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### Resistance to Stem Rust Race TTKSK Maps to the *rpg4/Rpg5* Complex of Chromosome 5H of Barley

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#### **ABSTRACT**

Steffenson, B. J., Jin, Y., Brueggeman, R. S., Kleinhofs, A., and Sun, Y. 2009. Resistance to stem rust race TTKSK maps to the *rpg4/Rpg5* complex of chromosome 5H of barley. Phytopathology 99:1135-1141.

Race TTKSK (Ug99) of the wheat stem rust pathogen (*Puccinia graminis* f. sp. *tritici*) is a serious threat to both wheat and barley production worldwide because of its wide virulence on many cultivars and rapid spread from eastern Africa. Line Q21861 is one of the most resistant barleys known to this race. To elucidate the genetics of resistance in this line, we evaluated the Q21861/SM89010 (Q/SM) doubled-haploid population for reaction to race TTKSK at the seedling stage. Segregation for resistance:susceptibility in Q/SM doubled-haploid lines fit a 1:1 ratio (58:71 with  $\chi^2 = 1.31$  and P = 0.25), indicating that a single gene in Q21861 confers resistance to race TTKSK. In previous studies, a recessive gene (rpg4) and a partially dominant gene (Rpg5)

were reported to control resistance to *P. graminis* f. sp. *tritici* race QCCJ and *P. graminis* f. sp. *secalis* isolate 92-MN-90, respectively, in Q21861. These resistance genes co-segregate with each other in the Q/SM population and were mapped to the long arm of chromosome 5H. Resistance to race TTKSK also co-segregated with resistance to both rusts, indicating that the gene conferring resistance to race TTKSK also lies at the *rpg4/Rpg5* locus. This result was confirmed through the molecular analysis of recombinants previously used to characterize loci conferring resistance to race QCCJ and isolate 92-MN-90. The 70-kb region contains *Rpg5* (a nucleotide-binding site leucine-rich repeat serine/threonine-protein kinase gene), *rpg4* (an actin depolymerizing factor-like gene), and two other genes of unidentified function. Research is underway to resolve which of the genes are required for conferring resistance to race TTKSK. Regardless, the simple inheritance should make Q21861 a valuable source of TTKSK resistance in barley breeding programs.

Barley is an important crop in the Upper Midwest region of the United States and Prairie Provinces of Canada, with >4.8 million ha produced annually (13). Although stem rust (caused by Puccinia graminis f. sp. tritici Eriks. & Henn. and P. graminis f. sp. secalis Eriks. & Henn.) has historically been a problem on the crop in the past, it has not caused any widespread epidemics for over 70 years (27,40). Since 1942, effective control of stem rust has been achieved through the deployment of cultivars with the durable resistance gene Rpg1 but other factors, such as a largely resistant wheat crop and barley's shorter maturation period coupled with the late arrival of rust inoculum to the northern planted crop, likely contributed to the long-lasting disease control (33). Rpg1 is effective against many races of P. graminis f. sp. tritici; however, some with virulence for this gene have been reported on cultivars from the field (33). One of the most recent cases involved QCCJ, a race thought to have originated in western Canada in the late 1980s (21,33). Race QCCJ became established in the Great Plains region in 1989 and damaged some barley fields planted to cultivars carrying Rpg1 in both the United States and Canada (33). It has since become a minor component of the stem rust population in the United States and Canada after susceptible wheat cultivars were no longer grown in the central Great Plains. A new threat to barley is TTKSK (original described race designation: TTKS with isolate synonym of Ug99), a race first characterized from Uganda in 1999 (26). Race TTKSK has a wide virulence spectrum and is capable of attacking >70% of the

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world's wheat cultivars (30). Recent data also indicate that it is virulent for many barley cultivars, including those bred with Rpg1 (34,36). Race TTKSK represents a serious threat to wheat and barley production worldwide because of its wide virulence, ability to cause severe yield loss (8), and rapid spread from Uganda. Since its initial discovery in 1999 (26), TTKSK has spread to other countries in eastern Africa (Kenya, Ethiopia, and Sudan), crossed the Red Sea into Yemen in 2006, and is now confirmed in Iran (23,30,42). It may only be a short time before it spreads into the vast cereal production areas of southern Asia and beyond. In the short term, fungicides offer the best control solution where no resistant cultivars are available; however, this strategy is prohibitively expensive for many of the region's subsistence growers and raises some environmental concerns. The deployment of resistant cereal crops is the most effective strategy for combating the threat posed by race TTKSK. A global effort to breed wheat for resistance to race TTKSK was launched after an international expert panel convened to address the problem in 2005. Through this initiative, now called the Borlaug Global Rust Initiative, significant progress has been made in identifying new and native sources of stem rust resistance and transferring the resistance genes into adapted wheat cultivars (1,30,31). Comparatively little research has been advanced on barley, due to its smaller worldwide acreage and lower rank as a food crop for people. Yet, in the very countries where TTKSK now exists (Ethiopia and Yemen), barley is one of the most important food crops, especially for people living at high elevation (12). Wheat and barley are often cultivated in the same areas. To effectively control stem rust, it is essential that both crops be bred for resistance, otherwise inoculum from one crop could initiate epidemics on the other crop. Furthermore, the effective population size of P. graminis f. sp. *tritici* on both crops must be held in check, otherwise the chance for dangerous mutants to arise will increase (28).

