

Inheritance of Resistance to Bacterial Blight in 21 Cultivars of Rice

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ABSTRACT

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Genetic analysis for resistance to bacterial blight (*Xanthomonas oryzae* pv. *oryzae*) of 21 rice (*Oryza sativa* L.) cultivars was carried out. These cultivars were divided into two groups based on their reactions to Philippine races of bacterial blight. Cultivars of group 1 were resistant to race 1 and those of group 2 were susceptible to race 1 but resistant to race 2. All the cultivars were crossed with TN1, which is susceptible to all the Philippine races of *X. oryzae* pv. *oryzae*. F₁ and F₂ populations of hybrids of group 1 cultivars were evaluated using race 1 and F₁ and F₂ populations of hybrids of group 2 cultivars were evaluated using race 2. All the cultivars showed monogenic inheritance of resistance. Allelic re-

lationships of the genes were investigated by crossing these cultivars with different testers having single genes for resistance. Three cultivars have *Xa4*, another three have *xa5*, one has *xa8*, two have *Xa3*, eight have *Xa10*, and one has *Xa4* as well as *Xa10*. Three cultivars have new, as yet undescribed, genes. Nep Bha Bong To has a new recessive gene for moderate resistance to races 1, 2, and 3 and resistance to race 5. This gene is designated *xa26(t)*. Arai Raj has a dominant gene for resistance to race 2 which segregates independently of *Xa10*. This gene is designated as *Xa27(t)*. Lota Sail has a recessive gene for resistance to race 2 which segregates independently of *Xa10*. This gene is designated as *xa28(t)*.

Additional keywords: durable resistance, genetic engineering, independent segregation, isogenic lines, marker aided selection, molecular markers, pyramided lines.

Bacterial blight, caused by *Xanthomonas oryzae* pv. *oryzae*, is one of the most serious diseases of rice. It is widespread throughout Asia and also has been reported to occur in Australia, the United States, and several rice-growing countries of Latin America and Africa (16). The disease may cause damage at seedling stage resulting in complete wilting or death of affected tillers. The infection at maximum tillering stage results in blighting of leaves. The yield losses in severely infected fields generally range from 20 to 30% but may reach up to 80% (24,28).

Chemical control of bacterial blight in the monsoon climate of Asia is impractical. Additionally, no effective bactericide is commercially available for disease control. Therefore, the preferred strategy for disease management is through varietal resistance. Large germ plasm collections have been screened to identify sources of resistance for breeding programs and many resistant cultivars have been genetically analyzed to identify diverse genes for resistance. To date, 22 genes for resistance have been identified (2,3,11,12,15,23,25–27,34). Two of the genes, *Xa1* and *Xa21*, have been cloned (30,33). Several of these genes have been incorporated into improved rice cultivars which are now widely grown (14). However, new races of the pathogen continue to evolve that can overcome the resistance conveyed by the major genes (16,19).

Mew and Vera Cruz (17) reported four races of bacterial blight in the Philippines and Vera Cruz and Mew (32) reported six races. Two additional races were described by Nelson et al. (21). Some of these races may have originated as a result of deployment of cultivars with major genes. Others may have existed but were identified when additional differentials were employed. In order to maintain a dynamic breeding program for bacterial blight resis-

tance for such a variable pathogen, a constant supply of new genes is needed. This study was undertaken to identify additional genes for resistance to bacterial blight.

MATERIALS AND METHODS

Twenty-one rice cultivars were analyzed (Table 1). These cultivars were divided into two groups based on their reaction to *X. oryzae* pv. *oryzae*. Cultivars of group 1 are resistant to Philippine race 1 and a few other races. Cultivars of group 2 are susceptible to race 1 but resistant to races 2 and 5. This division is important because two genes, *Xa4* and *xa5*, which are commonly found in resistant cultivars, convey resistance to races 1 and 5. On the other hand, cultivars susceptible to race 1 but resistant to races 2 and 5 have *Xa10* for resistance. Seven testers were employed for genetic analysis (Table 2). These lines included six isogenic lines with single genes for resistance and TN1, which is susceptible to all Philippine races of *X. oryzae* pv. *oryzae*.

All the cultivars were crossed with TN1. F₁ and F₂ populations from the crosses of group 1 cultivars were inoculated with race 1, whereas the F₁ and F₂ populations from the crosses of group 2 cultivars were inoculated with race 2. The gene for resistance of each cultivar was predicted on the basis of their reaction pattern to six Philippine races of bacterial blight. Thereafter, the test cultivars were crossed with various testers having the known genes for resistance to verify our predictions.

