Alteration in Lignin Biosynthesis Restricts Growth of *Fusarium* spp. in Brown Midrib Sorghum

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**ABSTRACT**


To improve sorghum for bioenergy and forage uses, *brown midrib (bmr)*6 and -12 near-isogenic genotypes were developed in different sorghum backgrounds. The *bmr6* and *bmr12* grain had significantly reduced colonization by members of the Gibberella fujikuroi species complex compared with the wild type, as detected on two semiselective media. *Fusarium* spp. were identified using sequence analysis of a portion of the translation elongation factor (TEF) 1-α gene. The pathogens *Fusarium thapsinum*, *F. proliferatum*, and *F. verticillioides*, *G. fujikuroi* members, were commonly recovered. Other frequently isolated *Fusarium* spp. likely colonize sorghum asymptomatically. The χ² analyses showed that the ratios of *Fusarium* spp. colonizing *bmr12* grain were significantly different from the wild type, indicating that *bmr12* affects colonization by *Fusarium* spp. One *F. incarnatum-F. equiseti* species complex (FIESC) genotype, commonly isolated from wild-type and *bmr6* grain, was not detected in *bmr12* grain. Phylogenetic analysis suggested that this FIESC genotype represents a previously unreported TEF haplotype. When peduncles of wild-type and near-isogenic *bmr* plants were inoculated with *F. thapsinum*, *F. verticillioides*, or *Alternaria alternata*, the resulting mean lesion lengths were significantly reduced relative to the wild type in one or both *bmr* mutants. This indicates that impairing lignin biosynthesis results in reduced colonization by *Fusarium* spp. and *A. alternata*.

Additional keywords: caffeic acid O-methyltransferase, cinnamyl alcohol dehydrogenase, *Fusarium equiseti*.

Sorghum (*Sorghum bicolor* (L.) Moench) is a valuable crop due to its adaptability to drought-stressed conditions (16). Recently, interest in sorghum grain and biomass has been piqued because of their usability as feed stock for bioenergy industries (54). Genetic modification to reduce lignin content in sorghum and other potential bioenergy crops has promise for greater utilization by increasing biomass digestibility (32). However, lignin is important for structural support and water transport within the plant (7). Additionally, lignin is involved in defense against pathogens and insects by providing a physical barrier (18,62), and lignin biosynthesis is induced in response to biotic stresses (26).

Two different mutated genes in the sorghum lignin biosynthesis pathway have allowed the development of forage and grain lines with significantly reduced lignin content (46). These mutations, *brown midrib (bmr)*6 and *bmr12*, result in the phenotype of reddish-brown leaf midveins which are associated with reduced lignin content and modified lignin composition (46,48). *Bmr6* encodes cinnamyl alcohol dehydrogenase (CAD) (53,57) and *Bmr12* encodes caffeic acid O-methyltransferase (COMT) (5). CAD converts the aldehyde group of monolignols to an alcohol for incorporation into the lignin polymer, while COMT methylates the 5-hydroxyl group of the hydroxyl coniferol substrates forming sinapyl groups (24). Both *bmr6* and *bmr12* are caused by nonsense mutations that result in the absence of detectable gene products (5,48,57).

When *bmr6*, *bmr12*, and near-isogenic wild-type lines were grown at two locations in field trials, density of *bmr6* grain was reduced (46), which could be caused by infection by grain-molding pathogens (34). However, when some of this same grain was screened, results indicated that *bmr* genotypes could have significantly fewer colonies of *Fusarium* spp. per seed (20). Among species identified using morphological characters were *Fusarium thapsinum* (formerly known as *F. moniliforme*), *F. proliferatum*, and *F. subglutinans*, members of the species complex Gibberella fujikuroi (31,44). Greenhouse inoculation of peduncles of *bmr6* and *bmr12* plants with *F. thapsinum* resulted in mean lesion lengths significantly smaller than those on near-isogenic wild-type plants (20). When considering genetic background, one or both low-lignin lines had significantly smaller mean lesions as a result of inoculation with *F. thapsinum* in four of six genetic backgrounds. Therefore, this research indicated that *bmr* resulted in reduced colonization by *Fusarium* spp but genetic background influenced the interaction with low-lignin plants (20).

Several *Fusarium* spp. in the *G. fujikuroi* species complex are potentially pathogenic on sorghum (31). Infections of stalks or roots can reduce yield and grain quality while lodging can make harvest difficult (6). Infections of panicles can result in grain of reduced quality and usability (36). *F. thapsinum* and other *Fusarium* spp., including those considered nonpathogenic, are able to colonize sorghum without producing symptoms (20,21,30). Many *Fusarium* spp. produce mycotoxins (31,41). Accumulation of these metabolites are often exacerbated when grain is stored or stover is ensiled (1,13). Deterioration by grain-molding fungi and contamination by mycotoxins may be a particular problem in cultures in which sorghum is used in foods and beverages as well as for livestock feed, and grain is stored in traditional containers on rural farms (14,55). Long-term exposure to mycotoxins
can result in life-threatening diseases to animals and possibly to humans (60).

Because grain of \textit{bmr} genotypes exhibited reduced colonization by \textit{Fusarium} spp. and peduncles had reduced colonization by \textit{F. thapsinum}, the hypothesis that these genotypes had specific interactions with \textit{Fusarium} spp. was proposed. \textit{Fusarium} isolates from grain of field-grown \textit{bmr} and wild-type plants were identified to species using sequence analysis of the translation elongation factor (\textit{TEF}) 1-\textalpha{} gene. Five \textit{Fusarium} genotypes and one \textit{Alternaria} sp. commonly found in sorghum fields, plants, or debris were inoculated onto peduncles of \textit{bmr} plants and near-isogenic wild-type plants under greenhouse conditions. Results of this study may elucidate differential colonization of grain and stalks of \textit{bmr} and near-isogenic wild-type plants.

**MATERIALS AND METHODS**

\textit{Fusarium} and \textit{Alternaria} spp. colonization of field-grown grain from \textit{bmr} and wild-type plants. Plants of the wild type, \textit{bmr6}, and \textit{bmr12} in genetic backgrounds BTX623, BTX631, and Redlan and wild-type, \textit{bmr6}, \textit{bmr12}, and \textit{bmr6}, \textit{bmr12} double mutant plants in backgrounds Wheatland and RTx430 were grown in two-row plots, 7.6 m long and spaced 76 cm apart, at University of Nebraska Field Laboratories in Lincoln and Ithaca, NE in 2003 (49,50). During 2004, \textit{bmr6}, \textit{bmr12}, double-mutant, and wild-type near-isogenic lines in Wheatland and RTx430 backgrounds were grown at both locations.