In barley, at least eight genes for resistance to stem rust have been described. Those identified on the basis of their reaction to races of the wheat stem rust pathogen, P. graminis f. sp. tritici, include Rpg1 derived from cvs. Peatland (CIho 5267), Chevron (CIho 1111), and Kindred (CIho 6969) (25,29,33); Rpg2 from Hietpas 5 (CIho 7124) (24); Rpg3 from PI 382313 (14,15); rpg4 from Q21861 (PI 584766) (17); and Rpg6 from 212Y1, a barley line with a Hordeum bulbosum introgression (9). Another gene, aside from RpgI, was identified in Peatland (designated as RpgU) and confers moderate resistance to race QCCJ in the field (10). Three genes have been identified on the basis of their reaction to the rye stem rust pathogen, P. graminis f. sp. secalis. A single dominant gene for resistance was identified in the barley accession Skinless (20), and a single recessive gene (rpgBH, formerly designated S gene) was found in Black Hulless (CIho 666) (37). Additionally, a partially dominant gene for resistance to P. graminis f. sp. secalis was reported in Q21861 by Sun et al. (39) and later provisionally designated as RpgQ (38). This gene was thought to be the same as rpg4 because the resistance conferred by RpgQ to rye stem rust co-segregated with resistance to wheat stem rust race QCCJ conferred by rpg4 (39). However, recent high-resolution mapping efforts have shown that RpgQ is a different but closely linked gene to rpg4 on the long arm of chromosome 5H and was given the new locus symbol of Rpg5 (3).

In barley, research is urgently needed on identifying sources of resistance to race TTKSK and transferring resistance genes into adapted breeding lines. Preliminary evaluations of *Hordeum* germplasm have identified several sources of resistance to race TTKSK in various cultivars, landraces, and wild barley accessions (34,36). Breeding line Q21861 was found to be one of the most resistant barley lines to race TTKSK at both the seedling stage in greenhouse and at the adult plant stage in the field (B. Steffenson and Y. Jin, *unpublished*). Information on the genetics of TTKSK resistance in Q21861 is important for developing the most efficient strategy for stem rust resistance breeding. Thus, the objective of this study was to determine the number and chromosomal location of genes controlling resistance to race TTKSK in Q21861.

#### MATERIALS AND METHODS

**Plant materials.** The doubled-haploid (DH) population of Q21861/SM89010 (Q/SM) was used for this study. It was produced by anther culture from  $F_1$  plants of the two parents and consists of 129 lines (35). Q21861 is a two-rowed breeding line of unknown parentage that was originally selected in 1983 from a breeding nursery established at Centro Internacional de Mejoramiento de Maíz y Trigo in Mexico. It carries the stem rust resistance genes Rpg1, rpg4, and Rpg5 (3,17,35) as well as other disease resistance genes (11,35). SM89010 is a two-rowed malting barley line from the University of Saskatchewan breeding program. It was derived from the cross Nairn × Manley and is susceptible to stem rust.

Plant growth conditions, inoculation protocol, and infection/incubation period. Evaluations of the Q/SM population for resistance to race TTKSK were done at the United States Department of Agriculture–Agricultural Research Service Cereal Disease Laboratory during the winter months. Three to five seeds from each DH line and the parents were planted in plastic cones (21 cm in depth and 3.8 cm in diameter) filled with a 50:50 mixture of soil and Metro Mix 200 (vermiculite, peat moss, perlite, and sand) and fertilized with Osmocote 14-14-14 (Scott's Company, Marysville, OH) (1.4 g/cone) and Peters Dark Weather 15-0-15 (Scott's Company) (150 g per 3.78 liter at 1/16 dilution). Plants were grown in a greenhouse at 22°C with supplemental lighting provided by 1,000-W sodium vapor lamps for 14 h per

day. Isolate 04KEN156/04 of race TTKSK was initially increased on a susceptible wheat host, collected, desiccated, and stored in tubes at -80°C until needed. On the day of inoculation, the tubes of rust were heat shocked at 45°C for 10 min, rehydrated in 80% KOH, placed in gelatin capsules containing a lightweight mineral oil carrier (Soltrol 170; Phillips Petroleum, Bartlesville, OK), and applied to 9-day-old plants with fully expanded primary leaves. The concentration of inoculum used was 4.28 mg/ml of lightweight mineral oil. Inoculation protocols were as described by Sun and Steffenson (38), except that 0.004 mg of rust was applied per plant instead of 0.035 mg. After inoculation, plants were placed in mist chambers at 21°C for 16 h in the dark. During this time, the plants were kept moist by periodic misting from ultrasonic humidifiers. Then, light was provided by fluorescent bulbs, and the chamber doors were partially opened to reduce the build up of heat. After 3 to 4 h, the leaf surfaces were completely dry, and plants were returned to the greenhouse under the conditions described above.

The experiment was planted in a completely randomized design with one replicate and was repeated once. Three to five plants were assessed per replicate. A few DH lines gave variable reactions between experiments and were tested again in two additional experiments.

**Disease assessment.** Fourteen to seventeen days after inoculation, the infection types (ITs) on each DH line and the parents were assessed using a 0 to 4 scale. The IT scale used for barley is a modification of the one developed for wheat by Stakman et al. (32) and is based primarily on uredinial size as described by Miller and Lambert (22). Barley frequently exhibits two or more ITs on a single leaf (i.e., a mesothetic reaction) when infected with P. graminis (38); thus, all of the ITs observed on each DH line were recorded in order of their prevalence. To simplify the presentation of data, the two most common ITs (i.e., the IT mode) are presented in Table 1. Comparisons were made on the basis of the IT mode because the two most common ITs usually comprised >85% of all those observed on individual DH lines. Additionally, the lowest and highest ITs (i.e., the IT range) observed also are given to indicate the degree of variation in stem rust reactions. Classification of progeny into resistant and susceptible categories was based on the reaction of the parents to race TTKSK. The genetics of resistance to both P. graminis f. sp. tritici (races QCCJ and MCCF) and P. graminis f. sp. secalis (isolate 92-MN-90) was previously investigated in the Q/SM population (35,39). Raw, unpublished IT data from these studies are presented in Table 1. These IT data were compared with those collected in this study, by inspection, to race TTKSK to assess the agreement of general reaction phenotypes (i.e., resistance versus susceptibility).