The inoculation method of Kauffman et al. (9) was employed for inoculation of hybrid progenies. Pure cultures of the bacterial strains were maintained in the bacterial blight laboratory at the Plant Breeding, Genetics, and Biochemistry Division, International Rice Research Institute (IRRI). The inoculum was prepared and diluted by mixing bacterial cultures in distilled water. Using a spectrophotometer, the absorbance (*A*) of the inoculum was adjusted to *A* = 0.05 (620 nm). This value corresponds to a concentration of $\approx 10^8$ cells per ml. Plants were inoculated at

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maximum tillering (40 days after transplanting) or booting stage (55 to 60 days after transplanting) and were scored for disease reaction 14 days after inoculation. Both visual assessment and lesion length measurements of three inoculated leaves of at least three plants of each F₁ population were carried out. Each plant of F₂ populations was classified visually as either resistant (lesion length <10 cm) or susceptible (lesion length >10 cm).

RESULTS

Inheritance of resistance. The reactions to race 1 of F₁ and F₂ populations of crosses of group 1 cultivars with TN1 are given in Table 3. The F₁ hybrids of cvs. Bazail 975, Karia, Minh-Soc, Nep Bha Bong To, and Latu were susceptible and their F₂ populations segregated in a ratio of one resistant to three susceptible, indicating monogenic recessive control of resistance in these cultivars. The F₁ hybrids of cvs. WC1263, HR 12, SLO 2, Ragu Sail, Trang Chum, and Ba Kieu were resistant and their F₂ populations segregated in a ratio of three resistant to one susceptible, indicating that each of them have a dominant gene for resistance to race 1.

The reactions to race 2 of F₁ and F₂ populations of crosses of group 2 cultivars with TN1 are given in Table 4. The F₁ hybrids of all cultivars except Lota Sail were resistant and F₂ populations segregated in a ratio of three resistant to one susceptible, indicating monogenic dominant control of resistance to race 2 in these cultivars. The F₁ hybrid of Lota Sail with TN1 was susceptible and the F₂ population segregated in a ratio of one resistant to three susceptible, indicating Lota Sail has a recessive gene for resistance to race 2.

SLO 2 also showed resistance to race 2; therefore, the F₁ and F₂ populations of its cross with TN1 also were evaluated for resistance to race 2 (Table 4). The F₁ hybrid was resistant, and the F₂ population segregated in a ratio of three resistant to one susceptible, indicating that it has a dominant gene for resistance to race 2.

Allele tests. WC1263, HR 12, and Ragu Sail have the reaction pattern to six races typical of rice cultivars with *Xa4*; therefore, these cultivars were crossed with IRBB4. SLO 2 also was crossed with IRBB4 to determine if its resistance to race 1 may also be due to *Xa4*. The F₁ hybrids of these four crosses were resistant to

race 1, and the F₂ population did not segregate for susceptibility, showing that resistance to race 1 in these cultivars is due to *Xa4* (Table 5).

The reaction pattern of Bazail 975, Karia, and Latu to six races was similar to that of cultivars with *xa5* (i.e., resistant to races 1, 2, 3, and 5 and susceptible to races 4 and 6). Therefore, these cultivars were crossed with IRBB5. The F₁ hybrids were resistant to race 1 and F₂ populations did not segregate for susceptibility (Table 5), showing that these three cultivars have *xa5* for resistance.

Trang Chum and Ba Kieu were found to have a dominant gene for resistance to race 1, and their reaction pattern to six races was similar to that of cultivars with *Xa3* (i.e., moderately resistant to races 1, 2, and 3, resistant to race 5, and susceptible to races 4 and 6); therefore, they were crossed with IRBB3. The F₁ hybrids were resistant to race 1 and F₂ populations did not segregate for susceptibility. These results show that Trang Chum and Ba Kieu have *Xa3* for resistance (Table 5).