For 2003, nitrogen fertilizer was applied preplant at both locations at 112 kg ha\(^{-1}\). At the Lincoln location, propachlor (2-chloro-N-(1-methylthyl)-N-phenylacetamide) and atrazine (6-chloro-N-ethyl-N′-(1-methylthyl)-1,3,5-triazene-2,4-diamine) were applied at 3.56 and 1.1 kg ha\(^{-1}\), respectively, immediately after planting for weed control. At Ithaca, atrazine was applied at 2.2 kg ha\(^{-1}\) immediately after planting, followed by an application of quinclorac (3,7-dichloro-8-quinolinicarboxylic acid) and atrazine at 0.37 and 1.1 kg ha\(^{-1}\), respectively, \(\pm 14\) days postemergence. At Ithaca, supplemental irrigation was applied via overhead sprinklers on 24 July (2.5 cm) and 4 (5.0 cm), 7 (5.0 cm), 14 (2.5 cm), and 28 (2.5 cm) August. In 2004, nitrogen fertilizer was applied prior to planting at both locations at 157 kg ha\(^{-1}\) and herbicides were applied as in 2003. Supplemental irrigation (2.5 cm) was applied at Ithaca on 3, 12, and 19 August. These field plantings also were yield studies; therefore, no artificial inoculations were conducted and irrigation was applied to avoid water stress. Grain yields, test weights, and fiber and grain content were previously reported (46,50,56). In 2003, lodging scores approached zero. Visual inspection of grain samples from both years showed no obvious signs or symptoms of grain molding.

For each plot from both locations, grain were sampled from 10 randomly chosen heads throughout both rows and combined in a seed envelope. Seed from each plot were surface sterilized by washing for 2 min in 95% ethanol followed by a 10-min wash in 1% sodium hypochlorite with 0.01% Tween 20. Five seeds from each plot sample were asceptically placed onto one each of two peptones (Becton, Dickinson and Co. [BD], Sparks, MD) agar media containing either chloramphenicol and the fungicide dichloran (dichloran chloramphenicol peptone agar [DCPA]) (Sigma-Aldrich, St. Louis) (2) or streptomycin (Fisher Scientific, Fair Lawn, NJ) and the fungicide pentachloronitrobenzene (PCNB) ("Terrachlor"; Uniroyal Co., Middlebury, CT) (40). DCPA is semiselective for \textit{Fusarium} spp., \textit{Alternaria} spp., and other dark-spored ascomycetes, while PCNB is semiselective for \textit{Fusarium} spp. In our experience, using a combination of these two media appears to give a more complete picture of grain colonization by \textit{Fusarium} and \textit{Alternaria} spp., because the luxuriant growth of \textit{Alternaria} spp. on DCPA may overgrow some slower-growing \textit{Fusarium} spp. Alternatively, \textit{Alternaria} spp. are nearly completely selected against on PCNB (20). For analysis of grain from single-mutant lines, in 2003, four plots each of five genetic backgrounds for wild-type, \textit{bmr6}, and \textit{bmr12} genotypes were grown at two locations so that, in total, 200 seeds of each \textit{bmr} genotype were screened on each medium. For analysis of grain of double-mutant lines in the background of Wheatland and RTx430 and comparison with single-mutant and wild-type lines, in 2003, four plots of each genotype and each genetic background were planted whereas, in 2004, five plots of each genotype and each genetic background were planted. Thus, in total, 180 seeds of each genotype were screened on each medium.

Each fungal colony growing from an individual seed and onto the agar medium was transferred to one-half strength potato dextrose agar (PDA) (prepared with potato dextrose broth) (BD), \(\approx 3\) and 0.1% of seed had two colonies growing onto DCPA and PCNB, respectively. In these cases, each distinctive colony was transferred to PDA. Fungi were identified using colony morphology on PDA and conidiophore structures and conidial types and morphologies on minimal media (22,42). Putative \textit{Fusarium} spp. were transferred and grown on 1.5% agar medium containing 80 mM potassium chloride. Putative \textit{Alternaria} isolates were transferred by point inoculation to a 2% water agar medium, followed by aseptic placement of an \(\approx 1\)-cm\(^2\) filter paper square (Whatman no. 1; Whatman International Ltd., Maidstone, England) over the inoculation site. \textit{Fusarium} spp. were categorized as members of the \textit{G. fujikuroi} species complex using nomenclature previously described (31,44). Number of fungi per seed was enumerated in each of the following categories: total fungi, \textit{Alternaria} spp., \textit{Fusarium} spp. in the \textit{G. fujikuroi} species complex, and other \textit{Fusarium} spp.

The field experimental design was a split plot with whole plots arranged in a randomized complete block, with four replications in 2003 (single-mutant and double-mutant analyses) and five replications in 2004 (double-mutant analyses). Whole plots for single-mutant analyses were genetic background (RTx430, BTX623, BTX631, Redlan, and Wheatland) and subplots were \textit{bmr} genotype (the wild type, \textit{bmr6}, and \textit{bmr12}). Whole plots for double-mutant analyses were genetic background (RTx430 and Wheatland) and \textit{bmr} genotype (wild-type, \textit{bmr6}, \textit{bmr12}, and \textit{bmr6}, \textit{bmr12}). Analyses of colonization of grain from single-mutant lines compared with wild-type lines was conducted for each fungal category obtained on each medium. Analyses of colonization of grain from double-mutant lines compared with single-mutant lines and wild-type lines was conducted for each fungal category, and for each medium. Data were analyzed using Proc Mixed with the following model:

- **CLASS LOCATION REP GENE LINE**
- **MODEL DEPENDENT VARIABLE = GENE|LINE**
- **RANDOM LOCATION REPLOCATION**
- **REPEATED/GROUP = LOCATION**
- **LSMEANS GENE LINE/PDIFF**

Least squares means (LSM) and standard errors (SEs) are reported.

**Molecular identification of \textit{Fusarium} isolates from grain from field-grown \textit{bmr} and wild-type plants.** Molecular identification of representative field isolates was conducted, as described by Geiser and associates (23). This methodology utilizes small but consistent base pair differences in the sequence of the 5′ region of the \textit{TEF} gene that distinguish each species. The resulting \textit{FUSARIUM-ID} database has "excellent representation" of species members of \textit{G. fujikuroi} (23).

To conduct this analysis, isolates grown from grain in one plot and selected on either DCPA or PCNB medium were chosen for further analysis if they had different colony morphologies on PDA or conidial morphology on KCl and were considered to be representatives of different \textit{Fusarium} spp. These representative isolates from each plot were single-conidium purified. Isolates collected similarly from a study conducted in 2002 (20) also were included.
in this study. Sequence analyses were conducted on 105 of the 850 isolates collected and morphologically speciated from 2002, 2003, and 2004 field-grown grain.