Genetic and physical mapping. Restriction fragment length polymorphism (RFLP) markers were developed for the Q/SM population as described by Druka et al. (6). Sequence tag site (STS) markers were developed using the Q21861 genomic sequence from the rpg4/Rpg5 region (accession no. EU878778) (3). Regions with polymorphism between Q21861 and SM89010 were utilized for polymerase chain reaction (PCR)-based marker development. The simple sequence repeat (SSR) markers were developed from cv. Morex bacterial artificial chromosome (BAC) sequence (accession no. EU812563). Primer sequences for STS and SSR markers are shown in Table 2. A single nucleotide polymorphism (SNP) was scored in the RB SSR1 amplicon by sequencing the PCR product from the Q/SM recombinants. The RB SSR1 amplicon length is identical between Q21861 and SM89010; however, SM89010 contains an additional CAA repeat and is missing one CAT repeat compared with Q21861. Molecular analysis of the rpg4/Rpg5 region used seven recombinants from Q/SM based on 129 gametes within the flanking region defined by RFLP markers ABG391 and MWG740. BAC physical maps were developed according to Druka et al. (6) and Brueggeman et al. (3).

TABLE 1. Infection type mode (IT-M) and range (IT-R) for doubled-haploid (DH) progeny from the Q21861/SM89010 population to *Puccinia graminis* f. sp. *tritici* races TTKSK, QCCJ, and MCCF and *P. graminis* f. sp. *secalis* isolate 92-MN-90 at the seedling stage<sup>a</sup>