Minh-Soc and Nep Bha Bong To were found to have a recessive gene for resistance to race 1 and their reaction pattern was similar to that of *xa8* to six races; therefore, they were crossed with IRBB8. Nep Bha Bong To showed segregation with IRBB8, so it also was crossed with IRBB5. The F₁ hybrid of Minh-Soc with IRBB8 was resistant and the F₂ population did not segregate for susceptibility (Table 5). These results indicate that Minh-Soc has *xa8* for resistance. The F₁ hybrids of Nep Bha Bong To with IRBB5 and IRBB8 were susceptible and the F₂ populations

TABLE 2. Reactions of various testers to the six Philippine races of *Xanthomonas oryzae* pv. *oryzae*

Tester	Gene	Reaction to race ^a					
		1	2	3	4	5	6
TN1	...	S	S	S	S	S	S
IRBB3	<i>Xa3</i>	MR	MR	MR	S	R	S
IRBB4	<i>Xa4</i>	R	S	S	S	R	S
IRBB5	<i>xa5</i>	R	R	R	S	R	S
IRBB7	<i>Xa7</i>	MR	R	R	S	R	S
IRBB8	<i>xa8</i>	MR	MR	MR	S	R	S
IRBB10	<i>Xa10</i>	S	R	S	S	R	S

^a R = resistant, MR = moderately resistant, and S = susceptible.

TABLE 1. Rice cultivars used in the study and their reactions to the six Philippine races of *Xanthomonas oryzae* pv. *oryzae*

Cultivar	Accession no. ^b	Origin	Reaction to <i>X. oryzae</i> pv. <i>oryzae</i> races ^a					
			1	2	3	4	5	6
Group 1								
WC1263	11057	India	R	S	S	S	R	S
HR 12	674	India	R	S	S	S	R	S
SLO 2	678	India	R	R	S	S	R	S
Ragu Sail	26774	Bangladesh	R	S	S	S	R	S
Bazail 975	32816	Bangladesh	R	R	R	S	R	S
Karia	6702	Fiji	R	R	R	S	R	S
Latu	18093	Indonesia	R	R	R	S	R	S
Minh-Soc	16731	Vietnam	MR	MR	MR	S	R	S
Trang Chum	16778	Vietnam	MR	MR	MR	S	R	S
Ba Kieu	16855	Vietnam	MR	MR	MR	S	R	S
Nep Bha Bong To	17005	Vietnam	MR	MR	MR	S	R	S
Group 2								
Aghanisail	25827	Bangladesh	S	R	S	S	R	S
Arai Raj	26536	Bangladesh	S	R	S	S	R	S
Huson Dhaw	26644	Bangladesh	S	R	S	S	R	S
Khar Mao	26687	Bangladesh	S	R	S	S	R	S
Lota Sail	26719	Bangladesh	S	R	S	S	R	S
Pelatie	30598	Sierra Leone	S	R	S	S	R	S
Ganga Sagar	31650	Bangladesh	S	R	S	S	R	S
Bhua Balam	31729	Bangladesh	S	R	S	S	R	S
Kanchanchari	31837	Bangladesh	S	R	S	S	R	S
Surma Sail	31941	Bangladesh	S	R	S	S	R	S

^a R = resistant, MR = moderately resistant, and S = susceptible.

^b International Rice Research Institute accession numbers.

segregated in a ratio of seven resistant to nine susceptible (Table 5). These results show that the recessive gene for resistance to race 1 of Nep Bha Bong To is nonallelic to and independent of *xa5* and *xa8*.

Eleven cultivars of group 2 showed a reaction pattern to six races that is similar to that of IRBB10 (i.e., resistance to races 2 and 5 but susceptible to races 1, 3, 4, and 6). These 11 cultivars were crossed with IRBB10. The F₂ population of Arai Raj from its cross with IRBB10 segregated for susceptibility; therefore, it also was crossed with IRBB7. Genetic analysis indicated that Lota Sail has a recessive gene for resistance to race 2, so it also was crossed with IRBB5 and IRBB8. The F₁ hybrids of SLO 2, Aghanisail, Huson Dhaw, Pelatie, Bhua Balam, Kanchanchari, Surma Sail,

Khar Mao, and Ganga Sagar with IRBB10 were resistant and F₂ populations did not segregate for susceptibility (Table 6). These results show that these nine cultivars have *Xa10* for resistance to race 2.

The F₂ populations of Arai Raj with IRBB10 and IRBB7 segregated in a ratio of 15 resistant to 1 susceptible, indicating that the dominant gene of Arai Raj for resistance to race 2 is different from and independent of *Xa10* and *Xa7*. The F₂ populations of Lota Sail from crosses with IRBB10, IRBB5, and IRBB8 segregated in resistant:susceptible ratios of 13:3, 7:9, and 7:9, respectively. These data indicate that the recessive gene of Lota Sail for resistance to race 2 is different from and independent of *Xa10* as well as *xa5* and *xa8*.

