A Codominant Randomly Amplified Polymorphic DNA (RAPD) Marker Useful for Indirect Selection of Bean Golden Mosaic Virus Resistance in Common Bean

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Abstract. Bean golden mosaic virus (BGMV) is a devastating disease of common bean (*Phaseolus vulgaris* L.) in tropical America. The disease is effectively controlled by combinations of genetic resistances. The most widely deployed source of resistance to BGMV is a recessive gene (*bgm-1*) derived from the dry bean landrace cultivar Garrapato (Mexico) that conditions a nonmosaic partial resistance response to the pathogen. To expedite introgression of partial resistance into snap bean for southern Florida and other susceptible dry bean market classes for the Caribbean and Central American regions, a RAPD marker tightly linked to *bgm-1* has been identified. Two contrasting DNA bulks, one consisting of five BGMV-resistant and the other five susceptible F_6 recombinant inbred lines, were used to screen for polymorphic fragments amplified by 300 decamer primers in the polymerase chain reaction. RAPDs generated between the bulks were analyzed across F_2 populations segregating for the marker and the gene. One codominant RAPD marker ($R2_{570/530}$) tightly linked to the recessive resistance gene *bgm-1* was found. The 530-base pair (bp) fragment was linked in repulsion with *bgm-1* and the other 570-bp fragment was linked in coupling. No recombinants between $R2_{570/530}$ and *bgm-1* were observed among 91 F_2 progeny from one dry bean population, and there were two recombinants (4.2 cM) observed among 48 F_2 progeny combined across four snap bean populations. Assays of $R2_{570/530}$ across susceptible germplasm and lines likely to have the 'Garrapato'-derived partial resistance to BGMV have revealed that the codominant marker is gene-pool nonspecific and maintains its original linkage orientation with the recessive *bgm-1* gene through numerous meioses. The codominant marker is useful for rapidly introgressing partial resistance to BGMV into susceptible germplasm.

Bean golden mosaic virus (BGMV) is the most serious viral disease of common bean (*Phaseolus vulgaris* L.) in Latin America (Gálvez and Morales, 1989). Bean golden mosaic is caused by a geminivirus (Maxwell et al., 1994) vectored by the whiteflies *Bemisia tabaci* (Bellows and Perring) and *B. argentifolii* (Genn.) (Gálvez and Morales, 1989). BGMV has been reported in southern Florida where it has devastated snap bean production (Blair et al., 1995).

Genetic resistance to BGMV has proven to be an effective form of disease control (Beebe, 1994; Morales and Singh, 1991). The first BGMV resistant cultivars ('Jutiapan', 'Ostua', and 'Quetzal', etc.) have been developed by International Center for Tropical Agriculture (CIAT) and Agricultural Institute of Science and

Technology (ICTA), Guatemala, breeders for Central America. These cultivars derived reduced mosaic and superior yielding resistance from 'Porrillo Sintético' (G4495) and diffuse mottling resistance from Turrialba 1 (Beebe and Pastor-Corrales, 1991). Newly released cultivars, such as 'Dorado' (formerly DOR 364), 'Don Silvio' (formerly DOR 482), and 'Turbo III', have culminated ≈ 16 years of breeding for BGMV resistance (S. Beebe, personal communication). These cultivars have greater resistance to BGMV obtained from various combinations of partial resistances derived from 'Porrillo Sintético' and 'Turrialba 1', partial "nonmosaic" resistance of 'Garrapato' (G 2402), and tolerance of 'Honduras 46' (G 4791). Generally, in our breeding program and elsewhere (CIAT, 1995), new lines and cultivars with high levels of BGMV resistance carry the 'Garrapato' source of partial resistance in their pedigree.

Lack of expression of mosaic symptoms from 'Garrapato', following inheritance studies involving the breeding line A429, was conditioned by a single recessive gene (Blair and Beaver, 1993). In our paper, we have assigned a tentative symbol, *bgm-1*, to represent this recessive resistance gene. To date, *bgm-1* is widely used as a source of partial BGMV resistance. The resistance is considered partial because low yield and pod deformation are observed under moderate to high levels of disease incidence. Introgression of *bgm-1* is an important first step in breeding new BGMV-resistant cultivars. Combining *bgm-1* with other resistance genes that condition high yield is also important (Beebe, 1994).

For introgression purposes, the recessive nature of *bgm-1*

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requires a generation of selfing after every second to third backcross generation to obtain homozygous resistant BC_nF_2 parents for the next backcross cycle. Other obstacles associated with breeding for BGMV resistance include lack of uniform and consistent year-to-year disease pressure in the field. Greenhouse programs that use viruliferous whiteflies (Beaver et al., 1994) or a mechanical form of inoculation (Morales and Niessen, 1988) provide uniform inoculum, but risk introducing the virus into new regions, are time-consuming and do not provide reliable yield data.