	TTKSK			92-MN-90			QCCJ			MCCF		
H line	IT-M	IT-R	Allele	IT-M	IT-R	Allele	IT-M	IT-R	Allele	IT-M	IT-R	Alle
21861	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	0;1-	0; to 1	Rpg1
189010	32	1 to 3	Rpg4/rpg5	23-	2 to 3–	rpg5	33–	2 to 3+	Rpg4	3–3	3– to 3	rpg1
M001 _	33+	3 to 3+	<i>Rpg4/rpg5</i>	3–32	1 to 3	rpg5	33–	2 to 3+	Rpg4	10;	0; to 2	Rpg1
M002	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	3–3	3– to 3	rpg1
M003	0;	0;	rpg4/Rpg5	0;1	0 to 1	Rpg5	0;1	0 to 1	rpg4	33–	3- to $3$	rpg1
M004 _	3	3	Rpg4/rpg5	23-	2 to 3–	rpg5	33–	2 to 3+	Rpg4	33–	3– to 3+	rpg1
M005	0;1-	0; to 1–	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	0;1–	0 to 1–	Rpg
M008	0;1–	0; to 1–	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	0;1–	0; to 1–	Rpg.
M010	3–2	2 to 3–	Rpg4/rpg5	213-	1 to 3–	rpg5	3–3	2 to 3	Rpg4	0;1	0; to 1	Rpg.
M011	32	2 to 3+	Rpg4/rpg5	213–	1 to 3–	rpg5	33–	2 to 3+	Rpg4	33-	3– to 3	rpg1
M012	3	3– to 3	Rpg4/rpg5	3–2	2 to 3–	rpg5	33–	2 to 3+	Rpg4	12-	1 to 2–	Rpg
M013	3+	3+	Rpg4/rpg5	3–2	1 to 3–	rpg5	33+	3 to 3+	Rpg4	21	1 to 3–	Rpg.
M014	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0 to 2	rpg4	3–3	3– to 3	rpgI
M015	0;1-	0; to 1–	rpg4/Rpg5	0;1	0; to 1	Rpg5	0;1	0 to 2	rpg4	33–	2 to 3	rpg1
M016	0;1-	0; to 1–	rpg4/Rpg5	0;1	0; to 1	Rpg5	00;	0 to 0;	rpg4	3–3	3– to 3	rpg1
M017	3–	3-	Rpg4/rpg5	213-	1 to 3	rpg5	3–2	2 to 3	Rpg4	33–	3– to 3+	rpg1
M018	0;1	0; to 1	rpg4/Rpg5	0;1	0; to 1	Rpg5	0;	0 to 1	rpg4	3–3	3- to $3$	rpg1
M019 _	3	3– to 3	Rpg4/rpg5	3–2	2 to 3–	rpg5	33–	3– to 3+	Rpg4	33–	3- to $3+$	rpg1
M020	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	33–	3- to $3$	rpg1
M023	3	3	Rpg4/rpg5	3–2	2 to 3	rpg5	33+	3– to 3+	Rpg4	33–	3– to 3	rpg1
M024	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 0;	rpg4	10;	0; to 2	Rpg.
M026	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 0;	rpg4	0;1–	0; to 1–	Rpg.
M028 _	3	3	Rpg4/rpg5	3–2	2 to 3–	rpg5	33+	3– to 3+	Rpg4	0;1-	0; to 1–	Rpg.
M029	10;	0; to 1	rpg4/Rpg5	0;1	0; to 2	Rpg5	0;1	0 to 2	rpg4	33-	2 to 3	rpg1
M030	3+	3– to 3+	<i>Rpg4/rpg5</i>	23-1	1 to 3	rpg5	33–	2 to 3+	Rpg4	0;1	0; to 1	Rpg.
M031 _	33+	3 to 3+	<i>Rpg4/rpg5</i>	213-	0; to 3–	rpg5	33+	2 to 3+	Rpg4	33-	3- to $3$	rpg1
M033	0;1	0; to 1	rpg4/Rpg5	00;	0 to 1–	Rpg5	0;1	0 to 2	rpg4	3–3	3- to $3$	rpg1
M034	3	3	Rpg4/rpg5	23-1	1 to 3–	rpg5	33+	3– to 3+	Rpg4	33-	3- to $3+$	rpg1
M035	3	3	Rpg4/rpg5	213-	1 to 3-	rpg5	3–3	2 to 3+	Rpg4	3–3	3– to 3	rpg1
M036	3	3	Rpg4/rpg5	23-1	1 to 3-	rpg5	33-	2 to 3+	Rpg4	33-	3– to 3	rpg1
M037	0;1-	0; to 1–	rpg4/Rpg5	0;1	0 to 1	Rpg5	0;1	0; to 2	rpg4	33-	3– to 3	rpg1
M038	3	3	Rpg4/rpg5	3-21	1 to 3-	rpg5	33-	3- to $3+$	Rpg4	33-	3– to 3	rpg1
M039	0;1-	0; to 1–	rpg4/Rpg5	0;1	0; to 1	Rpg5	00;	0 to 1-	rpg4	0;1-	0; to 1–	Rpg.
M041	0;1-	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 0;	rpg4	10;	0; to 2–	Rpg.
M042	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	3–3	2 to 3	rpg1
M043	33-	3– to 3	Rpg4/rpg5	23-1	1 to 3-	rpg5	33-	2 to 3+	Rpg4	33-	3- to $3+$	rpg1
M045	3	3- to 3+	Rpg4/rpg5	23-1	1 to 3-	rpg5	33+	3- to 3+	Rpg4	12	1 to 2	Rpg.
M046	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	0;1	0; to 2–	Rpg.
M047	33+	3 to 3+	Rpg4/rpg5	3–2	2 to 3-	rpg5	33+	2 to 3+	Rpg4	23-	1 to 3-	Rpg
M048	33+	3 to 3+	Rpg4/rpg5	21	1 to 2	rpg5	33+	2 to 3+	Rpg4	33-	3- to 3	rpg1
M049	0;1-	0; to 1–	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0 to 1	rpg4	3–3	3– to 3	rpg1
M050	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 0;1	rpg4	33-	3- to $3$	rpg1
M052	0;1-	0; to 1–	rpg4/Rpg5	00;	0 to 1-	Rpg5	00;	0 to 0;	rpg4	3–3	2 to 3	rpg1
M054	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	0;1	0; to 2–	Rpg.
M055	33-	3– to 3	Rpg4/rpg5	213-	1 to 3-	rpg5	33-	2 to 3+	Rpg4	0;1	0; to 1	Rpg.
M056	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	33-	3– to 3	rpg1
M057	10;	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	3–3	3– to 3	rpg1
M058	33+	3 to 3+	Rpg4/rpg5	3–3	3– to 3	rpg5	33-	2 to 3+	Rpg4	21	1 to 3–	Rpg.
M059	33-	3– to 3	Rpg4/rpg5	23-1	1 to 3	rpg5	33+	2 to 3+	Rpg4	33-	3– to 3	rpg1
M060	3–	3–	Rpg4/rpg5	23-1	1 to 3-	rpg5	33-	2 to 3+	Rpg4	33-	3– to 3	rpg1
M061	3	3– to 3+	Rpg4/rpg5	3-23	1 to 3	rpg5	33+	2 to 3+	Rpg4	33–	2 to 3	rpg1
M062	3	3- to 3+	Rpg4/rpg5	213-	1 to 3-	rpg5	33+	3- to 3+	Rpg4	12	1 to 3–	Rpg.
M063	3	2+ to 3	Rpg4/rpg5	23-1	1 to 3-	rpg5	33-	3- to 3+	Rpg4	0;1	0; to 1	Rpg.
M065	3+	3+	Rpg4/rpg5	3–2	2 to 3–	rpg5	3+3	3 to 3+	Rpg4	0;1	0; to 2	Rpg.
M066	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 1	rpg4	0;1	0; to 1	Rpg.
M067	10;	0; to 2	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	33-	3– to 3	rpgl
M068	3	3	Rpg4/rpg5	23–1	1 to 3–	rpg5	33–	2 to 3+	Rpg4	33–	3– to 3	rpg1
M069	3	3	Rpg4/rpg5	3–3	2 to 3	rpg5	33+	3- to 3+	Rpg4	10;	0; to 2–	Rpg.
M070	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 2	rpg4	33-	3– to 3	rpgl
M071	3–	3–	Rpg4/rpg5	23–1	1 to 3–	rpg5	33–	2 to 3	Rpg4	0;1	0; to 1	Rpg.
M073	3	3	Rpg4/rpg5 Rpg4/rpg5	3–2	2 to 3–	rpg5 rpg5	33 <del>-</del>	3 to 3+	Rpg4 Rpg4	21	1 to 3–	Rpg.
M074	3	3 to 3+		3–2 3–21			33+ 33-	2 to 3+		21	1 to 3– 1 to 2	
	0;1	0; to 2–	Rpg4/rpg5		1 to 3–	rpg5	21	0; to 3–	Rpg4	33-	3– to 3	Rpg
M075		•	rpg4/Rpg5	0;1-	0 to 1-	Rpg5		•	rpg4			rpg1
M076	3	3	Rpg4/rpg5	3–21	0; to 3–	rpg5	33-	2 to 3+	Rpg4	3–2	2 to 3	Rpg
M077 M078	3 33–	3	Rpg4/rpg5	213-	1 to 3–	rpg5	33+	2 to 3+	Rpg4	33–	3– to 3	rpgl
	4.4_	3– to 3	<i>Rpg4/rpg5</i>	23-1	1 to 3–	rpg5	3–3	2 to 3	Rpg4	3–3	3- to $3+$	rpg1

<sup>&</sup>lt;sup>a</sup> ITs are based on the 0 to 4 scale of Stakman et al. (32) as modified for barley by Miller and Lambert (22). The two (or three in the case of 92-MN-90) most common ITs observed on accessions (i.e., IT-M) are given in addition to the lowest and highest types observed (i.e., IT-R). Allele = recognized allele. The raw IT data for *P. graminis* f. sp. *secalis* isolate 92-MN-90 and the two *P. graminis* f. sp. *tritici* races (QCCJ and MCCF) have not been previously published, but are from the studies of Sun et al. (39) and Steffenson et al. (35), respectively. Interactions highlighted in gray indicate the presence of the respective resistance alleles in the QSM line.