TABLE 3. Reaction of F₁ hybrids and F₂ populations to race 1 of *Xanthomonas oryzae* pv. *oryzae* from crosses of group 1 cultivars with susceptible cv. TN1

Cross	F ₁ plants	Reaction to race 1 ^a				
		F ₂ population			χ^2	P value
		R	S	Ratio		
TN1 × WC1263	R	284	92	3:1	0.06	0.75–0.90
TN1 × HR 12	R	298	110	3:1	0.84	0.25–0.50
TN1 × SLO 2	R	274	87	3:1	0.16	0.50–0.75
TN1 × Ragu Sail	R	311	116	3:1	1.07	0.25–0.50
TN1 × Bazail 975	S	123	351	1:3	0.23	0.50–0.75
TN1 × Karia	S	91	228	1:3	2.11	0.10–0.25
TN1 × Latu	S	139	362	1:3	2.01	0.10–0.25
TN1 × Minh-Soc	S	145	427	1:3	0.04	0.75–0.90
TN1 × Trang Chum	R	127	41	3:1	0.03	0.75–0.90
TN1 × Ba Kieu	R	245	67	3:1	2.07	0.10–0.25
TN1 × Nep Bha Bong To	S	96	347	1:3	2.62	0.05–0.10

^a R = resistant and S = susceptible.

TABLE 4. Reaction of F₁ hybrids and F₂ populations to race 2 of *Xanthomonas oryzae* pv. *oryzae* from crosses of group 2 cultivars with susceptible cv. TN1

Cross	F ₁ plants	Reaction to race 2 ^a				
		F ₂ population			χ^2	P value
		R	S	Ratio		
TN1 × SLO 2	R	137	47	3:1	0.03	0.80–0.90
TN1 × Aghanisail	R	117	34	3:1	0.50	0.25–0.50
TN1 × Huson Dhaw	R	384	148	3:1	2.25	0.10–0.25
TN1 × Pelatie	R	285	87	3:1	0.52	0.25–0.50
TN1 × Bhua Balam	R	315	104	3:1	0.007	0.90–0.95
TN1 × Kanchanchari	R	437	157	3:1	0.65	0.25–0.50
TN1 × Surma Sail	R	265	86	3:1	0.05	0.75–0.90
TN1 × Khar Mao	R	147	54	3:1	0.37	0.50–0.75
TN1 × Ganga Sagar	R	321	109	3:1	0.03	0.80–0.90
TN1 × Arai Raj	R	227	57	3:1	3.68	0.05–0.10
TN1 × Lota Sail	S	75	280	1:3	2.84	0.05–0.10

^a R = resistant and S = susceptible.

TABLE 5. Reaction of F₁ hybrids and F₂ populations to race 1 of *Xanthomonas oryzae* pv. *oryzae* from the crosses of group 1 cultivars with different testers

Cross	F ₁ plants	Reaction to race 1 ^a				
		F ₂ population			χ^2	P value
		R	S	Ratio		
IRBB4 × WC1263	R	504	0	1:0	0.0	1.00
IRBB4 × HR 12	R	479	0	1:0	0.0	1.00
IRBB4 × SLO 2	R	493	0	1:0	0.0	1.00
IRBB4 × Ragu Sail	R	487	0	1:0	0.0	1.00
IRBB5 × Bazail 975	R	485	0	1:0	0.0	1.00
IRBB5 × Karia	R	472	0	1:0	0.0	1.00
IRBB5 × Latu	R	314	0	1:0	0.0	1.00
IRBB8 × Minh-Soc	R	477	0	1:0	0.0	1.00
IRBB3 × Trang Chum	R	465	0	1:0	0.0	1.00
IRBB3 × Ba Kieu	R	454	0	1:0	0.0	1.00
IRBB5 × Nep Bha Bong To	S	193	277	7:9	1.38	0.10–0.25
IRBB8 × Nep Bha Bong To	S	180	295	7:9	6.62	<0.05

^a R = resistant and S = susceptible.

Designation of new genes. Of the 11 cultivars of group 1, 4 (i.e., WC1263, HR 12, SLO 2, and Ragu Sail) were each found to have single dominant genes for resistance that are allelic to *Xa4*. Bazail 975, Karia, and Latu each have single recessive genes for resistance which are allelic to *xa5*. Minh-Soc has a single recessive gene for resistance which is allelic to *xa8*. The recessive gene *xa8* was first identified in rice cultivar PI231129 (27). Since then, we have genetically analyzed several hundred cultivars for resistance to bacterial blight and Minh-Soc is the only other cultivar found to have *xa8*. Trang Chum and Ba Kieu each have single dominant genes for resistance that are allelic to *Xa3*. The cultivar Nep Bha Bong To has a recessive gene for resistance that is nonallelic to two other recessive genes, *xa5* and *xa8*. It segregated independently of *xa5* but may be linked to *xa8* because the χ^2 value for independent segregation is quite high. Therefore, the recessive gene for resistance of Nep Bha Bong To appears to be a new gene and, following the rules of gene nomenclature in rice, this gene is designated as *xa26(t)*.

