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The Multigeneic *Rhg1* Locus: A Model For The Effects on Root Development, Nematode Resistance and Recombination Suppression.

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Abstract

Soybean (*Glycine max* L. Merr.) resistance to any population (HgType) of *Heterodera glycines* I., the soybean cyst nematode (SCN), requires a functional allele at *rhg1*. An apoptosis-like response in the giant cells formed at the nematode feeding site results about 24-48 h after feeding commences. The response may be mediated by the 3 genes within the *rhg1* locus, a receptor like kinase (RLK), a laccase and an ion anti-porter . This study aimed to identify the role of the genes. Used were near isogeneic lines that contrasted at their *rhg1* alleles(NILs). Five features of the *rhg1* locus, the candidate genes and their nascent proteins were elucidated. First, evidence for a syntenic gene cluster on Lg B1 was found. Second, the effectiveness of SNP probes for distinguishing homeolog sequence variants on LgB1 from alleles at the *rhg1* locus on LgG was shown. Third, analysis of polymorphism among heterozygotes found NIL 34-33 segregating at the *rhg1* locus showed that the resistant allele was dominant in NILs and segregated in phase with a modifier. Fourth, the total absence of recombination events among the NILs between the RLK and other 2 genes eliminated the possibility of a monogeneic *rhg1* locus. Cosegregation of an unlinked locus was detected and a mechanism for segregation distortion inferred. Evidence for the presence transcripts and proteins encoded by the three genes at the *rhg1* locus was shown. Finally, an effect on root development was discovered. A model for multigeneic resistance based on developmental control of root growth is presented.

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Introduction

Soybean (*Glycine max* L. Merr.) seed yield losses due to root infestation by *Heterodera glycines* I. soybean cyst nematode (SCN) have been severe [1]. Losses have occurred since the crop was domesticated, since SCN has been the most widespread and damaging soybean pathogen worldwide. Soybean resistance to SCN was found in about 1% of pre-domesticated and early domesticated Plant Introductions (PI) [2, 3]. The development of partially resistant cultivars by gene introgression and other disease management measures have limited the costs of SCN infestation to the US soybean producers to about $1 billion/year in seed yield losses.

Soybean cyst nematode, like many plant parasitic nematodes, lives as an obligate endoparasite of plant roots that can use many alternate hosts [4, 5] . Over the past 50 years in the US, the number of Hg Types (ex. races) of SCN that are recognized has expanded from 4 in the 1960's to 16–20 of a possible 1,024 to date [6, 7]. Directed breeding for cultivar resistance will continue to select for new Hg types, so new resistance genes and new alleles will be continually needed. Resistant cultivars have a Female Index (FI) of less than 10% of the cyst numbers on susceptible cultivars in parallel tests [2]. Though arbitrary, the 10% measure of resistance often approximates to the economic loss thresh-hold in US soils (about 25 cyst/ 100 cm^3 of soil). Genetic diversity is found both in the field and in the commonly used inbred cyst populations like PA3, Hg type 0. PA3 was derived from a field population by incomplete inbreeding. Therefore, recombination due to sexual reproduction, transposon derived genome plasticity and/or mutation may continue to generate diversity in this population and other populations of the same Hg Type [8].

Variation among the host plant roots response to SCN has been associated with light, temperature and genetic purity such that genetic identities and environmental conditions must be rigorously controlled during host pathogen assays [2,3]. In both resistant and susceptible cultivars, the interaction between the nematode and soybean root passes through several discernable phases [9,10, 11]. However, resistance or susceptibility of the soybean is not induced until females establish a feeding site. The feeding site develops to provide a giant cell with its own secondary root-type vasculature providing nutrients to the female cyst. Cell to cell contact occurs at the syncitia through a stylet sufficiently narrow to prevent the passage of proteins and other molecules greater than 20 kD. Membrane to membrane contact was inferred. Secretions from several glands of SCN contain plant growth regulators and other bioactive factors.

Inheritance of resistance to SCN was first reported in the PI 'Peking' , and three recessive gene symbols (*rhg1–rhg3*) were assigned to the underlying loci [6]. One locus, *rhg1*, provides the major portion of resistance to SCN Hg type 0 (race 3) and Hg type 1.3.5.6.7.8, (race 14) across many genotypes whether they were derived from Peking, 'PI437654', 'PI88788', 'PI209332' or 'PI90763' [12] . However, given the evolution of the soybean nematode interaction is ancient and complex [4] it is likely that the locus contains several genes with each contributing partly to the activity of the locus [13].

The cytological studies suggests the *rhg1* driven Peking-type resistances share mechanisms of giant cell breakdown (pronounced necrosis and cell wall appositions) not seen in PI88788 type resistances in response to Hg type [9]. The differences in the mechanism of giant cell breakdown in Peking and PI88788 may derive from distinct alleles at *rhg1* and/or other defense-associated loci [2,14]. The *rhg1* locus was repeatedly located to a sub-telomeric region of the soybean molecular linkage group G by many studies [12-20]. However, some mapped resistance sources have an *rhg1*-like locus (required for resistance to all races) at another location in SCN resistant PIs, including Lg B1 [21], mid LgG, [22] and LgB2 [23]. Therefore, functional paralogs of *rhg1* may exist among the duplicated regions of the soybean genome [24].