DNA was extracted from ground lyophilized mycelium (19.29) and the S' region of TEF was polymerase chain reaction amplified using primers EF-1 and EF-2 (23). Amplification products were sequenced, and sequences were compared with those in the FUSARIUM-ID database (http://isolate.fusariumdb.org) (23). Among the sequenced isolates were 23 isolates that appeared to be members of the Fusarium incarnatum-F. equiseti species complex (FIESC) as previously defined by O’Donnell and associates in 2007 (45). FUSARIUM-ID and GenBank (4) were used to genotype 22 putative members of FIESC. Two isolates had the greatest similarity (93 to 96%) to Fusarium sp. no. 1321, formerly known as F. xanii sp. cf. equiseti, in FUSARIUM-ID. Using GenBank, members of this group, which will be known as FIESC no. 1321 in this study, were found to be most highly similar (92 and 96%) in the TEF region to haplotype FIESC 4-a (accession no. EF453007) (45). Seven isolates had the greatest similarity (≥99%) to F. pallidoroseum isolate no. 1320 in FUSARIUM-ID. These isolates, which will be referred to as FIESC no. 1320 for this study, when compared in GenBank, had high similarity to haplotype FIESC 3-a (98 to 99%) (accession no. EF453042) (45). Thirteen isolates had highest sequence identities (93 to 98%) to Fusarium sp. no. 1319, formerly known as F. xanii sp. cf. bullatula, in FUSARIUM-ID; members of this group will be called FIESC no. 1319 for this study. When compared in GenBank, these isolates were also most highly similar (91 to 93%) to FIESC 3-a. Sequences were deposited into GenBank (accession nos. EF152431, GQ339783 to GQ339793, and GU116575 to GU116584).

Due to a high frequency of zeros causing nonnormal distribution of data, and small sample sizes, analyses of Fusarium genotypes collected from single mutant grain (2002 and 2003) was conducted with χ² tests using PROC FREQ (SAS Institute, Cary, NC). Non-normality of the data was addressed using the exact option in PROC FREQ. Quoting SAS (SAS Institute): “Exact tests are appropriate when a data set is small, sparse, or heavily tied.”

Ratios of numbers of each Fusarium genotype obtained from bmrl6 or bmrl12 grain was compared with ratios obtained from wild-type grain, which was considered the expected. In most categories of Fusarium spp., genetic background (e.g., RTx430, BTx623, BTx631, Redlan, and Wheatland) was not significant, except for numbers of F. thapsinum. Therefore, numbers from each genetic background were combined for analyses. Ratios of Fusarium genotypes colonizing double-mutant grain compared with wild-type grain (2003 and 2004), and single-mutant grain compared with wild-type grain, in two genetic backgrounds (RTx430 and Wheatland) were similarly approached; however, χ² from some comparisons approached infinity, presumably due to some Fusarium spp. categories having one or more zeros. Therefore, pairwise comparisons within a Fusarium genotype were conducted; again, comparisons in which both the wild type and bmr genotype had “zeros” were not attempted due to infinite ratios. In this case, by using pairwise analyses, it was established that, within each Fusarium genotype category, there were no significant affects of genetic background. Thus, backgrounds were combined and pooled when making χ² comparisons of numbers of each Fusarium genotype obtained from bmrl6, bmrl12, or bmrl6, bmrl12 double-mutant grain with those found on wild-type grain, similar to as described above.

Phylogenetic analysis of field isolates and genotypes putatively in FIESC. TEF sequences corresponding to bases 16 to 660 in sequence from Fusarium sp. NRRL 43740 haplotype 3-a (accession no. EF453042) were compared with sequences of seed isolates collected from field studies in 2002, 2003, and 2004 and, presumably, members of FIESC. Also included were FIESC haplotypes 2-a (accession no. EF453019), 3-a, and 4-a (accession no. EF453007); and, from the FUSARIUM-ID database, F. pallidoroseum isolate no. 1320 and Fusarium spp. nos. 1319 and 1321. These sequences were chosen because, during BLAST searches, sequences from this study were highly similar to each database sequence. To represent Fusarium spp. outside of FIESC, sequences from F. graminearum Schwabe (M02-7035S-2) and F. thapsinum (M03-11337S-2), isolated from bmrl12 grain, also were included (GenBank accession nos. GU116585 and GU116586).

Lengths of sequences were 639 to 651 bases. The DNA sequence alignment was performed using ClustalW function of MacVector (MacVector, Inc., Cary, NC). The unrooted phylogram was constructed using the neighbor-joining method with 1,000 bootstrap replications; the distances were calculated using the Tamura-Nei algorithm. Gaps distributed proportionally and nodes with bootstrap confidence values <65% were collapsed.

Greenhouse inoculations of bmrl6, bmrl12, and wild-type Redlan plants. The following fungal isolates were used in inoculation studies. F. solani (Mart.) Appel & Wollenw. emend. Syd. & Hans, isolate FRC-M-0900, was obtained from sorghum field soil (Texas). Stalk isolates were F. verticillioioides (Sacc.) Nirenberg (= F. moniliforme) (isolate FRC-M-1141) (Missouri) and F. thapsinum Kliitch, Leslie, Nelson et Marasas sp. nov. isolate (FRC-M-3790) (California) (referred to hereafter as F. th.-1). These three isolates were obtained from the Fusarium Research Center, Pennsylvania State University. F. armeniacum (G. A. Forbes, Windels & L. W. Burgess) L. W. Burgess & Summerell (formerly known as F. acuminatum) (isolate H02-781L-5b) and FIESC no. 1320 (formerly known as F. pallidoroseum or F. semitectum) (isolate M03-11261L-9) were isolated from sorghum leaves (Nebraska). F. thapsinum (isolate H03-11-9) (21) (referred to hereafter as F. th.-2) and Alternaria alternata (Fr.:Fr.) Keissl. (isolate H02-755S) (20) were obtained from sorghum grain (Nebraska). Species or genotype identifications were confirmed using sequence analysis of conserved DNA regions (GenBank accession nos. EF152426, EF152427, EF152428, EF152429, EU016680, and GU116571 to GU116574). Inocula were prepared by inoculating sterile toothpicks in 5 ml of sterile potato dextrose broth with a mycelium-covered agar plug from a PDA plate grown for 7 days. The broth-and-toothpick cultures were incubated for 2 weeks at 25°C prior to inoculations (25).

For greenhouse inoculations, plants of cv. Redlan near isogenic for bmrl6, bmrl12, and the wild type were grown by sowing seed into 25.4-cm-diameter pots containing standard soil mix. Seedlings were culled to two per pot; two plants of the same genotype were in each pot in order to utilize greenhouse space more efficiently while reducing the opportunity for error that would be associated with seeding multiple genotypes in a single pot. Cv. Redlan was chosen because, in a previous study, lesions on wild-type plants were intermediate in length following inoculation with F. th.-1 compared with lesions produced on five other genotypes (20). Peduncles of plants 2 weeks after anthesis (defined as approximately half the anthers exerted) were inoculated with fungal-infested toothpicks or toothpicks incubated in sterile broth (25). Plants within a single pot were inoculated with two different fungi or a fungal culture and broth control. Eighteen days following inoculation, peduncles were split longitudinally and the length of the discolored region was measured. This measurement will be referred to as “lesion length.”