Eight cultivars of group 2 (i.e., Aghanisail, Huson Dhaw, Pelatie, Bhua Balam, Kanchanchari, Surma Sail, Khar Mao, and Ganga Sagar) each have a single dominant gene allelic to *Xa10*. In addition, SLO 2 also has *Xa10*. Although cv. Arai Raj has the reaction pattern to six races that is similar to that of *Xa10*, it turned out to have a single dominant gene for resistance which is nonallelic to and independent of *Xa10* as well as *Xa7*. This dominant gene for resistance to races 2 and 5 is designated as *Xa27(t)*.

Similarly, cv. Lota Sail has a reaction pattern to six races that is also similar to that of *Xa10*. However, it has a single recessive gene for resistance to races 2 and 5 which segregates independently of *xa5*, *xa8*, and *Xa10*. This gene is designated as *xa28(t)*.

The genetic analysis of 21 cultivars has resulted in identification of three new genes for resistance to bacterial blight. The genes for resistance of these cultivars are summarized in Table 7.

DISCUSSION

Insufficient durability of resistance to bacterial blight is a major concern. Durable resistance is "resistance that remains effective while a cultivar possessing it is widely cultivated" (8). Introduction of resistant cultivars often has resulted in shifts of virulence in populations of *X. oryzae* *pv.* *oryzae* (16). Many cultivars with *Xa4* for resistance to bacterial blight have been cultivated widely in the Philippines and elsewhere in Asia. The frequency of pathotypes virulent to *Xa4* increased over time and became dominant in the Philippine populations of the pathogen in response to the use of *Xa4* (18).

One strategy to prolong the useful life of major gene resistance is to pyramid two or more major genes in a single cultivar (10).

This strategy would be desirable if, through various mechanisms (20), it would be more difficult for a pest to overcome several resistance genes in the host than to overcome a single resistance gene.

It is difficult to pyramid resistance genes by phenotypic selection because the presence of one major gene obscures the effects of other genes. Molecular markers that are closely linked to major genes can be used to identify individuals carrying more than one resistance gene in a segregating population. Fortunately, a molecular genetic map of rice densely populated with various types of molecular markers has been prepared (4) and many genes for disease and insect resistance have been tagged with molecular markers (13). Another advantageous situation in regard to bacterial blight is the availability of near-isogenic lines with single genes for resistance (22). DNA marker-aided selection was used to pyramid four bacterial blight resistance genes, *Xa4*, *xa5*, *xa13*, and *Xa21*. Breeding lines with two, three, and four resistance genes were developed and tested for resistance to bacterial blight. The pyramided lines showed a wider spectrum and a higher level of resistance than lines with only a single gene (5). Pyramided lines were shared with national rice improvement programs of several countries in Asia, where they are being used in local breeding programs (29).

The newly discovered genes are being transferred to near-isogenic background and, at the same time, are being tagged with

TABLE 7. Genes for bacterial blight resistance in the rice cultivars analyzed

Cultivar	Gene for resistance
WC1263	<i>Xa4</i>
HR 12	<i>Xa4</i>
SLO 2	<i>Xa4</i> + <i>Xa10</i>
Ragu Sail	<i>Xa4</i>
Bazail 975	<i>xa5</i>
Karia	<i>xa5</i>
Latu	<i>xa5</i>
Minh-Soc	<i>xa8</i>
Trang Chum	<i>Xa3</i>
Ba Kieu	<i>Xa3</i>
Nep Bha Bong To	<i>xa26(t)</i>
Aghanisail	<i>Xa10</i>
Huson Dhaw	<i>Xa10</i>
Pelatie	<i>Xa10</i>
Bhua Balam	<i>Xa10</i>
Kanchanchari	<i>Xa10</i>
Khar Mao	<i>Xa10</i>
Surma Sail	<i>Xa10</i>
Ganga Sagar	<i>Xa10</i>
Arai Raj	<i>Xa27(t)</i>
Lota Sail	<i>xa28(t)</i>