Genes underlying resistance to Hg type 0 (PA3, race 3) have been mapped with greatest accuracy using recombinant inbred lines (RILs) and near isogenic lines (NILs) derived from the cross of 'Essex' by 'Forrest' [13, 24-27]. Forrest provided a unique set of tools for genomics. Forrest introgressed only resistance to Hg type 0 from Peking [28] that also resists two other Hg types. Only *rhg1* and *Rhg4* were introgressed into Forrest [19] . Several NIL populations segregating for *rhg1* and/or *Rhg4* were developed from the cross of Essex by Forrest [13, 29-31]. Genomic analysis identified three genes and their intergeneic regions within the 42 kbp identified as the locus [13]. Within the region the Forrest genes [13] showed with many allelic differences compared to susceptible genotypes 'A3244' [32] and 'Williams 82' (www.phytozome.net) with nine alleles recognized among PIs and four among resistant PIs.

The action of the *rhg1* resistance alleles has complex effects. First, *rhg1* alone was necessary, but not alone sufficient, for resistance to all known Hg-biotypes [12] . Second, some *rhg1* alleles restrict seed germination [33]. Third some interacting alleles are needed prevent zygote death are co-inherited (located on LgM as judged by RFLP markers) [15]. Fourth, some *rhg1* alleles inhibit seed yield at harvest in the absence of the disease [34-36]. Therefore, multiple gene or locus interactions were inferred that both underlie resistance and also alter plant development [13].

The three genes within the markers bounding the *rhg1* locus in Forrest included the RLK, a laccase and a predicted sodium/hydrogen antiporter [13]. Immediately outside the locus (on the basis of recombination events) were two predicted proteins of unknown function. Only the RLK and laccase and the 46.1 Kd predicted proteins were present in EST collections derived from roots. These three genes and their integeneic regions may interact to provide resistance to SCN.

Here, a molecular basis for resistance to SCN is inferred from five features of the *rhg1* locus, the RLK candidate gene and its nascent protein.

Results and Discussion

The soybean genome is hypothesized to be the product of a diploidized tetraploid. Therefore a detailed molecular analysis of the *rhg1* locus required that paralogs and syntenic gene clusters be identified. Probes developed from BAC 73P06 (Figure 1) were used.

Syntenic paralogs of the *rhg1 locus*

To identify *rhg1* paralogs, primers specific to the conserved regions of *rhg1* leucine-rich repeat and *rhg1*- kinase domain were designed. The conserved regions were determined by aligning DNA sequences from known *rhg1*-like genes in different soybean cultivars and other plant species. PCR-amplified products were radiolabeled and used as probes against the Forrest BIBAC libraries. The hybridizing BAC colonies were confirmed by Southern hybridizations to purified DNA. There were five positive clones for the LRR probe (Supplemental Figure 1) and three were also positive from the kinase hybridizations. One of the identified clones (B21d09) contained the *rhg1*, found on scaffold 121 whereas the other three clones contained RLK paralogs (B10a18, B55i16 and H38f23). B10a18 and B55i16 were on Lg A1 scaffold 15; H38f23 was on Lg B1 and scaffold 139 (69,100-144,000). Significantly, this BAC and scaffold 139 contained a complete set of syntenic genes for a second *rhg1* locus(the RLK, laccase both antiporters, the kinase and the helicase; Figure 1). The DNA markers sequence paralogs were present but more diverged except TMD1 that was highly conserved. Sequence analysis of the alleles in Williams 82, 'Asgrow' 3244 and Forrest RLK at *rhg1* showed 99% amino sequence identity. Among the paralogs DNA sequence identity was

high (~92% in geneic region; Supplemental Figure 2). Amino acid identity was 84% in the LRR, 86% in the transmembrane domain and 94% in the kinase domain. The laccase and the antiporter also showed 85-96% amino acid identity.

Both of the RLK paralogs were located by BLAT of microsatellite markers and BES to sequence scaffolds and were in regions where loci with functions similar to *rhg1* were located Lg A1 (the RLK) [21] and LG B1 (the syntenic cluster) [23]. The paralogs may encode proteins that recognize novel race biotypes or substitute for *rhg1* following activation in certain PIs or crosses [21-23].

Allele discrimination

Since paralogs with homeolog sequence variants (HSVs) appeared to exist for each gene in the cluster it was important to distinguish alleles precisely and separately from HSVs. In the NIL population RLK alleles were distinguished using a SNPs from the LRR region (Supplemental Figure 3); and the SIUC-TMD1 marker (Figure 2). In the intergeneic region between the RLK and laccase the SNP probe 10893 was used (Figure 1). Within the laccase was used an indel in the first intron, G10 probe from exon 2-3 and SNP probe 37583 in exon 6. In the anti-porter probe G11 and SNP probe 375821 was used. Each probe could detect polymorphism among the alleles of the three genes at *rhg1* in the ExF derived NILs but not at the paralogous loci.

Dominant, recessive or co-dominant nature of *rhg1*

In NIL 34-33 segregating at the *rhg1* locus, 4 plants were heterozygous at the TMD1 (Figure 2; Table 1) and Sac5 markers (results not shown). The frequency of heterozygousity among F5:13 generation seed was surprisingly high [13,31]. The existence of these plants suggests that fixation is selected against or heterozygosity is selected for, at this locus in these and related NILs.

The cyst scores, for all plants in the NIL population corresponded with the respective alleles at the *rhg1* locus. For the four heterozygous plants, polymorphic at TMD1, the cyst score correspond to those for resistant plants. Therefore, the *rhg1* locus was dominant in this set of NILs infested with this HgType.