Marker-assisted selection provides an ideal strategy for transferring and combining monogenic resistance traits in common bean (Kelly, 1995). Indirect selection of a recessive resistance gene based on a closely linked randomly amplified polymorphic DNA (RAPD) marker (Welsh and McClelland, 1990; Williams et al., 1990) can reduce the number of generations required in a backcross breeding program and limit the need for greenhouse and field testing. Our objective was to obtain a gene-pool-nonspecific (Young and Kelly, 1996) RAPD marker tightly linked to the recessive *bgm-1* gene from 'Garapato' that conditions partial nonmosaic resistance to BGMV. Identifying a dominant RAPD in repulsion-phase linkage with *bgm-1* is desired for rapid backcrossing of partial resistance to BGMV into susceptible types (Johnson et al., 1995).

Materials and Methods

Plant material. Random $F_{5,6}$ recombinant inbred lines (RILs) segregating for the recessive *bgm-1* gene from 'Garapato' that conditions nonmosaic partial resistance to BGMV were derived from a T446/A429 hybridization using single-seed descent. The RILs were used with DNA bulked segregant analysis (Michelmore et al., 1991) to initially screen 300 hundred arbitrary decamer primers in the polymerase chain reaction (PCR) for identifying RAPDs putatively linked to the *bgm-1* locus. T446 is an advanced pinto bean breeding line with high yield potential in the tropics. A429 is an advanced pinto breeding line from CIAT with upright architecture and combined BGMV resistances from 'Porriño Sintético' and 'Garapato'.

RAPD protocols. DNA was extracted using a modified "mini-

prep" procedure (Afanador et al., 1993), quantified by fluorometry (Hofer TKO 100, Hofer Scientific, San Francisco), and standardized to $10 \text{ ng} \cdot \mu\text{L}^{-1}$. Two contrasting DNA bulks were formed, one with five F_6 RILs with uniform nonmosaic resistance and the other with five F_6 RILs with uniform susceptibility. Uniformity of disease reaction of selected RILs initially was based on natural infection in the field and then later confirmed by greenhouse inoculations using whiteflies viruliferous for the Puerto Rican strain of BGMV (Beaver et al., 1994). DNA bulks were screened with 300 decamer primers (Operon Technologies, Alameda, Calif.).

The PCR consisted of 25- μL reactions containing two units of Stoffel fragment DNA polymerase (Perkin Elmer Cetus, Norwalk, Conn.), $1 \times$ Stoffel buffer, 5 mM MgCl_2 , 200 M of each dNTP, 0.2 M primer, and 25 ng template DNA. A DNA thermal cycler (model 480; Perkin Elmer) was used. The thermo-cycling profile consisted of three initial cycles of denaturing at 94°C for 1 min, annealing at 35°C for 1 min, and extension at 72°C for 2 min, followed by 31 cycles of 94°C for 10 s, 40°C for 20 s, and 72°C for 2 min, with a 1-s autosegment extension for each cycle (Haley et al., 1994). A final extension cycle of 72°C for 5 min was conducted before cooling the samples to 4°C . Amplified products were separated on 1% agarose gels containing ethidium bromide ($0.5 \text{ g} \cdot \text{mL}^{-1}$) for 5 h at $3 \text{ V} \cdot \text{cm}^{-1}$ constant voltage.

Linkage analyses. RAPDs observed between contrasting DNA bulks were assayed across individual lines that comprised the bulks. RAPDs that cosegregated among lines within bulks then were screened across five F_2 populations that segregated for the *bgm-1* gene. One dry bean population consisted of 91 F_2 progeny derived from a XAN 176/T446/A429 hybridization. The T446/A429 parent was an F_5 line with nonmosaic resistance obtained from the previously mentioned RIL population. A population of 48 progeny was combined across the other four F_2 populations involving 'Quest', 'Mirada', 'Seville', and 'Duchess' snap beans crossed with A429, because the heterogeneity χ^2 for 1:3 segregation for disease reaction amongst them was nonsignificant. Phenotypic disease reactions of F_2 populations were determined by greenhouse inoculations using viruliferous whiteflies (Beaver et al., 1994). Only one RAPD (codominant) marker was useful following F_2 cosegregation analysis.

F_2 cosegregation analysis.