#### RESULTS AND DISCUSSION

Q21861 was highly resistant to race TTKSK and consistently exhibited a very low IT mode of 0; (Table 1). In contrast, SM89010 was susceptible, exhibiting an IT mode of 32. Q/SM DH progeny could be easily classified into two distinct

groups based on their IT mode. The resistant group included 58 progeny with IT modes from 0; to 0;1 (rarely 10; and 12), and the susceptible group included 71 progeny with IT modes from 3– to 3+ and combinations thereof (rarely 23– and 3–2). The 58:71 segregation for resistance:susceptibility in the Q/SM DH population fit a 1:1 ratio ( $\chi^2 = 1.31$ , P = 0.25),

 $TABLE\ 1.\ (continued\ from\ preceding\ page)$ 

	TTKSK			92-MN-90			<b>QCC</b> J			MCCF		
DH line	IT-M	IT-R	Allele	IT-M	IT-R	Allele	IT-M	IT-R	Allele	IT-M	IT-R	Allele
QSM079	3	2 to 3+	Rpg4/rpg5	3-23	1 to 3–	rpg5	33+	2 to 3+	Rpg4	33-	3- to 3+	rpg1
QSM080	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 0;	rpg4	0;1-	0; to 1–	Rpg1
QSM082	3	3	Rpg4/rpg5	213–	1 to 3–	rpg5	33+	3- to 3+	Rpg4	33-	3- to 3+	rpg1
QSM083 QSM084	3 33–	3– to 3+ 3– to 3	Rpg4/rpg5 Rpg4/rpg5	3–3 12	2 to 3– 0; to 2	rpg5	3–3 3–2	2 to 3+ 2 to 3-	Rpg4	10; 0;1–	0; to 2– 0; to 1	Rpg1
QSM085	33–	3– to 3	Rpg4/rpg5	33–	2 to 3	rpg5 rpg5	3 <del>-</del> 2	2 to 3=	Rpg4 Rpg4	0;1-	0; to 1	Rpg1 Rpg1
QSM086	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	33–	3– to 3	rpg1
QSM087	0;1	0; to 1	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0 to 2	rpg4	33–	3– to 3	rpg1
QSM088	3	3– to 3	Rpg4/rpg5	213-	1 to 3-	rpg5	3–3	2 to 3+	Rpg4	3–3	3– to 3	rpg1
QSM089	0;1-	0; to 1–	rpg4/Rpg5	0;1-	0 to 1–	Rpg5	00;	0 to 2–	rpg4	12-	0; to 2	Rpg1
QSM090	23–	2 to 3–	Rpg4/rpg5	23–1	1 to 3–	rpg5	33+	3- to 3+	Rpg4	3–3	3– to 3	rpg1
QSM091 QSM093	3 0;	3 0;	Rpg4/rpg5 rpg4/Rpg5	3–2	2 to 3– 0 to 0;	rpg5 Rpg5	33– 00;	2 to 3+ 0 to 1	Rpg4 rpg4	3–3	1 to 3 2 to 3	Rpg1
QSM093 QSM094	33+	3 to 3+	Rpg4/rpg5	3–21	1 to 3–	rpg5	33–	2 to 3+	Rpg4	10;	0; to 1	rpg1 Rpg1
QSM095	0;1-	0; to 1–	rpg4/Rpg5	00;	0 to 0;	Rpg5	0;1	0; to 1	rpg4	0;1-	0; to 1	Rpg1
QSM097	0;1	0; to 1	rpg4/Rpg5	0;	0; to 1–	Rpg5	00;	0 to 1–	rpg4	0;1-	0; to 1–	Rpg1
QSM098	0;1	0; to 1	rpg4/Rpg5	0;1	0; to 1	Rpg5	00;	0 to 1	rpg4	0;1	0; to 1–	Rpg1
QSM099	3–	3–	Rpg4/rpg5	3–3	2 to 3	rpg5	33–	2 to 3+	Rpg4	33-	3– to 3	rpg1
QSM100	0;1-	0; to 1–	rpg4/Rpg5	0;1	0; to 1	Rpg5	00;	0 to 1	rpg4	0;1-	0; to 1	Rpg1
QSM101 QSM103	0;1 0;1	0; to 1 0; to 1	rpg4/Rpg5 rpg4/Rpg5	0;1 0;1	0; to 1 0 to 1	Rpg5 Rpg5	00; 0;1	0 to 1 0; to 2	rpg4	0;1– 10;	0; to 1 0; to 2–	Rpgl
QSM103 QSM104	3–	3–	Rpg4/rpg5	3–2	2 to 3–	rpg5	33–	2 to 3+	rpg4 Rpg4	33-	3– to 3	Rpg1 rpg1
QSM105	33–	2 to 3	Rpg4/rpg5	213-	1 to 3–	rpg5	3–2	2 to 3+	Rpg4	3–3	3– to 3	rpg1
QSM106	0;1	0; to 1	rpg4/Rpg5	0;1	0; to 1	Rpg5	00;	0 to 1–	rpg4	0;1-	0; to 1–	Rpg1
QSM107	0;	0;	rpg4/Rpg5	0;1	0; to 1	Rpg5	0;1	0 to 2	rpg4	3–3	3– to 3	rpg1
QSM108	33–	3– to 3	Rpg4/rpg5	3–21	1 to 3–	rpg5	3–2	2 to 3+	Rpg4	3–3	3– to 3	rpg1
QSM109	3+	3+	Rpg4/rpg5	3–21	1 to 3–	rpg5	3–3	2 to 3+	Rpg4	3–3	3– to 3	rpgl
QSM110 QSM111	3– 33–	2 to 3 3– to 3	Rpg4/rpg5	10;23– 3–	0; to 3– 2 to 3	rpg5	3–2 33–	1 to 3 2 to 3+	Rpg4	3–3 33–	3– to 3 3– to 3	rpg1