Both recessive and co dominant roles have been assigned for the *rhg1* locus. In both past and recent studies with PIs, resistance encoded by *rhg1* was reported as recessive [2,14] whereas previously in NILs, the *rhg1* locus was reported to be co-dominant [19] but without single plant to marker allele associations. Co-dominant and recessive roles of plant disease resistance loci are rare and unusual [37, 38]. However, on the basis of the segregation pattern at the intrageneic TMD1 (intron) and Sca5 (promoter) markers, in NIL 34-33 background, the *rhg1* locus was shown to be dominant (Figure 2; Table 2). The discrepancies in dominance among different populations may be associated with the genetic background the gene resides in and may result from interactions among genes at the *rhg1* locus and/or modifier genes at other loci [15]. The phenotype at the rice blast resistant locus, *Xa3* is also influenced by the genetic background. The gene at the locus behaves differently in different genetic backgrounds, even displaying dominance reversal in one case [39].

Inhibition of root growth by alleles of *rhg1* in the NILs

When counting the cysts with prior knowledge of the allele at *rhg1* it was noted that root mass and vigor appeared to differ among genotypes. Measurements of root mass showed a significant difference among NILs that were associated with the allele at *rhg1* or linked loci (Figure 3; Table 3). Across several experiments, both NILs that were pure breeding susceptible and NILs that segregated some susceptible lines had higher root masses than their SCN resistant counterparts. This phenomenon might underlie the global association of resistance to SCN with low seed germination, seedling vigor, stand formation and ultimately seed yield [33,35].

The recombination events found among the six Hg Type 0 susceptible PIs [13] suggests that the action of *rhg1* requires elements to the distal side of the RLK intron, possibly one or all of the 3 polymorphisms found in the intracellular kinase of the complete RLK. Some mutations in the kinases of other plant RLKs are known to be lethal [40,41]. Kinase mutants can be lethal in many cases [42, 43]. Therefore, it may be the kinase at the *rhg1* locus that underlies restricted root growth in resistant genotypes directly or after some sort of interaction.

The three genes at *rhg1* are expressed in both resistant and susceptible soybean roots

The presence of the RLK at *rhg1* mRNA and protein in roots was confirmed by RT-PCR and Western hybridization (Figure 4). The *rhg1* transcript was detected under both inoculated and non-inoculated conditions in the both the resistant cultivar Forrest and resistant NIL34-23 and the susceptible cultivar Essex and susceptible NIL34-3.

The quantitative PCR used to determine differences in transcript abundance between infected and uninfected cultivars showed the mRNA was increased about 2 fold following SCN inoculation.

Expression judged by examination of EST libraries in silico, cDNA libraries by hybridization and mRNA populations by RT-PCR showed the RLK (Figure 4), laccase [44] and 46.1 Kd hypothetical transporter protein (unpublished) were transcribed in both non-infested roots and SCN-infested roots. However, paralogs were detected for each gene as judged by multiple amplicons from cDNA with laccase probes (G10) [44] and antiporter probes (G11; not shown). Therefore, genes in the syntenic *rhg1* paralog locus might influence *rhg1* activity by cooperation or competition.

Evidence for segregation distortion at *rhg1* from the absence of recombination events

Using the complete set of microsatellite and SNP probes across *rhg1* no recombination events have been found between TMD1 and SIUC Satt75 in the resistant haplotype within the region encompassing the 3 genes at *rhg1*. Used have been the ExF RIL population (n = 100) [19]; the set of SCN resistant PIs (n=112) [13]; NILs with recombination events between Satt309 and Satt214 collected from two populations ExF 34 (2,000) and ExF11 (2,000) [31]; and RILs with recombination events between Satt309 and Satt038RILs collected from the RxH population (n=975) and FxH population (n = 725) [45]. Here, the region from the RLK to the Na H antiporter was analyzed all available markers [13, 42] polymorphic in ExF (Figure 1). Again no recombination events were found. An absence of recombination events in a region can

have one of several causes among them; lack of homology; regional inversions; condensed heterochromatin; and recombinant allele lethality. The first three are not occurring at the *rhg1* locus since DNA sequences are collinear over 100 kbp, even though the locus was an introgression from a PI [13]. Therefore, recombinant allele lethality is the most likely cause.

A hypothetical model for recombinant allele lethality was developed. The kinase domain of the RLK at *Rhg1* was proposed as the killer element and the hypothetical protein was proposed as the target locus kept in the resistant state when the RLK is conferring SCN resistance (Figure 5). The laccase [44] trapped between these two genes and the intergenic regions are held in phase by the locus. It is possible the lethal nature of the resistant linkat is not fully suppressed (leaky) and results in the inhibition of root growth observed (Figure 3).