The experimental design consisted of six replications containing 12 pots each, with each pot containing two plants. Individual plants were considered the experimental units. Within each replication, each of three bmrl genotypes was randomly assigned to four pots and one of eight inoculation treatments assigned to each plant so that each bmrl–inoculation treatment combination occurred once per replication. Pairings of inoculation treatments within pots were randomized for each replication. Replications were blocked by location within the greenhouse. The experiment
(set of six replications) was repeated thrice across time. When the three repetitions of the experiment were combined for analyses, there were significant time period and treatment–time period effects. Therefore, LSMeans for each repetition were analyzed separately using PROC MIXED with the following model:

CLASS ENVIRON REP BMR INOCULUM;
MODEL DEPENDENT VARIABLE = BMR|INOCULUM;
RANDOM REP POT(REP);
LSMEANS BMR|INOCULUM/DIFF;
LSMs and SEs are reported.

During the third repetition of the experiment, portions of each peduncle were analyzed for fungal growth. After measuring lesion lengths from plants in each block, one-half of a longitudinally split peduncle was surface sterilized in 1% sodium hypochlorite and rinsed in sterile distilled water, three times. Workers changed disposable gloves and sprayed gloved hands with 70% ethanol before handling each peduncle. Using surface-sterilized scalpels and disposable gloves and sprayed gloved hands with 70% ethanol before handling each peduncle.

RESULTS

_Fusarium and Alternaria_ spp. colonization of field-grown grain from _bmr_ and wild-type plants._ Colonization of grain from _bmr6_, _bmr12_, and near-isogenic wild-type plants (grown in 2003) was analyzed by selecting fungi grown from random seed from each sample on PCNB and DCPA media. Across genetic backgrounds, _bmr_ genotypes had a significant effect on total numbers of colonies when selected on either medium (P ≤ 0.02) (Table 1). The _bmr_ genotype also had a significant effect on numbers of _G. fujikuroi_ isolates per seed when selected on PCNB (P = 0.04); isolates per grain from _bmr12_ plants were significantly less than those from wild-type seed (Table 1). When selected on DCPA medium, _bmr_ genotype had a significant effect on _G. fujikuroi_ isolates per seed for colonization of _bmr6_ grain than when compared with wild-type plants (P = 0.01). Numbers of _Alternaria_ spp. selected on DCPA per seed in _bmr6_ grain were significantly fewer than those from _bmr6_ grain (P = 0.03). There was no apparent significant effect of _bmr_ genotype on numbers of other _Fusarium_ spp. in this analysis when fungi grown from seed were selected on either medium (Table 1).

Fungal colonies growing from field-grown grain of _bmr6_, _bmr12_ double-mutant plants, as well as _bmr6_, _bmr12_, and wild-type plants in the genetic backgrounds Wheatland and RTx430 (2003 and 2004), also were selected on DCPA or PCNB (Table 2). In this analysis, there was no indication that combining mutations had a greater effect than each individual mutation in near-isogenic lines on colonization by the fungi assayed in this study. When selected on DCPA, grain from _bmr_ genotypes in the background Wheatland had significantly fewer _G. fujikuroi_ isolates than the wild type while selection on PCNB of RTx430 _bmr12_ grain resulted in significantly fewer _G. fujikuroi_ isolates compared with other _bmr_ genotypes in this background (Table 2). The _bmr_ genotype did not have a significant effect on colonization of grain by other _Fusarium_ spp. in either background. However, significantly more other _Fusarium_ spp. were recovered on PCNB medium from Wheatland _bmr12_ grain than from RTx430 _bmr12_ grain (P = 0.04) (Table 2). When considering numbers of _Alternaria_ spp., colonies per seed selected on DCPA were significantly higher in wild-type RTx430 grain than from near-isogenic _bmr12_ and _bmr6_ _bmr12_ double-mutant grain; however, differences in colonization of wild-type and _bmr_ grain in Wheatland background were not significant (Table 2). _Fusarium_ spp. obtained from wild-type grain, as determined by morphological characters (42), were _F. moniliforme_ (e.g., unable to distinguish whether _F. verticillioides_ or _F. thapsinum_), _F. proliferatum_ (Matsush.) Nirenberg ex Gerlach & Nirenberg, _F. solani_, _F. subglutinans_ (Wollenw. & Reinking) P. E. Nelson, _F. thapsinum_, _F. verticillioides_, and unidentified _Fusarium_ spp. _Fusarium_ spp. obtained from _bmr6_ grain were similar to those from wild-type grain, except no _F. proliferatum_ isolates were identified morphologically. _Fusarium_ spp. isolated from _bmr12_ grain and identified by morphological characteristics were _F. aquaeductuum_ (Radlk. & Rabenh.) Lagerh., _F. moniliforme_, _F. proliferatum_, _F. verticillioides_, and unidentified _Fusarium_ spp. As determined by morphological characteristics, _F. avenaeceum_ (Fr.) Sacc., _F. equiseti_ (Corda) Sacc.

<table>
<thead>
<tr>
<th><em>bmr</em> genotype</th>
<th>Medium</th>
<th>Total colonies</th>
<th><em>Alternaria</em> spp.</th>
<th>Gibberella fujikuroi</th>
<th>Other <em>Fusarium</em> spp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>DCPA</td>
<td>0.92 ± 0.07</td>
<td>0.72 ± 0.05</td>
<td>0.09 ± 0.03</td>
<td>0.02 ± 0.01</td>
</tr>
<tr>
<td><em>bmr6</em></td>
<td>DCPA</td>
<td>0.94 ± 0.07</td>
<td>0.74 ± 0.05</td>
<td>0.02 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td><em>bmr12</em></td>
<td>DCPA</td>
<td>0.75 ± 0.07</td>
<td>0.61 ± 0.05</td>
<td>0.05 ± 0.03</td>
<td>0.02 ± 0.01</td>
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<tr>
<td>Wild type</td>
<td>PCNB</td>
<td>0.15 ± 0.05</td>
<td>na</td>
<td>0.09 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td><em>bmr6</em></td>
<td>PCNB</td>
<td>0.06 ± 0.05</td>
<td>na</td>
<td>0.06 ± 0.03</td>
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<tr>
<td><em>bmr12</em></td>
<td>PCNB</td>
<td>0.08 ± 0.05</td>
<td>na</td>
<td>0.04 ± 0.03</td>
<td>0.02 ± 0.01</td>
</tr>
</tbody>
</table>

1 Dichloran chloramphenicol peptone agar (DCPA) is semiselective for _Fusarium_ spp., _Alternaria_ spp., and other dark-spored ascomycetes and pentachloro-nitrobenzene (PCNB) medium is semiselective for _Fusarium_ spp.