TABLE 6. Reaction of F₁ hybrids and F₂ populations to race 2 from the crosses of group 2 cultivars with different testers

Cross	F ₁ plants	Reaction to race 2 ^a					
		F ₂ population					P value
		R	S	Ratio	χ^2		
IRBB10 × SLO 2	R	473	0	1:0	0.0	1.00	
IRBB10 × Aghanisail	R	490	0	1:0	0.0	1.00	
IRBB10 × Huson Dhaw	R	377	0	1:0	0.0	1.00	
IRBB10 × Pelatie	R	500	0	1:0	0.0	1.00	
IRBB10 × Bhua Balam	R	494	0	1:0	0.0	1.00	
IRBB10 × Kanchanchari	R	540	0	1:0	0.0	1.00	
IRBB10 × Surma Sail	R	493	0	1:0	0.0	1.00	
IRBB10 × Khar Mao	R	477	0	1:0	0.0	1.00	
IRBB10 × Ganga Sagar	R	479	0	1:0	0.0	1.00	
IRBB10 × Arai Raj	R	437	36	15:1	1.50	0.10–0.25	
IRBB7 × Arai Raj	R	429	20	15:1	2.47	0.10–0.25	
IRBB10 × Lota Sail	R	348	99	13:3	3:39	0.05–0.10	
IRBB5 × Lota Sail	S	179	213	7:9	0.58	0.25–0.50	
IRBB8 × Lota Sail	S	119	170	7:9	0.78	0.25–0.50	

^a R = resistant and S = susceptible.

molecular markers. Thus, these genes can be employed in rice improvement programs either through conventional breeding approaches or through molecular marker-aided selection or gene pyramiding.

Genetic engineering also allows pyramiding of resistance genes. Resistance genes, if cloned, can be introduced into the rice genome in combination with existing resistance genes or with other novel genes.

As an example, *Xa21* was cloned from a near-isogenic rice line, IRBB21, by Song et al. (30). It was introduced into IR72, which has *Xa4* for resistance (31). Thus, transgenic IR72 has *Xa4* and *Xa21* pyramided together. This pyramided line has been evaluated under field conditions in China and showed a broad spectrum of resistance.

To date, 49,752 entries from the germ plasm collection maintained at IRRI have been evaluated for resistance to bacterial blight and 11.1% were found to be resistant (6). Approximately 700 have been genetically analyzed and this has resulted in the identification of 21 genes for resistance. Three genes (*xa19*, *xa20*, and *Xa25*) have been induced through mutagenesis. Two dominant genes, *Xa21* and *Xa23*, were transferred to cultivated germ plasm from closely related wild species *Oryza longistaminata* and *O. rufipogon*, respectively. Most of the distantly related wild species are resistant to bacterial blight. Crosses between these wild species and cultivated rice have been accomplished through embryo rescue technique (7) and genes for bacterial blight resistance have been transferred to cultivated rice (1).

Two bacterial blight resistance genes have been cloned. *Xa21* was isolated by map-based cloning and found to be member of a complex locus located on chromosome 11 (30). As mentioned earlier, this gene has been transferred back to rice through genetic engineering. *Xa1*, the second resistance gene cloned from rice, is located on chromosome 4 (33). Unlike *Xa21*, *Xa1* is a single-copy gene and is pathogen and wound inducible.

The concept of tissue-specific expression of major genes could be applied not only to genes isolated from other organisms but also to the manipulations of natural resistance genes such as *Xa1* and *Xa21*. When cloned major genes are engineered for tissue-specific expression, there will be less selection pressure on the pathogen exerted by the major genes. If the major genes for bacterial resistance could be engineered for expression in the leaves from booting stage until 10 to 15 days after flowering, their expression probably could protect the crop from yield losses and would be more durable.

As discussed above, various genetic strategies for reducing the yield losses from bacterial blight include identification of genes for resistance from the primary gene pool, transfer of genes for resistance from the closely as well as distantly related germ plasm, mapping and tagging of genes with molecular markers, transfer of genes to the elite genetic background through conventional and molecular marker-aided selection, pyramiding of genes through molecular marker-aided selection, fine mapping and cloning of genes for resistance, introduction of cloned genes through transformation into elite cultivars under tissue-specific expression, and pyramiding the cloned genes with naturally occurring genes. The identification of new resistance genes reported in this study is thus a basic step in the galaxy of approaches to minimize the losses caused by *X. oryzae* pv. *oryzae* in rice.

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