QSM111 QSM112	0;1	0; to 1	Rpg4/rpg5 rpg4/Rpg5	00;	0 to 0;	rpg5 Rpg5	0;1	0 to 2	Rpg4 rpg4	0;1	0; to 1	rpg1 Rpg1
QSM113	0;1	0; to 2	rpg4/Rpg5	0;1	0; to 2	Rpg5	0;1	0; to 2	rpg4	0;1	0; to 1	Rpg1
QSM114	33–	3– to 3	Rpg4/rpg5	33–	2 to 3	rpg5	33–	2 to 3+	Rpg4	3–3	3– to 3	rpg1
QSM115	0;1	0; to 2	rpg4/Rpg5	00;	0 to 1	Rpg5	00;	0 to 2	rpg4	33–	3– to 3	rpg1
QSM116	12	0; to 2	rpg4/Rpg5	0;1	0 to 1	Rpg5	0;1	0; to 2	rpg4	3–3	3– to 3	rpg1
QSM117	0;1-	0; to 1–	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	3–3	3– to 3	rpg1
QSM118 QSM119	3+ 0;1	3+ 0; to 2	Rpg4/rpg5 rpg4/Rpg5	3–3	3– to 3 0;	rpg5 Rpg5	3 0;1	2 to 3+ 0; to 3-	Rpg4 rpg4	3–3 33–	3– to 3+ 3– to 3	rpg1 rpg1
QSM110 QSM120	3	3- to 3+	Rpg4/rpg5	3–21	1 to 3–	rpg5	3–3	2 to 3	Rpg4	0;1	0; to 1	Rpg1
QSM121	33+	3 to 3+	Rpg4/rpg5	23–1	1 to 3	rpg5	33–	2 to 3	Rpg4	33–	3– to 3	rpg1
QSM122	33-	3– to 3	Rpg4/rpg5	3–3	2 to 3	rpg5	33+	3- to 3+	Rpg4	3–3	3– to 3	rpg1
QSM123	3	3	Rpg4/rpg5	3-21	1 to 3–	rpg5	33–	3- to 3+	Rpg4	33–	3– to 3	rpg1
QSM124	33+	3 to 3+	Rpg4/rpg5	23–1	1 to 3	rpg5	3–2	2 to 3+	Rpg4	0;1-	0; to 1	Rpg1
QSM125 OSM126	0;	0; 3– to 3	rpg4/Rpg5	00;	0 to 1 1 to 3–	Rpg5	00; 3–3	0 to 2 2 to 3	rpg4	33– 3–3	3– to 3	rpg1
QSM126 QSM127	33– 3+	3-10 3 3+	Rpg4/rpg5 Rpg4/rpg5	3–21 3–3	2 to 3	rpg5 rpg5	3-3 33-	2 to 3+	Rpg4 Rpg4	0;1-	3– to 3 0; to 1–	rpg1 Rpg1
QSM128	3	3– to 3+	Rpg4/rpg5	33–2	2 to 3	rpg5	33–	2 to 3	Rpg4	0;1-	0; to 1	Rpg1
QSM129	3–3	2 to 3	Rpg4/rpg5	3–21	1 to 3–	rpg5	3–3	2 to 3	Rpg4	0;1-	0; to 1	Rpg1
QSM130	3-3	3– to 3	Rpg4/rpg5	213-	1 to 3-	rpg5	33-	2 to 3+	Rpg4	3–3	3– to 3	rpg1
QSM132	3–3	3– to 3	Rpg4/rpg5	3–21	1 to 3–	rpg5	3–2	1 to 3–	Rpg4	3–3	3– to 3	rpg1
QSM133	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	0;1-	0; to 1	Rpg1
QSM134 QSM135	33-	2 to 3 0; to 2–	Rpg4/rpg5 rpg4/Rpg5	3–21	1 to 3– 0 to 1–	rpg5 Rpg5	33- 0;1	2 to 3+ 0; to 2	Rpg4	33– 33–	3– to 3+ 3– to 3	rpg1
QSM135 QSM136	0;1 0;1	0; to 2– 0; to 1	rpg4/Rpg5	00; 0;1	0 to 1– 0; to 1	крд5 Rpg5	00;	0; to 2 0 to 1	rpg4 rpg4	3–3	2 to 3	rpg1 rpg1
QSM137	0;1	0; to 1	rpg4/Rpg5	0;1	0; to 1	Rpg5	10;	0; to 2	rpg4	0;1	0; to 1	Rpg1
QSM138	0;1-	0; to 1–	rpg4/Rpg5	0;1	0 to 1	Rpg5	00;	0 to 0;	rpg4	33–	3– to 3	rpg1
QSM139	3–3	2 to 3	Rpg4/rpg5	3–3	2 to 3	rpg5	3–2	2 to 3	Rpg4	12	0; to 2	Rpg1
QSM140	0;	0;	rpg4/Rpg5	0;	0;	Rpg5	0;1	0; to 3	rpg4	0;1	0; to 2	Rpg1
QSM141	0;	0;	rpg4/Rpg5	00;	0 to 1	Rpg5	00;	0 to 1	rpg4	3–3	2 to 3	rpg1
QSM142 QSM143	0;1	0; to 1	rpg4/Rpg5	0;1 213–	0 to 1 1 to 3–	Rpg5	0;1 33–	0; to 2 2 to 3+	rpg4 Rpg4	0;1– 12	0; to 1– 0; to 2	Rpg1
QSM143 QSM144	3 3+3	2 to 3+	Rpg4/rpg5 Rpg4/rpg5	3–21	1 to 3– 1 to 3–	rpg5 rpg5	33– 33–	2 to 3+	Rpg4 Rpg4	3–3	3– to 3	Rpg1 rpg1
QSM144 QSM145	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1–	rpg4	0;1	0; to 1	Rpg1
QSM146	3–	3–	Rpg4/rpg5	3–	2 to 3	rpg5	3–3	2 to 3	Rpg4	12-	0; to 2–	Rpg1
QSM147	3–3	3– to 3	Rpg4/rpg5	3–3	3– to 3	rpg5	3–2	2 to 3–	Rpg4	10;	0; to 2	Rpg1
QSM148	0;	0;	rpg4/Rpg5	00;	0 to 0;	Rpg5	00;	0 to 1	rpg4	0;1-	0; to 1–	

indicating that a single gene in Q21861 confers resistance to race TTKSK.