2 Least squares means and standard errors are shown. Within each category of fungal counts and for each medium, values with differing letters are statistically significant at P ≤ 0.05; na indicates not applicable because PCNB is strongly selective against _Alternaria_ spp.

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sensu Gordon, *F. moniliforme*, *F. proliferatum*, *F. thapsinum*, *F. verticillioides*, and unidentified *Fusarium spp.* were recovered from grain of double-mutant plants.

**Molecular identification of Fusarium isolates from grain from field-grown bmr and wild-type plants.** Sequence analysis of the 5′ TEF region from representative *Fusarium* isolates from single-mutant *bmr* genotypes and near-isogenic wild-type genotypes (2003) and from a previously reported study (2002) (20) were genotyped by comparison with sequences of type isolates in FUSARIUM-ID. Eleven different species or genotypes were distinguished. The most frequently isolated from wild-type grain were *F. thapsinum*, FIESC no. 1319, and *F. proliferatum*. Other commonly isolated species or genotypes recovered from wild-type grain were FIESC no. 1320, *F. verticillioides*, *F. graminearum*, and *F. subglutinans*. Four species were recovered infrequently (from one or two plots) from these trials. The distribution of fungal isolates recovered from wild-type, *bmr6*, and *bmr12* grain is illustrated in Figure 1. The χ² analyses, using ratios of FIESC *spp.* isolated from wild-type grain as “expected,” showed that numbers of each genotype found on *bmr12* were significantly different (exact χ², P = 0.05). Further analyses, comparing numbers of each species or genotype found on *bmr12* grain with that found on wild-type grain, showed that the difference could be attributed, in part, to numbers of FIESC *spp.* from wild-type grain but was not detected in *bmr12* grain. Colonization by *F. proliferatum* also was reduced in *bmr12* grain compared with the wild type (P = 0.06).

Sequence comparisons between 5′ TEF regions from isolates representing *Fusarium* *spp.* obtained from *bmr6*, *bmr12* double-mutant grain also were made with those from single-mutant *bmr6* and *bmr12* and wild-type *Wheatland* and RTx430 plants (2003 and 2004). *F. thapsinum* was the predominant species isolated and there were no significant differences detected in this analysis for colonization by *F. thapsinum* of wild-type, *bmr6*, *bmr12*, or *bmr6*, *bmr12* grain (P ≥ 0.14). Other species isolated from wild-type and *bmr6* grain included *F. proliferatum*, *F. verticillioides*, and FIESC no. 1320. Surprisingly, none of these other species were recovered from *bmr12* grain and only FIESC no. 1320 was found in *bmr6*, *bmr12* double-mutant grain, whereas FIESC genotype no. 1321 was isolated only from *bmr12* and double-mutant grain in RTx430 and Wheatland genetic backgrounds. Pairwise comparisons for each fungal species showed there were no significant differences of numbers isolated from each *bmr* genotype compared with those isolated from wild-type grain (P > 0.25).

**Phylogenetic analysis of field isolates and genotypes putatively in FIESC.** When phylogenetic analysis was performed on TEF sequences from putative FIESC members obtained from sorghum grain during this study, results indicated that one genotype, represented by a clone, was present in *bmr6* and wild-type grain but not in *bmr12* grain. TEF sequences were compared with sequences of FIESC haplotypes; representative isolates of *F. pallidoroseum* (no. 1320), *Fusarium* sp. cf. *equiseti* (no. 1321), and *Fusarium* sp. cf. *bullatum* (no. 1319) from FUSARIUM-ID; and *F. graminearum* and *F. thapsinum* isolates obtained from *bmr12* grain (Fig. 2). All the sorghum grain isolates that were putatively identified as FIESC were in the larger clade that includes haplotypes of FIESC 2-a, 3-a, and 4-a and *F. pallidoroseum*, *Fusarium* sp. cf. *equiseti*, and *Fusarium* sp. cf. *bullatum*, and separately from *F. graminearum* and *F. thapsinum*, indicating that they were, indeed, members of FIESC. This large clade separated into three branches. One branch had a small clade that included haplotypes FIESC 2-a and 4-a and *Fusarium* sp. cf. *F. incarnatum-F. equiseti* species complex.

![Fig. 1. Distribution of Fusarium genotypes isolated from wild-type and near-isogenic brown midrib (bmr6) and bmr12 grain. DNA from representative Fusarium grain isolates collected from each field plot were used for sequence analysis of polymerase chain reaction fragments from a region of the translation elongation factor 1-a gene; sequences were compared with those in the FUSARIUM-ID database. A χ² analysis of ratios of the genotypes found in bmr6 and bmr12 grain were compared with those in wild-type grain; **“*” indicates that χ² is significant (P = 0.05). FIESC = Fusarium incarnatum-F. equiseti species complex.**](Image)

### Table 2. Fungal colonization of grain from bmr double mutant plants and near-isogenic bmr6, bmr12 and wild-type Wheatland and RTx430 plants as determined by selecting on two media (2003 and 2004)

<table>
<thead>
<tr>
<th>Medium</th>
<th>Plant background</th>
<th>bmr Genotype</th>
<th>Total colonies</th>
<th>Alternaria spp.</th>
<th>Gibberella fujikuroi</th>
<th>Other Fusarium spp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>DCPA</td>
<td>Wheatland</td>
<td>Wild-type</td>
<td>0.95 ± 0.08</td>
<td>0.80 ± 0.05</td>
<td>0.05 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>Wheatland</td>
<td>bmr6</td>
<td>0.99 ± 0.08</td>
<td>0.84 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>Wheatland</td>
<td>bmr6, bmr12</td>
<td>0.81 ± 0.08</td>
<td>0.68 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>RTx430</td>
<td>Wild-type</td>
<td>0.84 ± 0.08</td>
<td>0.68 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>RTx430</td>
<td>bmr6</td>
<td>0.72 ± 0.08</td>
<td>0.60 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>RTx430</td>
<td>bmr12</td>
<td>0.39 ± 0.08</td>
<td>0.32 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>DCPA</td>
<td>RTx430</td>
<td>bmr6, bmr12</td>
<td>0.48 ± 0.08</td>
<td>0.39 ± 0.06</td>
<td>0.03 ± 0.03</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>Wheatland</td>
<td>Wild-type</td>
<td>0.06 ± 0.05</td>
<td>na</td>
<td>0.04 ± 0.04</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>Wheatland</td>
<td>bmr6</td>
<td>0.08 ± 0.05</td>
<td>na</td>
<td>0.06 ± 0.04</td>
<td>0.02 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>Wheatland</td>
<td>bmr12</td>
<td>0.08 ± 0.05</td>
<td>na</td>
<td>0.05 ± 0.04</td>
<td>0.04 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>Wheatland</td>
<td>bmr6, bmr12</td>
<td>0.09 ± 0.05</td>
<td>na</td>
<td>0.02 ± 0.04</td>
<td>0.02 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>RTx430</td>
<td>Wild-type</td>
<td>0.11 ± 0.05</td>
<td>na</td>
<td>0.08 ± 0.04</td>
<td>0.01 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>RTx430</td>
<td>bmr6</td>
<td>0.07 ± 0.05</td>
<td>na</td>
<td>0.06 ± 0.04</td>
<td>0.00 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>RTx430</td>
<td>bmr12</td>
<td>0.04 ± 0.05</td>
<td>na</td>
<td>0.03 ± 0.04</td>
<td>0.00 ± 0.01</td>
</tr>
<tr>
<td>PCNB</td>
<td>RTx430</td>
<td>bmr6, bmr12</td>
<td>0.05 ± 0.05</td>
<td>na</td>
<td>0.04 ± 0.04</td>
<td>0.00 ± 0.01</td>
</tr>
</tbody>
</table>