Conclusions

Suppression of recombination at *rhg1* was shown to center on the three genes at the core of the locus. Whether this causes the unusually high frequency of heterozygous plants or whether the helicase closely linked to the locus [13, 31] has local effects remains to be determined. A suppressor locus acting on *rhg1* was identified earlier [15]. The zygote or embryo lethal gene on Lg M that is co-inherited with *rhg1* [15] proved to have a homeolog that was near Satt594 on Lg G in ExF RILs (Table 1). Mapping the locus to a LG was difficult due to segregation distortion among resistant lines, but considering only susceptible lines carrying the susceptibility allele at *rhg1* the suppressor appears to be in the middle of LgG both in the map and in the sequence

scaffolds. The locus was fixed to the R haplotype in the NILs. Here we propose the locus be named suppressor of *rhg1* or *sup-rhg1.*

The discovery of a syntenic paralog to the gene cluster at *rhg1* raised significant barriers to reverse genetic approaches to the unequivocal proof that the RLK at *rhg1* candidate gene underlies part of the resistance to Hg type 0. Another barrier was the role of *rhg1* in normal plant development that can be inferred from the restricted root growth of NILs (Figure 3), some mutants in this gene (K. Meksem unpublished data) and the shoot effects measured in grafting experiments (A.J. Afzal published data). A third barrier to reverse genetics was the nematodes ability to inhibit RNAi activity (Dr. Chris Taylor unpublished data). Proof of *rhg1* function may require knock-outs of each of the paralogs (K. Meksem et al. unpublished data), or stable transformation to a new location (D. Simmonds and D.A. Lightfoot unpublished data), followed by measurements of genetic segregation. In each case, the analysis will be complicated by the co-dominant nature of the resistance gene in certain backgrounds [4, 20]. In fact, the possibility that the susceptible Essex allele, *rhg1*, promotes the establishment of parasitism by SCN must be explored.

The data presented here suggests the genes linked to the primary candidate RLK may encode factors involved in the modulation of *rhg1* activity. Possible roles include contributions to additive resistance; contributions to resistance in other resistance types (eg PI88788 and Toyohazu; R types 2 and 3) or contributions to the resistance to other Hg types [46]. Unlikely, in view of the susceptibility of segregation events within the interval from the RLK to Satt309 in both PI evolution and NIL segregation, is the hypothesis that the linked genes are factors necessary in

susceptible genotypes for SCN parasitism. The dominance of *rhg1* in NIL segregation also suggests the genes are active in resistant types and inactive in susceptible genotypes. In conclusion, the *rhg1* locus was inferred to be a complex of three genes assembled and co-inherited over long periods of selection for resistance to an pandemic pest, root parasitic nematodes [47].

The probes developed provide a high throughput alternative to satellite markers for marker assisted selection, three allelic discrimination tools were developed for Taqman primer probes (Supplemental Figure 3). The first Taqman probe (1040) could successfully discriminate resistant types 1 and 2 from susceptible haplotypes 2, 3 and 4. Marker 506 could distinguish R types 2 and 3 from other haplotypes. Marker 2050 distinguished among susceptible types. These tools will facilitate molecular breeding.

Materials and Methods

Plant materials

Many of the genetic materials were described previously [13, 29, 31]. Briefly, the seeds of RILs and the NIL populations derived from the cross of Essex by Forrest were obtained from Dr. Paul Gibson at Southern Illinois University at Carbondale in 1995 and were increased from 1995 to present at the Agronomy Research Center [48]. NIL populations were developed and maintained as described in [31]. Genetic identity and purity were checked after increase and before each experiment with 5–10 SSR markers/ line and DNA from 5–10 seeds/line. All lines are available on request as seed. Seeds of NIL 34-23 (resistant haplotype between markers Satt 214 to Satt 570) and NIL 34-3 (susceptible haplotype from the most telomeric marker Satt 214 to the Sat122-Satt 570 interval) were obtained at the F5:13 generation. Genotypes were

rhg1rhg1Rhg4Rhg4 for NIL 34-3 and *Rhg1Rhg1Rhg4Rhg4* for NIL 34-23 whereas NIL34-33 contained both those and *Rhg1rhg1Rhg4rhg4* in different plants.

Near isogenic line populations

Seed of soybean were obtained from the seed store at SIUC managed by Dr. Lightfoot. Seed of NIL 34-33 (polymorphic haplotype between markers Satt 214 to Satt 570) was obtained at the F5:13 generation. The three genotypes found within NIL 34-33 were; *rhg1Rhg1Rhg4Rhg4; rhg1rhg1Rhg4Rhg4* and *Rhg1Rhg1Rhg4Rhg4.*

SCN inoculations

Soybean plants were grown in 5 l buckets, each containing 20 cones in a randomized setup. Each bucket contained a 1:1 ratio of sand soil mix. The containers were placed in a water bath in the SIUC greenhouse. Growth conditions were a 14h light cycle, day time temperature of 30°C and a nighttime temperature of 22°C. The humidity was maintained at approximately 40-50% (v/v). Infection with Hg Type 0 SCN populations consisted of inoculating 2,000 eggs to each 4 day old seedling. Inoculated soybean plants were removed from the cones; 30 days post inoculation and cyst numbers counted.

The NIL experiments used single-plant replications. The cultivars 'Lee 74', 'Essex' and 'Hutcheson' were used as susceptible controls (Niblack et al., 2003). The differentials or indicator lines and the associated female indices (FI) were 'PI54840' (FI 7%), PI 88788 (FI 2%), PI90763 (FI 1%), PI437654 (FI 0%), 'PI 209332' (FI 1%), 'PI89772' (FI 2%) 'PI548316' (FI 8%) and 'PI548402' (FI 3%). Therefore, the standard

differentials showed this HG Type to be 0 [7] (Niblack et al. 2003) corresponding to race 3 [49].

