an LC score that differed with the growth environment, namely 4.5 in this study and 1.2 under greenhouse conditions (data not shown). This suggests that LC is easily influenced by environmental factors. We consider spore production to be more important than LC in the identification of resistance against soybean rust. LC showed a low correlation with the other four resistance characters, which appear to be more important for selection in the breeding program (Table 3). This, combined with apparent variation with environmental factors mentioned above and the fact that it shows not bimodal but continuous distribution (Figure 2), makes it unlikely to be a suitable criterion for resistance classification of genotypes. However, a clear difference in LC is considered to be useful for detecting mixed lesions derived from different pathogenic races.

The difference in the virulence among three rust populations was successfully revealed by the classification criteria for resistance which we developed. Therefore, it can be used for identifying the genes/varieties effective for the other rust populations. However, the evaluation methodology and the classification criteria mentioned in this paper are not universal in all phases of the breeding program. For example, DNA markers or visual classification of RB/TAN may be useful when we select the lines from the population where the major resistance genes segregate,

and Severity check may also be useful for introducing the minor genes and for evaluating effectiveness of fungicide application (Godoy et al., 2009). In addition, we observed high correlation among 4 characters related to resistance. This observation may enable us to simplify the process of evaluation. The criteria to decide resistance against soybean rust were determined by large number of phenotypes: four characters in 63 genotypes. However, phenotypic data of genotypes shown in this study are based on a one-time experiment, and thus we have to treat the data carefully by considering that.

Pathogenic differences among the 3 rust populations

The two Brazilian rust populations previously shown to have similar virulence in a different set of differential varieties in 2005 (Kato & Yorinori, 2008), showed some difference in their virulence in this study. Therefore, the differences observed here are likely to have resulted from a change in virulence, large changes in the race population, or both, which occurred from 2005 to 2008 in the two greenhouses. Phenotypes resistant to BRP-1 or BRP-2 were observed in varieties with the genes *Rpp2*, *Rpp3*, *Rpp4*, and *Rpp5* (entries 5 to 9) and in PI416764, but only two of these varieties, PI459025 (*Rpp4*) and cv. Shiranui (*Rpp5*), were resistant to both Brazilian populations (Figure 4). In other words, only two of the five major genes can be expected to be useful in resistance to Brazilian soybean rust. In addition, Brazilian populations caused more virulent reaction than Japanese populations in 13 standard varieties. With the infections of two Brazilian populations, two clearly different types of lesions were observed in the genotypes, PI416764 and PI587905, respectively for BRP-1 and BRP-2, in 13 standard varieties. However, such mixed lesions were not observed in JRP infection using 63 genotypes. This fact indicated that Brazilian rust populations may have not only higher but also more diverse virulence than the Japanese one. The most significant difference between the Japanese and Brazilian populations was observed in the 4 genotypes having *Rpp1* gene in this study. Pham et al. (2009) detected a susceptible reaction of *Rpp1* (PI200492) in the Brazilian isolate, but immunity and resistant reactions in the Indian and Louisiana isolates, respectively. Li (2009) also detected high resistance in PI200492 against 3 isolates from Mississippi. Thus, the resistant allele of *Rpp1* from PI200492 can be useful in some regions of Asia and the USA but not in Brazil.

Our results suggested that the resistant varieties or resistance genes useful in Brazil were limited. They also suggested that a breakdown of major resistance genes or large changes in race populations, as observed in the case of the *Rpp2* gene (from PI230970 and PI417125), are occurring. Therefore, a resistant cultivar that is universally effective against soybean rust should be developed by pyramiding many major resistance genes and by introducing horizontal resistance.

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