1 Dichloran chloramphenicol peptone agar (DCPA) is semiselective for *Fusarium spp.*, *Alternaria spp.*, and other dark-spored ascomycetes and pentachloronitrobenzene (PCNB) medium is semiselective for *Fusarium spp.*
2 Least squares means and standard errors are shown. Within each category of fungal counts, for each plant genotype and medium, values with differing letters are statistically significant at P ≤ 0.05; na indicates not applicable because PCNB is strongly selective against *Alternaria spp.*
Another branch, which included all sorghum FIESC isolates but two, split into two clades. Interestingly, one clade had isolates only from the wild type and bmr6 (6). The other clade included *Fusarium* sp. cf. *bullatum*, haplotype FIESC 3-a, and *F. pallidoroseum* and grain isolates from wild-type, bmr6, and bmr12 plants. The third branch within the larger clade included only two isolates, both from *bmr12* plants. This analysis indicates that a previously uncharacterized haplotype of FIESC existed, which preferentially colonized wild-type and bmr6 grain but not bmr12 grain.

**Greenhouse inoculations of bmr6, bmr12, and wild-type Redlan plants.** Three repetitions of an assay in which peduncles of plants were inoculated with different fungi were conducted. These experiments indicated that, when near-isogenic bmr6, bmr12, and wild-type plants were inoculated by *F. verticillioides* or *A. alternata*, as well as two *F. thapsinum* isolates, the mean lesion lengths could be significantly shorter on bmr plants than those resulting on wild-type plants. Lesions (areas of discoloration) produced on peduncles of bmr6, bmr12, and wild-type plants in cv. Redlan background following inoculation with one of two *F. thapsinum* isolates; with an isolate of *F. verticillioides*, *F. armeniacum*, and *A. alternata*; or with sterile broth were compared in three repetitions. Because of significant effects, each repetition of the experiment was considered independently. For the first repetition, bmr genotype, inoculum, and their interactions were significant (*P* ≤ 0.04).

Inoculations of wild-type plants by either *F. thapsinum* isolate, *F. verticillioides*, *F. solani*, *F. armeniacum*, or *A. alternata* resulted in mean lesion lengths significantly greater than the discoloration resulting from broth inoculation (Fig. 3A). Mean lesion lengths produced on bmr plants by *F. thapsinum*, *F. solani*, *F. armeniacum*, *A. alternata*, or *F. verticillioides* (bmr6 plants only) were significantly smaller than those produced on wild-type plants. During the second repetition of the experiment, only inoculum was significant (*P* < 0.01); although inoculation of wild-type plants by *F. thapsinum* and *F. verticillioides* resulted in mean lesion lengths significantly greater than discoloration resulting from broth inoculation, there were no significant differences between mean lesion lengths produced on wild-type and bmr plants (Fig. 3B).

During the third repetition of the experiment, results were significant for bmr genotype, inoculum, and their interactions (*P* ≤ 0.03). Mean lesion lengths resulting from inoculation of wild-type plants with either *F. thapsinum* isolate, *F. verticillioides*, *F. armeniacum*, and *A. alternata* were significantly greater than discoloration resulting from inoculation with broth (Fig. 3C). Inoculation of both bmr genotypes by *F. thapsinum* and *F. verticillioides*, or bmr6 plants with *F. thapsinum* and *A. alternata*, yielded mean lesion lengths significantly less than those following inoculation of the same fungi onto wild-type plants (Fig. 3C). To provide an indication of the extent of fungal growth in peduncles of inoculated plants, 1-cm sections from peduncles were plated and assessed for growth (+ or –) on DCPA medium. Percentage of

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**Fig. 2.** Phylogenetic analysis of sequences of a portion of translation elongation factor (TEF) 1-α gene from putative *Fusarium incarnatum-F. equiseti* species complex (FIESC) isolates from sorghum grain, type isolates from FUSARIUM-ID database, and haplotypes from GenBank (see Materials and Methods). FIESC isolates were isolated from grain from 2002, 2003, and 2004 field studies. “Wt” indicates an isolate from wild-type grain, “6” indicates an isolate from brown midrib (bmr6) grain, and “12” indicates an isolate from bmr12 grain. Isolates nos. 1319, 1320, and 1321 and haplotypes 2-a, 3-a, and 4-a had greatest similarity to isolates from this study when BLAST was conducted in FUSARIUM-ID or GenBank databases, respectively. TEF sequences from *F. graminearum* and *F. thapsinum*, isolated from bmr12 grain, were included as outgroups. Bootstrap values (percentage) for 1,000 replicates are indicated by each node.
peduncle sections from outside the lesion, at lesion borders, or within the lesion for bmr6 or bmr12 plants inoculated with fungi were compared with those from wild-type plants (Table 3). Fungal growth from bmr6 peduncle sections was not significant but sections from borders and from within lesions of bmr12 peduncles had significantly greater fungal growth compared with lesions on wild-type peduncles.

**DISCUSSION**

In numerous studies, lignin has been implicated in defense against pathogens and has been proposed as a physical barrier against penetration, further ingress, or diffusion of pathogen-produced toxins (8,17). Although induction of lignin synthesis is likely to be part of a generalized resistance response (43), the results presented herein suggest that impairing lignin biosynthesis does not lessen resistance to some pathogens. In particular, we propose that bmr12 may modify phenylpropanoid metabolism including and beyond lignin biosynthesis, thus restricting or excluding growth of a genotype of FIESC from grain of these plants.