DNA and RNA extraction

DNA was isolated following [50] modified as follows. Briefly, 100 mg of frozen plant tissue was ground, 600 µl preheated (65°C) extraction buffer, incubated at 65°C for 1 hour, cooled and centrifuged at 10,000 g for 15 minutes. The supernatant was decanted and 5 µl of RNase (5 mg/ml) was added at 37°C for one hour. The aqueous phase was extracted first by the addition of equal volume of phenol: chloroform: isoamyl alcohol (25:24:1) v/v, incubated for 2 min by slow inversion followed by centrifugation at 10,000 g for 15 minutes and secondly by the addition of equal volume of chloroform: Isoamyl alcohol (24:1). DNA was collected by the addition of iso-propanol to the supernatant, followed by centrifugation at 12,000 g for 5 min. The DNA pellets were washed twice with 70% (v/v) ethanol, dried and finally dissolved in 30 µl Tris buffer. Concentrations of DNAs were calculated by measuring absorbance at 260 and 280 nm.

Total RNA was isolated with Trizol (Invitrogen Cat. No. 15596-026, Carlsbad, CA, USA), according to the manufacturer's instructions. The RNA pellet was dissolved in 40µl of DEPC treated MQ water and quantified by measuring absorbance at 260 nm. First strand cDNA was synthesis carried out using oligo dT primers using a cDNA synthesis kit, according to manufacturer (Invitrogen). Presence of the *rhg1* mRNA was confirmed by PCR analysis using a *rhg1* intron flanking primers pair: *rhg1*-int-F-LRR (ATT TGA ATC AGA AGT CAG TGT) and *rhg1*-int-R-LRR (TCT GGT CTA ATC TCT TCC AGC; Supplemental Table 2).

NIL genotyping by microsatellite markers linked to *rhg1*

About 50 ng of DNA was used for microsatellite analysis on PAGE after [35]. The microsatellite markers from the RLK, SIUC-TMD1 and SIUC-Sac 5, were used to genotype resistant and susceptible segregants from NIL 34-33 [13] (Supplemental Table 2).   Amplification reactions for BARC-Satt markers for NIL analysis on agarose gels were performed after [51].

Taqman assays of alleles the RLK within the rhg1 locus

The SNP genotyping assay within the gene encoding the RLK was performed using a custom Taqman[TM] Kit. Three probes were designed 1486, 506 and 2040 to distinguish the 8 commonest alleles of the RLK (Supplemental Table 2). Only probe 1486 was polymorphic in Essex Forrest and the derived NILs. A 242 bp amplification reaction was carried out using an *rhg1* LRR forward primer: 5' CAG AGA ACA ACC TCC TTG 3' and an *rhg1* LRR reverse primer: 5' CAG AAC CTG AGA GGC TAT 3'; IDT DNA, Coralville, IA, USA) with the following discriminatory probe pair. Probe 1:5'-Fam-TAT TCC TTC AAG CAT TGC AAA CAT TTC CTC G-BHQ1-3' and Probe 2: 5' Hex -TAT TCC TTC AAG TAT TGC AAA CAT TTC CTC GC-BHQ1-3'. Primer and probe optimization were done by using different combinations of each pair and optimizing to optimal signal strength and balanced fluorophore intensity. The PCR reaction was carried out using a 3 step PCR protocol with one hold at 95°C for 10 minutes followed by 35 cycles that included a denaturation cycle of 95°C for 30 sec, annealing at 58°C for 10 seconds and an extension at 68°C for 20 sec.

SNP assays

Primers for SNPs within the *rhg1* locus on Lg G were used in fine melt curve assays using the ABI7900 with HTM software as described previously [52] with the following modifications. Briefly, genomic DNA was used; multiple amplicon sizes were detected on PAGE gels; melt curve data were normalized by both local (local background value was subtracted from the intensity value of sample) and global metrics. Three SNP primers used were as described in [53]. They were; AX196295 10893 at 54,040 bp between the laccase and RLK; AX196295 37583 CR-G at 62,107 in the laccase; and AX196295 37581 CR-G at 64,929 bp in the hypothetical gene (Supplemental data). Additional primers were designed to detect Essex to Forrest polymorphisms among the three genes and intergeneic regions within the *rhg1* locus.

Southern hybridization

Southern hybridizations were performed following the standard procedure described in [54]. Total genomic DNA was digested with restriction enzymes, separated by electrophoresis on a 0.8% (w/v) agarose gel and transferred onto a positively charged nylon membrane. After hybridization, the corresponding bands were visualized by exposure of X-ray film for 24–48 h.

Total root protein extraction

Protein from root material was isolated from infested and non-infested roots of Forrest and Essex after [55, 56]. Briefly, 2g of the finely ground frozen root was

resuspended in 5mL of Tris buffered phenol (pH 8.8) and 5mL of extraction buffer. The solution was vortexed vigorously and centrifuged at 5,000g for 15 minutes. After removal of the top phase (phenol) the bottom phase was back extracted with Tris buffered phenol (5ml) and an equal volume of the extraction buffer. Proteins were pelleted by centrifugation at 20,000g for 20 minutes and washed. The pellet was dried and resuspended in SDS loading buffer. Total protein concentration was determined using a non-interfering protein assay [57].

SDS–PAGE and Western hybridization

SDS–PAGE of total plant proteins from Essex and Forrest followed by Western hybridization was carried out according to [58] with the following modifications. For the Western hybridizations, a custom made antibody generated against a peptide CTL SRL KTL DIS NNA LNG NLP ATL SNL S from the LRR domain of RHG1 was used (Alpha diagnostics, San Antonio, Texas). As a secondary antibody, an anti rabbit IgG HRP was used (GE healthcare, Milwaukee, Wisconsin).