Chi-square tests for expected 1 (*bgm-1bgm-1*) partial nonmosaic resistance : 3 (*Bgm-1Bgm-1*; *Bgm-1bgm-1*) susceptible phenotypic segregation at the resistance gene locus and 1:2:1 genotypic segregation at the codominant RAPD marker locus were performed for both F_2 populations. Maximum-likelihood formulae were used to estimate recombination frequencies between *bgm-1* and the codominant RAPD marker in both F_2 populations (Suiter et al., 1983).

The codominant RAPD marker found linked to the BGMV resistance locus was assayed across susceptible lines and cultivars to determine the range of germplasm in which the marker would be useful for indirect selection (Miklas et al., 1993,

Table 1. Chi-square tests for phenotypic segregation at the *bgm-1* locus that confers partial resistance to bean golden mosaic virus and segregation at the $R2_{570/530}$ codominant RAPD marker locus and linkage estimates between the two loci.

F_2 population	No. of progeny	Observed	Expected	χ^2	P
Resistant gene					
(1 resistant : 3 susceptible)					
Dry bean	91	22:69	23:68	0.03	>0.80
Snap bean	48	6:42	12:36	4.00	>0.01
Codominant RAPD					
$R2_{570/530}$ (1-/+ : 2+/+ : 1+/-) ²					
Dry bean	91	22:47:22	23:45:23	0.10	>0.50
Snap bean	48	8:34:6	12:24:12	11.00	>0.01
Marker-gene		Linkage distance			
Dry bean (zero recombinants)		- cM			
Snap bean (two recombinants)		4.2 cM			
Across populations		1.4 cM			

²Phenotypes for the codominant $R2_{570/530}$ RAPD marker: -/+, 570-base pair (bp) fragment allele absent/530-bp fragment allele present, and the RAPD phenotype indicating homozygous resistance (*bgm-1bgm-1*) at the linked recessive gene locus; +/+, both fragment alleles present indicating heterozygous susceptible (*Bgm-1bgm-1*); and +/-, 570-bp fragment allele present/530-bp fragment allele absent indicating homozygous susceptible (*Bgm-1Bgm-1*).

1996a; Young and Kelly, 1996). To ascertain maintenance of the original linkage orientation of the codominant RAPD marker to the alleles at the *bgm-1* locus after numerous meioses, advanced lines and cultivars known to possess the *bgm-1* nonmosaic resistance from 'Garrapato' were likewise assayed for the resistance-linked codominant RAPD.

Results and Discussion

Only two RAPDs were tightly linked to the recessive *bgm-1* gene from 'Garrapato' that conditions nonmosaic partial resistance to BGMV (Table 1). These RAPDs were amplified by the same primer R2 (5'-CACAGCTGCC-3') and were codominant, with the 530 and 570 base pairs (bp) fragment length alleles of the marker in repulsion- and coupling-phase linkage (Miklas et al., 1996a) with the recessive resistance allele *bgm-1*, respectively (Fig. 1). The ability of the DNA bulks to detect a codominant RAPD marker is due to the allelic homozygosity at the *bgm-1* locus for each RIL used in the DNA bulk analysis and the uniformity and reliability of field and greenhouse screening tests used to detect the nonmosaic partial BGMV resistance conditioned by the *bgm-1* gene.

There were no recombinations between the codominant RAPD marker and *bgm-1* loci detected in the dry bean F_2 population, and only two recombinants were detected in the combined snap bean F_2 population (4.2 cM) (Table 1). In several other snap bean F_2 populations (data not reported), recombination averaged 11.0 cM between R2_{570/530} and the *bgm-1* gene. The difference in recombination frequency between the two population types might be attributed to higher recombination suppression occurring in one population vs. the other, which generally has been observed when linked loci have been transferred between, rather than within, common bean gene pools (Young and Kelly, 1996). See Singh et al. (1991) for a review of race and gene pool delineation of common bean germplasm. Observing a loose linkage in exotic F_2 snap bean populations suggested that linked loci might be of Andean origin or perhaps traced back to an outcross *Phaseolus* species, such as *P. coccineus*, with known accessions having apparent immunity to BGMV (Beebe and Pastor-Corrales, 1991). Based on seed and agronomic characteristics, 'Garrapato' belonged to the Durango race within the middle American gene pool, which apparently represented the origin of *bgm-1* and the codominant RAPD marker R2_{570/530}.