The Q/SM population was previously phenotyped for resistance to P. graminis f. sp. tritici races QCCJ and MCCF and also P. graminis f. sp. secalis isolate 92-MN-90 (35,39). Raw IT data recorded on Q/SM progeny from these evaluations are given in side-by-side comparisons with those taken for race TTKSK in Table 1. All of the progeny exhibiting low IT modes (0; to 12) to race TTKSK had similar low IT modes (00; to 10; and, rarely, 21) to P. graminis f. sp. tritici race QCCJ and P. graminis f. sp. secalis isolate 92-MN-90. Thus, resistance to race TTKSK co-segregated with resistance to race QCCJ and isolate 92-MN-90 in the Q/SM population. Similarly, progeny exhibiting high IT modes (23- to 3+) to race TTKSK also gave high IT modes to race QCCJ (3-2 to 3+3). A similar result was observed in response to isolate 92-MN-90 in that most of the 71 progeny exhibiting high IT modes to race TTKSK gave high IT modes of 23-1 to 33- to the rye stem rust isolate. However, 16 of the 71 TTKSK-susceptible progeny gave lower IT modes (10;23-, 12, 21, and 213-). These progeny were still considered susceptible because the typical resistant reaction to isolate 92-MN-90 is a very low IT of 00; to 0;1. Thus, progeny giving an IT mode higher than 0;1 or with 3type uredinia comprising part of its mesothetic reaction were considered susceptible. These phenotypic classifications were confirmed at the molecular level (see below).

The Q/SM population also was evaluated to race MCCF at high temperature (35). This was done to assess whether individual DH progeny also carried Rpg1 because this race, when used at high temperature, is useful for detecting lines with and without the gene. The high-temperature incubation was used because resistance gene rpg4, originally found to confer resistance to race QCCJ (17), also is effective against MCCF at low incubation temperatures. Using the high incubation temperature, rpg4 is rendered completely ineffective and, therefore, lines carrying Rpg1 can be clearly resolved based on their reaction to MCCF. The presence of Rpg1 had no effect on the reaction of DH progeny to race TTKSK. Similar results were previously reported in response to race QCCJ and isolate 92-MN-90 (17,35,39). These data demonstrate that, although Rpg1 confers resistance to many races of P. graminis f. sp. tritici (33), it is not effective against races TTKSK (34) or QCCJ (38). Additionally, Rpg1 is generally not effective against isolates of P. graminis f. sp. secalis as was previously reported (33,38).

Resistance to race QCCJ in line Q21861 is conferred by a single recessive-acting gene, rpg4 (17). This gene was initially mapped to the long arm of chromosome 5H using random amplified polymorphic DNA (RAPD) and RFLP markers (2). In a subsequent study, Sun et al. (39) found Q21861 to be resistant to *P. graminis* f. sp. *secalis* isolate 92-MN-90 and demonstrated that the resistance was due to a single partially dominant gene. Resistance to isolate 92-MN-90 co-segregated with resistance to race QCCJ, prompting the authors to conclude that resistance to

both formae speciales of P. graminis is controlled by a gene at the same locus as rpg4. In this study, resistance to race TTKSK cosegregated with resistance to both P. graminis f. sp. tritici race QCCJ and also P. graminis f. sp. secalis isolate 92-MN-90 in the 129 Q/SM progeny, indicating that the gene conferring resistance to race TTKSK also lies at the rpg4 locus. Molecular analysis of the region in the Q/SM population positioned the locus conferring resistance to all three rusts at one crossover proximal to ABG391 and six crossovers distal to MWG740 (Fig. 1). This covers a large genomic region of approximately 5.4 centimorgans. Preliminary analysis of recombinants previously used to characterize loci conferring resistance to QCCJ and 92-MN-90 (3) suggested that the TTKSK resistance locus also resides within the narrower 70kb region between ARD5112 and ARD5016. This region encodes four genes in the resistant parent: Rpg5, a nucleotide-binding site leucine-rich repeat serine/threonine-protein kinase gene; rpg4, an actin depolymerizing factor-like gene; and two other genes of unidentified function (3). Positioning of the TTKSK resistance locus within the same ARD5112-ARD5016 region suggests that one or more of these genes are responsible for conferring resistance to race TTKSK. Previous high-resolution genetic and molecular analyses of the rpg4/Rpg5 region suggested that although rpg4 and Rpg5 are separate genes, both may be required for resistance to race QCCJ, whereas Rpg5 functions independently to provide resistance to isolate 92-MN-90 (3). Brueggeman et al. (4) recently suggested that resistance to race TTKSK may also involve both rpg4 and Rpg5. We can neither confirm nor refute this hypothesis based on our current data; however, this aspect is being investigated by genetic, molecular, and virus-induced gene silencing analyses.