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Table Legends

Table 1: DNA marker segregation within the SCN resistant lines in the ExF34 derived NIL population lines 1-40 showing some resistance to HgType 0 the phenotype was associated only with segregation near Satt309. Female Index was the mean of 5 plants repeated once (10 plants total). A represents the Essex allele; B the Forrest allele and H identifies lines that are heterogenous and may contain heterozygous plants (eg. NIL34-33).

Table 2: Cyst score and phenotype for plants from Figure 2. The associations between phenotypic scores and marker scores are perfect for all plants. Plant 18 (*) and 19 were switched (**).

Table 3: Association of root growth in NILs with root development in seedlings at 28 days after germination with SCN infestation.

Figure Legends

Figure1: Marker map of the genomic region around *rhg1* and the homeolg of *rhg1* with locus ideograms. Marker anchors are shown for each. Scaffold 139 was anchored by Satt484 at 1,005,915 bp. The B73P06 left BES was homeologous to scaffold 139 at 121,470 bp and homologous to scaffold 121 at 1,675,980 bp. The right BES was at homologous to scaffold 121 at 1,759,500bp (83,520 bp) and weakly homeologous to scaffold 139 at 44,800 bp (76,670 bp). The H38F23 left BES was homeologous to scaffold 139 at 69,150 bp and homologous to scaffold 121 at 1,725,800 bp. The right
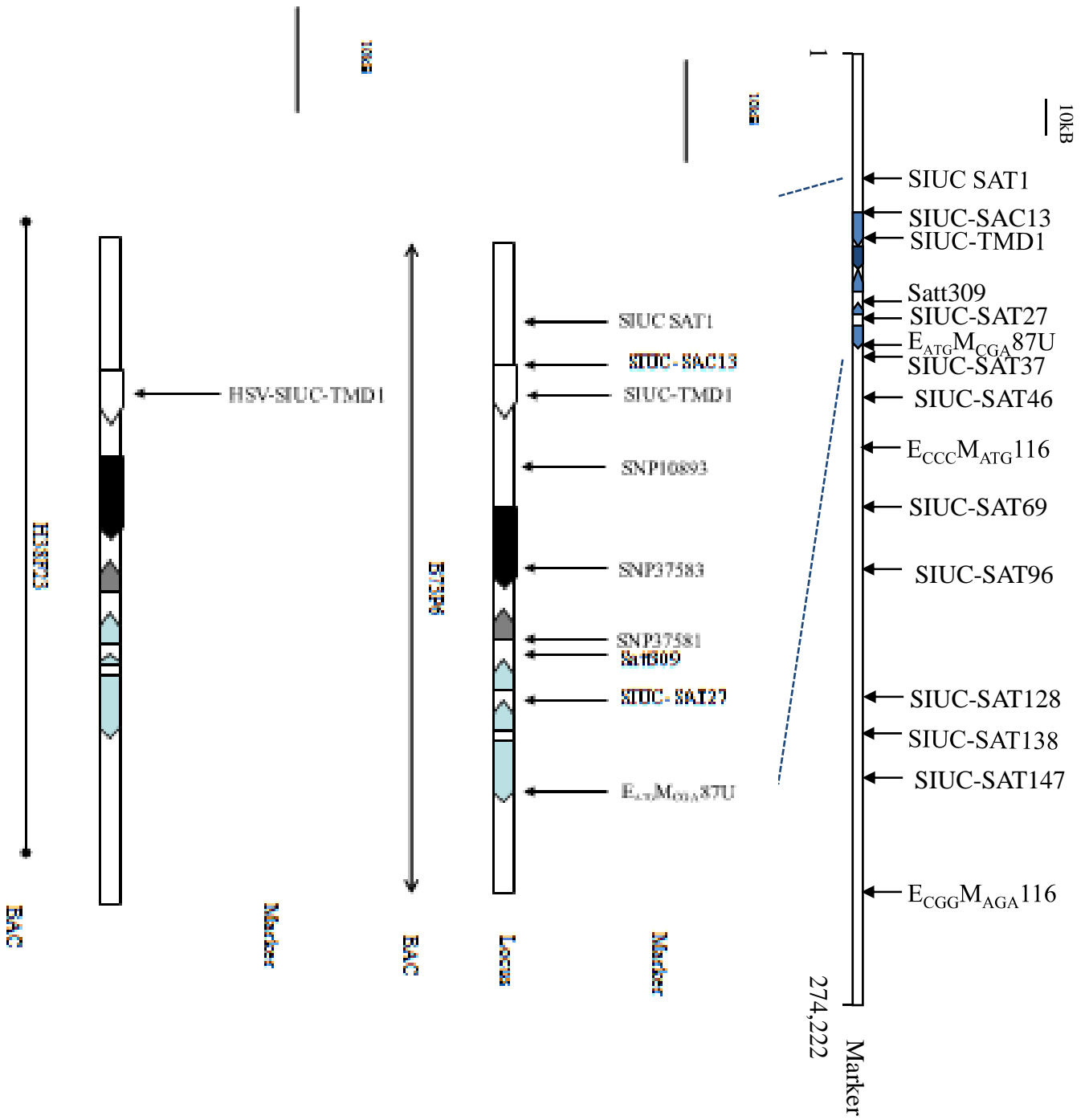
BES was at homologous to scaffold 139 at   144,000 bp (74,850 bp) and weakly
homeologous to scaffold 121 at 1,657,130 bp (68,670 bp). The marker Satt TMD1
(gi:56718383) has a homeolog of 858 bp at 79,000bp on scaffold 139 and a homolog of
762 bp on at 1,714,500 bp on scaffold 121 but the primers are specific to amplify only
the 762$\pm$ 3 bp amplicon. The marker TMD-indel amplified a fragment from both
homeologs (Supplemental Figure 3) of 303$\pm$ 15 bp and 362 bp.
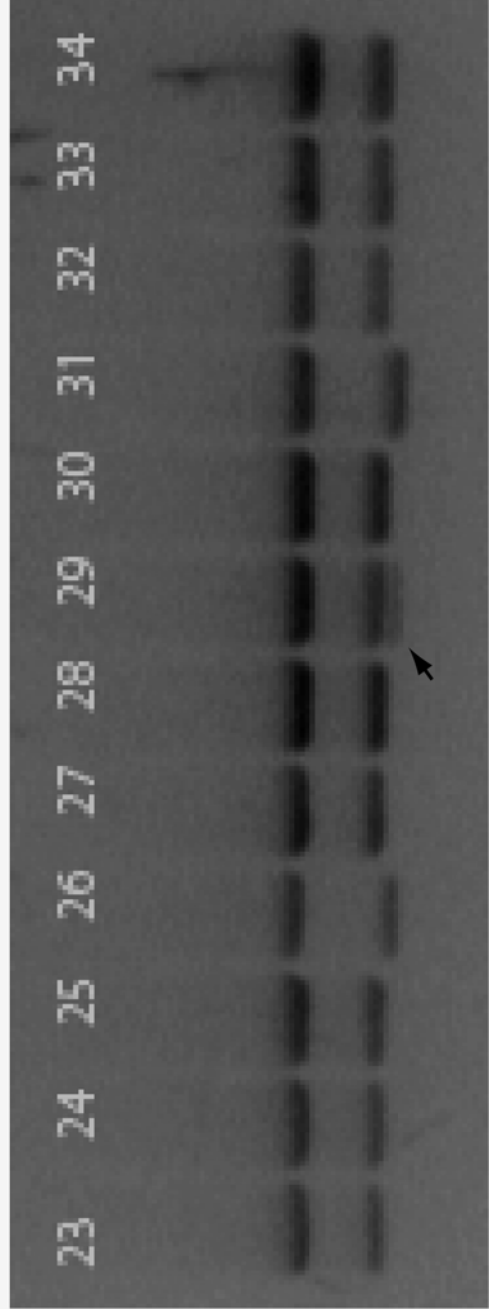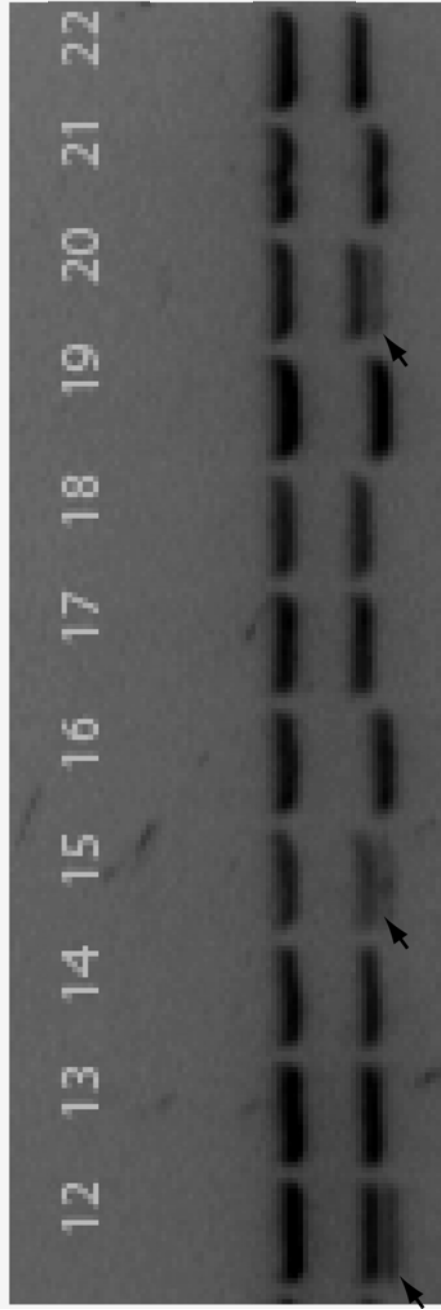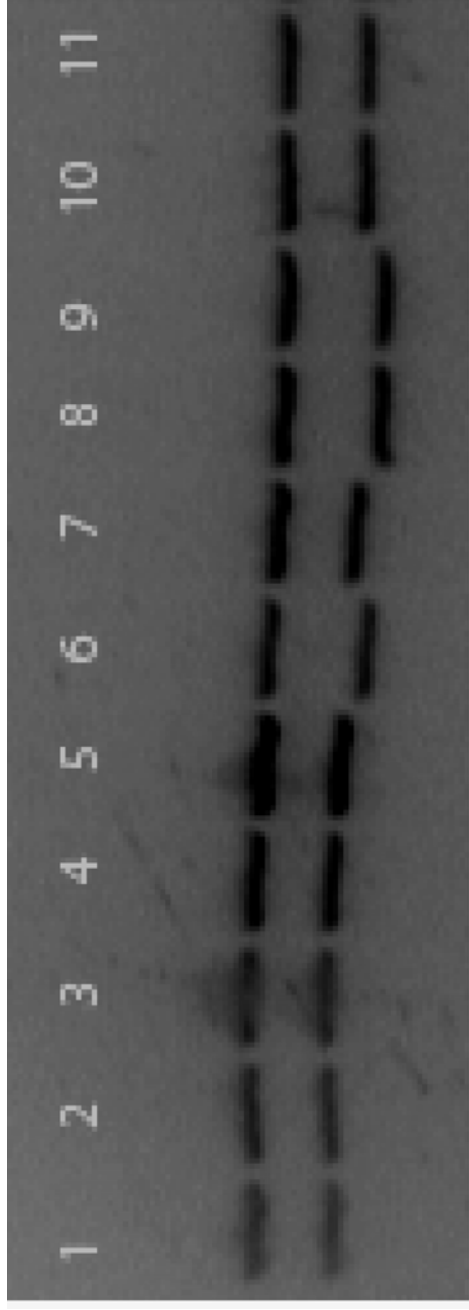
Figure 2: Analysis of NIL 34-33 with TMD1. Out of the 34 plants analyzed, 4 were
heterozygous (Lanes 12, 14, 20 and 29). The four heterozygous lines had a resistant
phenotype. SCN counts and gel scores for the plants are given in Table 3