Survey of susceptible common bean lines and cultivars for presence of the respective 570-bp and 530-bp fragment alleles of the codominant RAPD (Table 2) revealed that the marker was gene

Fig. 1. Occurrence of the R2_{570/530} codominant RAPD marker in snap bean. Snap parents (lanes 1 to 3 and 5), A429 parent (lane 4), and resistant control (lane 6). F_2 segregants: lanes 7 to 19. R2_{570/530} phenotype: -/+ , 570 base pair- (bp) fragment allele absent/530-bp fragment allele present, and the RAPD configuration indicating homozygous resistance (*bgm-1bgm-1*) at the linked recessive *bgm-1* gene locus; +/- both fragment alleles present indicating heterozygous susceptible (*Bgm-1bgm-1*); and +/+, 570-bp fragment allele present/530-bp fragment allele absent indicating homozygous susceptible (*Bgm-1Bgm-1*). M corresponds to the DNA molecular size marker *HindIII/EcoRI* digest of DNA.

Table 2. Survey of the R2_{570/530} codominant RAPD marker linked to the *bgm-1* resistance gene across susceptible germplasm, parents, and controls with 'Garrapato'-derived nonmosaic partial resistance to bean golden mosaic virus (BGMV).

Germplasm ^z	R2 _{570/530}
Kidney and miscellaneous (A)	
CELRK	+/- ^y
Linden	+/-
Montcalm	+/-
Pompadour G	+/-
Cacahuete 72	+/-
Royal Red	+/-
3M 152	+/-
DOR 303	+/-
Snap(A + MA)	
BelJersey-RR-14	+/-
Early Gallatin	+/-
Sprite	+/-
Pinto (MA)	
Arapaho	+/-
Bill Z	+/-
Othello	+/-
P 90521	+/-
BelDakMi-RR-1, -2	+/-
UI 114	+/-
PX 057	+/-
2-2186	+/-
3-4510	+/-
3-4537	+/-
Garrapato	-/+ ^x
JB 9226-5	-/+ ^x
TARS-VCI-4B	+/-
Black (MA)	
B 190	+/-
Turbo III	-/+ ^x
Negro 150	+/-
Porrillo Sintetico	+/-
Red(MA)	
PI 451895	+/-
Dorado	+/-
Don Silvio	-/+ ^x
Red Mexican 36	+/-
White (MA)	
Belmidak-RR-1	+/-
Belmidak-RR-2	+/-
Arroyo Loro	+/-
Seafarer	+/-
Schooner	+/-
Sanilac	+/-
C-20	+/-
NX 040	+/-
Mayflower	+/-
Parents	
Quest (snap)	+/-
Mirada (snap)	+/-
Seville (snap)	+/-
Duchess (snap)	+/-
XAN 176 (black)	+/-
T446/A429 (pinto)	-/+ ^x
A429 (pinto)	-/+ ^x

^zA and MA indicate germplasm of Andean and Middle American origin, respectively.

^yR2_{570/530} codominant RAPD marker genotype and corresponding genotype at the linked *bgm-1* BGMV resistance gene locus: -/+ homozygous resistant (*bgm-1bgm-1*); +/- heterozygous BGMV susceptible (*Bgm-1bgm-1*); and +/+ - homozygous BGMV susceptible (*Bgm-1Bgm-1*).

^xLines and cultivars with BGMV resistance derived from 'Garrapato', but often obtained through hybridizations with the dry bean A429.

pool nonspecific and, thus, useful for marker-assisted selection across a wide range of germplasm. A429 and ‘Garrapato’, the known source of *bgm-1*, carried only the resistance-linked 530-bp fragment allele in repulsion-phase linkage with the recessive resistance allele *bgm-1*. ‘Turbo III’ and ‘Don Silvio’, derived from A429 (CIAT, 1995) and carrying the nonmosaic *bgm-1* resistance from ‘Garrapato’, also contained only the resistance-linked 530-bp fragment allele. All susceptible lines and cultivars surveyed possessed only the susceptibility-linked 570-bp fragment allele in

coupling-phase linkage with *bgm-1*, indicating they were *Bgm-1Bgm-1* genotypes.

The original linkage orientation between the 570- and 530-bp fragment alleles of the codominant RAPD and linked *bgm-1* gene persisted through the numerous meioses involved in development of advanced lines with nonmosaic partial resistance from ‘Garrapato’ (Table 3). Recombination between the linked loci were detected in only two advanced lines, Red mottled-2 and DOR 482/9231-94. The line DOR 303/T968 appeared to be recombinant for the linked loci because it had a different source of nonmosaic partial resistance derived from DOR 303 instead of ‘Garrapato’.