All of the stem rust phenotyping for the Q/SM population was done on seedling plants under the controlled conditions of the greenhouse or growth chamber. A critical question to ask is whether the stem rust resistance found at the seedling stage would also be effective at the adult plant stage in the field. Adult plant resistance is essential because stem rust causes the most damage on barley after the heading stage. In field screening nurseries established at the Kenya Agricultural Research Institute in Njoro, Kenya where races TTKSK and TTKST were most common, Q21861 was highly resistant, exhibiting severities of 1 to 5% and mostly resistant to moderately resistant ITs (B. Steffenson, unpublished). The genetic basis of adult plant resistance to race TTKSK is being explored in the Q/SM population by mapping mRNA transcript abundance (i.e., expression quantitative trait loci) with the Barley1 Affymetric array (5) (R. Wise, personal communication), as was done previously for seedling resistance in the Steptoe/Morex population to race MCCF (7).

Different virulence types within the originally described race TTKS lineage (26) have been described in eastern and southern Africa (18,19,41). In addition to the hallmark virulence for resistance gene Sr31 (16), these types also carry virulence for Sr24 (race TTKST) (18) and Sr36 (race TTKSK) (19). Wheat

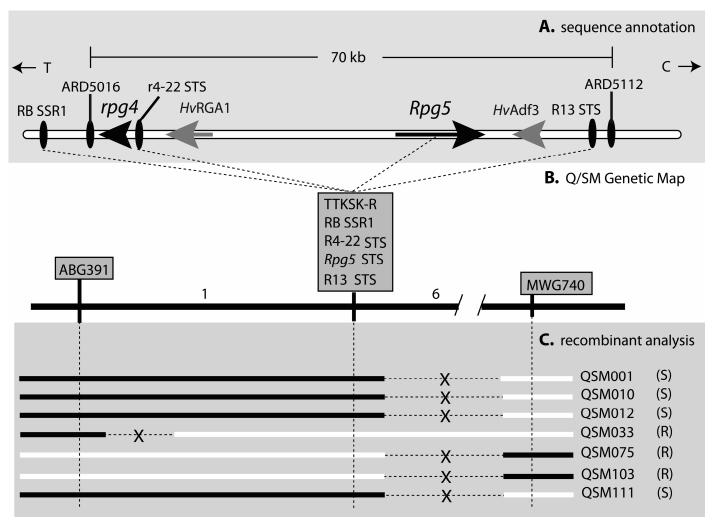
TABLE 2. Primer sequences for polymerase chain reaction (PCR)-based markers in the rpg4/Rpg5 region of barley chromosome 5H

			PCR products (bp) <sup>c</sup>			
Markers <sup>a</sup>	Primers	Sequence 5′–3′ <sup>b</sup>	Q21861	SM89010		
RB SSR1	RB_SSR1-F1	CACATCCACCCATGGTTGTTGAGAG	304	304		
	RB_SSR1-R1	CTTCACTGGTACCAGTTCGACCGAG				
R4-22 STS	r4-F22	CGGATATAGGGAGAAGGGTTTGATG	922	None		
	r4-R22	GTCGTCCATCTTGTACACCACGAAC				
Rpg5 STS	QK-F6	GGTGGATCGAAGAGAATGGAACTGC	677	None		
	R5-R7.3	GCAACCTTCATTCTGACAGACCATG				
R13 STS	A12-F13	CGCCCGACGAAAGAGAACGACAATG	1,337	None		
	A12-R13	GGGCCACCGACACTGTAGCACTC	•••	•••		

<sup>&</sup>lt;sup>a</sup> Markers are designated as simple sequence repeat (SSR) or sequence tag site (STS).

<sup>&</sup>lt;sup>b</sup> Primers are listed with 5'-3' sequences.

<sup>&</sup>lt;sup>c</sup> Amplicon sizes are shown for each marker. All the STS markers are dominant PCR markers only amplifying a product from the resistant parent Q21861.



**Fig. 1.** Genetic and molecular analysis of the TTKSK resistance locus on barley chromosome 5H. **A,** Sequence annotation of the rpg4/Rpg5 locus from the resistant line Q21861. The white horizontal bar represents genomic sequence from line Q21861. Arrows represent annotated genes, rpg4, Rpg5 (black), and other genes (gray). Black ovals represent the position of genetic markers. ARD5016 and ARD5112 are restriction fragment-length polymorphism markers reported in Brueggeman et al. (3) delimiting the rpg4/Rpg5 region. Dashed lines connect markers and genes from sequence annotation to positions on the genetic map and recombinant analysis below. All genes and markers are labeled above. The scale is shown above in kilobases. C and T represent direction of the centromere and telomere, respectively. **B,** Genetic map of the rpg4/Rpg5 region from the Q21861/SM89010 (Q/SM) population. Black vertical bars represent the position of genetic and phenotypic markers present within the attached gray boxes above. The number of recombinants between markers is shown above. **C,** Recombinant genotype analysis. Horizontal bars represent the genotype from seven lines with recombination defining the TTKSK resistance region from the Q/SM population. White horizontal bars represent the resistant Q21861 parent and black the susceptible SM89010 parent. The dashed horizontal lines with Xs denote regions where recombination occurred. Dashed vertical lines represent the positions of connected genetic markers from the genetic map above. The recombinant designations are labeled to the right with resistant (R) and susceptible (S), indicating response to race TTKSK.

lines carrying *Sr24* were initially considered to be one of the best sources of resistance against race TTKSK (18). However, the identification of races with virulence for *Sr24* in eastern Africa revealed the great vulnerability of many wheat cultivars worldwide, which are based on this resistance gene. Q21861 appears to possess a high level of adult plant resistance against both TTKSK and TTKST, two of the most common virulence types found in eastern Africa. Still, it is important to be vigilant and continually test this line against other virulence types emerging in the lineage. For the present, the stem rust resistance in Q21861 may be useful in barley breeding programs because of its effectiveness against the common African virulence types and simple inheritance. However, additional genes for resistance to race TTKSK must be identified to increase the diversity and longevity of resistance in deployed barley cultivars.

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