Figure 3: The RHG1 protein alters root development. Panel A; soybean NILS at 2
weeks pre-SCN inoculation show different root morphologies, post inoculation root
masses are not different (by 6 weeks). Therefore. *rhg1* inhibits germination and early
root growth. Panel B; the root morphologies co-segregated with the allele in the RLK at
the rhg1 locus as shown by the intragenic marker TMD1 (satellite in the intron).

Figure 4: Expression of the RLK at *rhg1* in soybean roots. Panel (A); Western
hybridization using an anti-RHG1 antibody from; Forrest root (a); Essex root (b);
expressed RHG1-LRR-Shrt (c); expressed RHG1-LRR-Long (d); and expressed RHG 4
(e). Panel (B); Agarose gel electrophoresis of cDNA amplified using *rhg1* LRR flanking
primers from RIL 34-23; non-infested control (1); SCN infested RIL 34-23 (2); SCN
infested RIL 34-3 (3); and Forrest genomic DNA (4). Negative control without template
is shown in lane 5.

Figure 5: Model for the function of the *rhg1* locus in resistance to SCN. Black arrows show positive interactions, blue arrows show inhibitions. In this model four phenotypic events are controlled by the 3 genes at *rhg1* and four unlinked genes *Rhg2-4* and *Rzd1* (*sup-Rhg1*). The apoptosis caused in the giant cell of the nematodes feeding sites will occur in plant root cells in the absence of *Rzd1* allele in the coupled phase with the RLK. Root growth inhibition occurs despite suppression of the RLK.
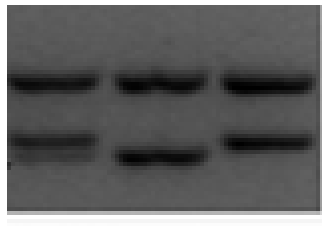
1

10kB

10kB

10kB

SIUC SAT1

SIUC-SAC13

SIUC-TMD1

Satt309

SIUC-SAT27

$E_{ATG}M_{CGA}87U$

SIUC-SAT37

SIUC-SAT46

$E_{CCC}M_{ATG}116$

SIUC-SAT69

SIUC-SAT96

SIUC-SAT128

SIUC-SAT138

SIUC-SAT147

$E_{CGG}M_{AGA}116$

274,222

Marker

HSV-SIUC-TMD1

H46F23

BAC

SIUC SAT1

SIUC-SAC13

SIUC-TMD1

SNP16893

SNP37583

SNP37581

Satt309

SIUC- SAT27

$E_{ATG}M_{CGA}87U$

B73P6

BAC

Locus

Marker

Marker

A.

B.
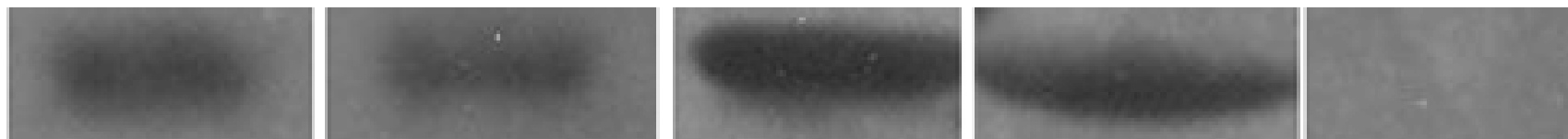
H   S   R

← Homeolog

← Alleles
←

(A)

(a)   (b)   (c)   (d)   (e)

(B)

(1)   (2)   (3)   (4)   (5)

Rhg1

Rhg2

Rhg3

Rhg4

Rzd

Receptor LRR domain

Kinase domain

Laccase

Antiporter

Gene

Nematode detection

Root growth inhibition

Isoflavone production

Vacuole secretion

Apoptosis

Nematode death