Since there were higher recombination frequencies for snap bean between the marker and gene loci, we recommend that separate and distinct F_1 -derived populations or lines be maintained throughout any $R_{2570/530}$ -assisted snap bean backcrossing program. To monitor recombination, we recommend every third or fourth generation to screen the disease reaction of selfed progeny from parents heterozygous or homozygous for the 530-bp fragment allele. Also noteworthy, *bgm-1* nonmosaic partial resistance by itself may not provide economic control of BGMV during moderate to severe epidemics. The most effective genetic control to BGMV is obtained through combined resistance. From field observations made in Puerto Rico, A429, which likely combines reduced mosaic and high yield from ‘Porriillo Sintético’ with *bgm-1* nonmosaic partial resistance from ‘Garrapato’, is more resistant than ‘Garrapato’ alone; moreover, ‘Don Silvio’, which likely combines the former two forms of resistance with an additional source, is slightly more resistant than A429.

The $R_{2570/530}$ codominant RAPD marker will be extremely useful for rapidly breeding BGMV resistance into all snap bean and susceptible dry bean market classes. First, the application of the marker is not limited by gene pool specificity (Table 1); therefore, it can be used for indirect selection of the *bgm-1* gene across all common bean germplasm. Second, the marker is highly repeatable and easy to score, and has been amplified across several different laboratories using similar but distinct DNA extraction and PCR protocols. However, when using lower quality DNA obtained from mini-prep extraction protocols, a Stoffel fragment or similarly engineered DNA polymerase may be critical to the reaction. Third, by selecting BC_1F_1 plants heterozygous for the marker as parents for subsequent backcrosses, it becomes unnecessary to develop and identify BC_nF_2 parents with nonmosaic partial resistance every second to third backcross. Last, linkage of a codominant RAPD marker to a recessive resistance gene can be manipulated to facilitate gene pyramiding (M. Bassett, personal communication). For example, advanced near-inbred lines can be developed with resistance in the form of reduced or delayed mosaic symptom expression to BGMV similar to ‘Dorado’ (Miklas et al., 1996b), while simultaneously maintaining heterozygosity at the $R_{2570/530}$ locus. On selfing these lines and then selecting individual plants homozygous for the 530-bp fragment allele, the *bgm-1* partial resistance can be effectively pyramided with any other source of BGMV resistance, be it independent, epistatic, or hypostatic to the phenotypic expression of the *bgm-1* gene.

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Table 3. Advanced breeding lines with *bgm-1*-conditioned nonmosaic partial resistance to bean golden mosaic virus (BGMV) characterized for the linked $R_{2570/530}$ codominant RAPD marker.^z

Advanced line	Reaction		$R_{2570/530}$
	Greenhouse ^y	Field ^y	
Arroyo Loro/DOR 482	R ^x		–/+ ^x
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	R		–/+
Arroyo Loro/DOR 482	S		+/+
Red Mottled-1	S		+/+
Red mottled-2	S		–/+ ^w
Red mottled-3	R		–/+
Red mottled-4	R		–/+
JB 9438-126	S		+/-
JB 9438-238	R		–/+
A429	R		–/+
DOR 483/BelNeb-RR-1		R ^v	–/+ ^v
MUS 83/DOR 483		R	–/+
DOR 475/Triumph		Seg.	–/+ ^u
DOR 475/Triumph		Seg.	–/+
DOR 483/A429		R	–/+
DOR 483/A429		R	–/+
A429/Triumph		R	–/+
DOR 483/BelNeb-RR-1		R	–/+
A429/Triumph		R	–/+
DOR 482/9231-94		R	+/- ^w
DOR 303/T968		R	+/-
DOR 483/A429		R	–/+
DOR 483/A429		R	–/+
I 9449-33		R	–/+
DOR 483		Seg.	+/+

^z $R_{2570/530}$ phenotype: –/+, 570-base pair (bp) fragment allele absent/530-bp fragment allele present, and the RAPD configuration indicating homozygous resistance (*bgm-1bgm-1*) at the linked recessive gene locus; +/+ both fragment alleles present indicating heterozygous susceptible (*Bgm-1bgm-1*); and +/-, 570-bp fragment allele present/530-bp fragment allele absent indicating homozygous susceptible (*Bgm-1Bgm-1*).

^yGreenhouse inoculations using whiteflies (Beaver et al., 1994) viruliferous for the BGMV strain endemic to Puerto Rico and field reactions obtained by natural infection.

^xFor each line, phenotypic greenhouse reaction to BGMV was averaged across three plants and the RAPD genotype was derived from pooled DNA.

^wLines with recombination between linked loci.

^vFor each line, phenotypic field reaction to BGMV was obtained on a plot basis, and the RAPD genotype was obtained from pooled DNA of three random plants grown in the greenhouse.

^uLines segregating could be either +/+ or –/+ depending on composition of the pooled DNA.

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