In a previous study, we showed that, when grain from two genetic backgrounds (RTx430 and Wheatland) grown at two locations was plated onto PCNB agar medium, numbers of other Fusarium spp. could be significantly reduced in bmr genotypes; numbers of G. fujikuroi isolates also could be significantly reduced but there appeared to be an effect of environment on colonization of grain (20). Even when plating larger numbers of grain from a given genetic background grown in different environments onto DCPA medium, the results for Fusarium spp. were less clear. We concluded that plating sorghum grain onto both media, side by side, which was done in the current study, appeared to present a more complete, if not more complicated, picture of infection or colonization of the grain mold fungi Alternaria and Fusarium spp. The modes of action of dichloran (in DCPA) or PCNB have not been reported. Tolerance of PCNB by Fusarium spp. was proposed to be due to a combination of reduced uptake as well as metabolism to less toxic compounds (39). PCNB appears to affect specific activity of glutathione-S-transferase in Fusarium spp. (12) as well as DNA excision repair in bacteria (11). In the present study, results from plating grain on both DCPA and PCNB to screen fungal growth from grain supported the contention that bmr genotype effects colonization by members of the G. fujikuroi species complex across genetic backgrounds (Table 1).

Representative Fusarium isolates from each plot from this and the previous study were genotyped by sequence analysis of a portion of the TEF gene, then compared with sequences from type isolates in the FUSARIUM-ID database. We were surprised to find that one genotype, having high similarities to FIESC isolate...

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**Fig. 3.** Mean lesion lengths resulting from inoculation of peduncles of cv. Redlan plants and near-isogenic brown midrib (bmr)6 and bmr12 plants with different sorghum fungi. Isolates that resulted in mean lesion lengths significantly greater than discolorations resulting from broth control following inoculation of wild-type plants are shown for three repetitions (A, B, and C). F. th.-1 and -2 are different Fusarium thapsinum isolates, and F. vert., F. sol., F. arm., and A. alt. indicate F. verticillioides, F. solani, F. armeniacum, and Alternaria alternata, respectively. Bars represent least squares means and standard errors; * indicates that mean lesion length is significantly less than that of the wild type following the same treatment (P ≤ 0.05).
no. 1319, did not appear to colonize bmr12 grain across plant genetic backgrounds. Several FIESC isolates from wild-type or bmr6 grain were most similar to genotype no. 1319, formerly known as *Fusarium* sp. cf. *bullatum*; therefore, for this study, they were referred to as FIESC no. 1319. Two of these isolates, from different locations in 2 years, had 98% similarity to FIESC no. 1319 (Wt-11 and Wt-12) (Fig. 2). The rest of the isolates had 93% similarity to FIESC no. 1319 (Wt-6 through Wt-10, Wt-11 through 6-4, and 6-6) and phylogenetic analysis indicated that these isolates, which apparently group within the FIESC, also group separately from genotype no. 1319 and from the three FIESC haplotypes most similar to sequences from the present study (2-a, 3-a, and 4-a) (45), indicating a previously unreported haplotype within FIESC. This new haplotype, isolated from wild-type and bmr6 grain, was excluded from colonizing bmr12 grain in our experiments (Figs. 1 and 2). Members of FIESC, identified morphologically in this study as *F. pallidosome* (also known as *F. semitectum*), *F. equiseti*, or simply *Fusarium* spp., probably colonize sorghum grain asymptomatically (47). Inoculation of peduncles with an FIESC no. 1319 isolate resulted in mean lesion lengths that were not significantly different from the broth control, while inoculation of developing flowers with the same isolate resulted in mature grain that was as healthy as uninoculated controls, as indicated by three measurements (22). In the present study, an FIESC no. 1320 isolate inoculated onto wild-type plants in three repetitions had mean lesion lengths not significantly different from those of the broth control (Fig. 3). These results suggest that at least some genotypes in FIESC are low in virulence on sorghum. However, FIESC isolates may cause deterioration of stored grain (38) or produce mycotoxins (27,63). In addition to apparently excluding at least one member of FIESC, evidence suggests that colonization by *F. proliferatum*, which is relatively common in wild-type and bmr6 grain, also was limited in bmr12 grain (Fig. 1). *F. proliferatum* is a member of species complexes that are known to cause grain mold and stalk rot diseases and can produce mycotoxins which reduce the quality and usability of sorghum grain (43,58,59).

**TABLE 3.** Percent of sections from fungal-inoculated peduncles having fungal growth as selected on dichloran chloramphenicol peptone agar (DCPA) medium.

<table>
<thead>
<tr>
<th>Peduncle section</th>
<th>Plant genotypea</th>
<th>bmr6</th>
<th>bmr12</th>
</tr>
</thead>
<tbody>
<tr>
<td>Outside lesion</td>
<td>Wild-type</td>
<td>40.5 (79)</td>
<td>48.8 (84)</td>
</tr>
<tr>
<td></td>
<td>bmr6</td>
<td>82.0 (78)</td>
<td>79.8 (84)</td>
</tr>
<tr>
<td></td>
<td>bmr12</td>
<td>88.3 (60)</td>
<td>93.1 (58)</td>
</tr>
</tbody>
</table>

* Peduncles inoculated with either *Fusarium thapsinum* (two isolates), *F. verticillioides*, *F. solani*, *F. armeniacum*, *F. incarnatum*-*F. equiseti* species complex no. 1320, or *Alternaria alternata* isolates, generated during the third repetition, were sectioned and plated to estimate extent of fungal growth. For each location along the peduncle, \( \chi^2 \) analysis of percentage of sections with fungal growth for each bmr genotype (bmr6 and bmr12) were compared with percentage of wild-type sections with fungal growth using PROC FREQ (SAS).

* Following measurement of lesions, peduncles from the third repetition were dissected (see Materials and Methods). From each peduncle, sections 1 cm above and below the lesion, 1-cm sections that included the upper or lower border of the lesion, and, if the lesion was >10 mm, 1-cm sections from within the lesion were cut and aseptically placed onto DCPA medium. Three days after plating, each section was scored for fungal growth onto the medium. Percentage of peduncle pieces with fungal growth are reported. For peduncles inoculated with broth, outside lesion sections from bmr6 plants had significantly more fungal growth (33.3%) compared with wild-type plants (0.0%) \( (P < 0.01) \). There were no significant differences in fungal growth from boundary sections from broth-inoculated lesions. There were too few “within-lesion” sections (lesions >1 cm) for comparison.

* Numbers in parentheses indicate number of sections tested. Percent growth from peduncle sections from each wild-type plant was compared with that from comparable sections from each similarly-treated bmr plant; * indicates percentages are significantly different from the wild type at \( P \leq 0.05 \).

Our previous study indicated that bmr grain had reduced colonization by *Fusarium* spp. and, when peduncles were inoculated with a virulent *F. thapsinum* isolate, mean lesion lengths on bmr plants were reduced compared with near-isogenic genotypes when six genetic backgrounds were compared (20). Other studies in which reduced lignin did not result in increased susceptibility to necrotrophic fungi have been conducted in smooth bromegrass and soybean. In the case of smooth bromegrass, clones with reduced lignin were more susceptible to a biotrophic fungus but not to two necrotrophic fungi (15). In soybean, it was shown that, among six cultivars with different levels of resistance to Sclerotinia stem rot, lignin content was positively correlated to disease severity (51). In the present study, it is likely that the reduced-lignin phenotypes in the near-isogenic bmr genotypes were due to defective genes in the lignin biosynthesis pathway (5,48,56). In general, these mutations have resulted in an overall reduction in colonization by *G. fujikuroi*. However, bmr12 also appeared to be selective against one or two *Fusarium* genotypes. Resistance genes that encode receptor proteins, which interact with specific microorganism-produced extracellular molecules (effectors) in a gene-for-gene manner, can block a potentially pathogenic genotype from infecting its host (9). However, in the present study, the bmr12 gene encoded for an impaired enzyme in the lignin biosynthetic pathway (9,57).

Biochemically, it is known that the bmr plants sampled for this study had reduced lignin content (as measured by acid detergent concentration [ADL]); at least one of the bmr genotypes from each plant background had significantly reduced ADL compared with that of its wild-type counterpart (46). In particular, both bmr6 and bmr12 lines in the Redlan background and another background commonly used in breeding, Wheatland, had significantly reduced ADL. Lignin pathway intermediates also were measured in bmr genotypes in RXT430 and Wheatland backgrounds (48). Genetic background appeared to interact with each bmr gene and in combination in the double mutants, affecting levels of different phenolics in sorghum stalks. In particular, levels of caffeic acid (CaF), an intermediate associated with the lignin biosynthetic pathway, were over three times higher in bmr6 or bmr12 Wheatland plants relative to the wild type (48). Additionally, wall-bound ferulic acid (FA), another lignin biosynthesis pathway intermediate, was >1.25 times greater in bmr12 Wheatland plants than in wild-type plants. These results show that there are differences in accumulations of phenolics in stalks of bmr plants compared with near-isogenic wild-type plants. In the present study, extent of lesion development in fungal-inoculated bmr12 peduncles appeared to be affected by plant response (Fig. 3C, Table 3). Discoloration resulting from inoculation of sorghum peduncles, the length of which was defined as “lesion length” in the present study, are due, at least in part, to plant pigments induced during fungal infection (10,33). In this study, extent of discoloration was significantly reduced in Redlan bmr12 following inoculation by either *F. thapsinum* or *F. verticillioides*. However, results in Table 3 suggest that extent of fungal growth may be slightly greater, but significantly so, beyond the region of discoloration in bmr12 plants versus wild-type plants, suggesting that bmr12 may interact with one or more factor such that the plant responds in a way which results in reduced pigmentation. Because of this interesting observation, we are conducting more extensive experiments to determine whether differential responses of bmr genotypes to pathogens are a direct or indirect result of altered accumulation of pigments and other phenolic compounds.

Previous studies have shown that Bmr6 and Bmr12 are highly homologous to their counterparts in other plants, indicating evolutionary conservation (5,56). Indeed, Bmr6 was shown to be more similar to its CAD gene orthologs in monocots (rice, maize, and sugarcane), dicots (*Arabidopsis*, tobacco, aspen, and eucalyptus), pine, and the lycophyte *Salaginella* than to other sorghum CAD genes (57). Although fewer COMT genes were available for com-
parison, the trend is the same; Bmr12 was more similar to lignin biosynthesis COMT genes in pine, maize, alfalfa, and *Arabidopsis* than to other COMT genes in sorghum (unpublished). These results suggest that Bmr6 and Bmr12 evolved primarily for lignin biosynthesis rather than for another function such as plant defense.

In spite of this, it is clear that lignin and related compounds could contribute to resistance. However, what about results from this and previous studies that suggest the contrary? Lignin biosynthesis is part of the phenylpropanoid pathway, which begins with the conversion of phenylalanine to cinnamic acid (CinA) by the enzyme phenylalanine ammonia lyase (PAL) (Fig. 4). CinA is a precursor of lignin monomers, salicylic acid (SA), and some phytoalexins, including 3-deoxyanthocyanidins produced by sorghum (24,33,37) (Fig. 4). FA, a derivative of feruloyl CoA or coniferaldehyde, and 3-deoxyanthocyanidin phytoalexins in sorghum grain are associated with resistance to fungi (Fig. 4) (3,61). During *Monilinia fructicola* infection of stone fruit, CafA, derived from caffeoyl CoA, reduced appressorium formation and inhibited production of the pathogenicity factors cutinase and endopolygalacturonidase (28). Because the pathways downstream of CinA are interconnected (Fig. 4), a mutation that affects one step in this phenylpropanoid metabolic network (e.g., lignin biosynthesis) could shift intermediates toward other branches, resulting in elevated levels of secondary metabolites such as aromatic phytoalexins or SA. *Arabidopsis* mutants impaired in COMT1, the gene orthologous to sorghum bmr12, were more susceptible to two necrotrophic fungi, one biotrophic fungus, and two bacterial pathogens (52). However, they were not more susceptible to the pathogenic nematode *Meloidogyne incognita*, even though infection of wild-type plants induced expression of COMT1, and they were more resistant to the biotrophic oomycete *Hyaloperonospora arabidopsis*, as measured by a significant decrease in asexual sporulation, unstable haustoria, and an increase in oospores (sexual spores). In mutant plants, it was shown that 5-hydroxyferuloyl malate (OH-FM) accumulated to 150 to 200 nmol g⁻¹ fresh tissue whereas it was undetectable in wild-type plants (52). OH-FM stimulated oospore formation in vitro in the nonobligate pathogenic oomycete *Phytophthora cactorum*. These results suggest that resistance in *comt*1 plants may be due to accumulation of OH-FM, and perhaps other metabolites, that significantly affect *H. arabidopsis* but not other pathogens. Thus, we propose that increases in antagonistic metabolites or constitutive expression of defense responses in sorghum *bmr12* grain may have resulted in inhibition of colonization by FIESC no. 1319 that may have been particularly sensitive to these factors.

To summarize, this study showed that *bmr* results in reduced colonization of sorghum grain by members of the *G. fujikuroi*...
species complex, across genetic backgrounds. Also, evidence is provided that a genotype of FIESC is restricted from colonizing bmr12 plants. Lesion lengths following inoculation of sorghum peduncles with *F. verticillioides* or *A. alternata*, as well as *F. thapsinum* (20), were reduced on bmr plants compared with near-isogenic wild-type plants.

**ACKNOWLEDGMENTS**

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**LITERATURE CITED**


