



Characterization of stem, stripe and leaf rust resistance in Tajik bread wheat accessions

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Abstract Stem rust [causal organism: *Puccinia graminis* f. sp. *tritici* (*Pgt*)], stripe rust [*Puccinia striiformis* f. sp. *tritici* (*Pst*)], and leaf rust [*Puccinia triticina* (*Pt*)] are important fungal diseases of wheat in Central Asia and worldwide. Therefore, identification of seedling and adult plant resistance (APR) genes is of major importance for the national wheat breeding program in many countries. The objectives of this study were to identify genes that confer seedling and APR resistances in widely grown wheat cultivars, landraces and advanced lines from Tajikistan. A total of 41 wheat accessions were inoculated with eleven races of *Pgt*, twelve races of *Pst* and nine races of *Pt* for postulation of *Sr* (stem rust), *Yr* (yellow or stripe rust), and *Lr* (leaf rust) resistance genes at the seedling stage. In addition, all of the accessions were tested in field trials for the response to stem rust and stripe rust. Genes for seedling stem rust resistance (i.e.

Sr5, *Sr6*, *Sr11*, *Sr31*, and *Sr38*), stripe rust resistance (*Yr9*, *Yr17*, and *Y27*), and leaf rust resistance (*Lr16* and *Lr26*) were postulated in the Tajik wheat. The presence of the pleiotropic APR genes *Sr2/Yr30/Lr27* (associated with pseudo-black chaff phenotype) and *Lr34/Yr18/Sr57* (associated with leaf tip necrosis phenotype), and also *Lr37* were assessed in the field and confirmed with linked molecular markers. In most of the wheat accessions, resistance genes could not be postulated because their infection types did not match the avirulence or virulence profile of the *Pgt*, *Pst* and *Pt* races tested. Six, seven, and nine accessions were identified that likely possess new genes for resistance to stem rust, stripe rust, and leaf rust, respectively, which have not been described previously. The research demonstrates the presence of effective seedling resistance and APR genes in widely grown wheat accessions that could facilitate further rust

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resistance breeding in the national wheat breeding program in Tajikistan.

Keywords Gene postulation · Molecular marker · Resistance gene · *Triticum aestivum*

Introduction

Bread wheat (*Triticum aestivum* L., $2n = 6x = 42$, ~ 17 Gb, BBAADD genome) is one of the most important and widely cultivated food crops, contributing substantially to the daily nutrition and food security of a large proportion of the world's population (Shiferaw et al. 2013). Unfortunately, many abiotic and biotic stresses limit wheat production across the globe. Among the most important biotic stresses of wheat are the three rust diseases, namely stem rust [caused by *Puccinia graminis* f. sp. *tritici* Erikss. & E. Henning (*Pgt*)], stripe rust [*Puccinia striiformis* Westend. f. sp. *tritici* Eriks. (*Pst*)], and leaf rust [*Puccinia triticina* Eriks. (*Pt*)]. Since ancient times, these rust diseases have caused many epidemics, resulting in significant and widespread crop losses (Kolmer 2005; Hovmøller et al. 2011; Szabo et al. 2014). Stem rust and stripe rust can cause complete crop loss, and losses due to leaf rust can be as high as 70% (Chen 2005; Huerta-Espino et al. 2011; Singh et al. 2015). In recent epidemics, yield losses ranging from 20 to 100% were reported for these three rust diseases in wheat growing regions worldwide (Huerta-Espino et al. 2011; Wellings 2011; Singh et al. 2015).

In Tajikistan, bread wheat is the most important food crop with respect to national food security (FAO 2015) but is constantly threatened by these three rust diseases. Epidemics of stripe rust occurred in 1952, 1958, 1966, 1997, 1998, 2003, 2010, and 2016, resulting in significant yield losses across the country (Eshonova et al. 2005; Rahmatov et al. 2011, 2012). Stem rust occurs mainly in the mountainous areas (Pett et al. 2005); however, when favorable environmental conditions prevail, the disease is capable of destroying the grain yield of wheat crops across all agroecological zones of Tajikistan. Leaf rust is more variable with respect to its impact on wheat in the country (Eshonova et al. 2005; Rahmatov et al. 2012).

Deployment of host genetic resistance is considered the most effective and low-cost management strategy

for rust diseases, particularly in developing countries (Ellis et al. 2014). To control these rust diseases in Tajikistan, the national wheat breeding program has developed several rust resistant wheat cultivars by utilizing advanced breeding lines from the International Winter Wheat Improvement Program and Centro Internacional de Mejoramiento de Maíz y Trigo (CIMMYT). Genetic resistance to rust diseases has been broadly categorized into “seedling resistance,” which is often conferred by single genes with major phenotypic effects across all growth stages of the plant (Flor 1971), and “adult plant resistance” (APR), which is often conferred by multiple genes with more subtle phenotypic effects during the later ontogenetic stages of plant development (Knott 1989). In selecting and breeding for rust resistance, seedling and adult plant phenotyping assays are routinely performed along with molecular marker assays if available for specific resistance genes (Juliana et al. 2017). One of the greatest challenges in breeding for rust resistance in wheat is the genetic variability of the rust pathogens. The virulence diversity of the three rust pathogens in Central Asia is high, particularly in Tajikistan (Kolmer and Ordoñez 2007; Berlin et al. 2015; Ali et al. 2017). For example, the first time *Pst* virulence was found for the yellow rust (*Yr*) resistance genes of *Yr1*, *Yr4* + , *Yr3 N*, *Yr9*, *Yr10*, *Yr17* and *Yr27* in Central Asia was in Tajikistan (Yahyaoui et al. 2012a, b). Eight barberry species have been reported in Tajikistan (Davlatov and Baikova 2011), which may play a role in disease epidemics and pathogen variation in the country since these species could potentially serve as alternate hosts for both *Pgt* and *Pst*.

Currently, more than 70 stem rust (*Sr*) resistance genes, 65 yellow rust (*Yr*) resistance genes and 79 leaf rust (*Lr*) resistance genes, including those with minor effects have been cataloged (McIntosh et al. 2017). Widely deployed cultivars with effective resistance genes can suffer yield losses when new, virulent races of the stem, stripe, and leaf rust pathogens emerge, leading to the “boom and bust” cycle of plant breeding (Pretorius et al. 2000; Huerta-Espino et al. 2011; Wellings 2011; Solh et al. 2012). Some of the most widely used and important resistance genes in wheat include *Sr13*, *Sr24*, *Sr31*, *Sr36*, *Sr38*, *SrTmp* and *SrIRS^{Amigo}* for stem rust, the *Yr2*, *Yr6*, *Yr7*, *Yr8*, *Yr9*, *Yr17* and *Yr27* for stripe rust, and *Lr9*, *Lr14a*, *Lr16*, *Lr17a*, *Lr24*, *Lr26* and *Lr39* for leaf rust all of which

have now been overcome by newly detected pathogen races (Huerta-Espino et al. 2011; Singh et al. 2015; Ali et al. 2017). The lack of knowledge regarding the presence of major effect seedling and minor effect APR resistance genes in Tajik wheat germplasm makes it difficult to make informed decisions with respect to breeding for stable resistance in the national wheat breeding program. Therefore, the aims of this study were to (1) evaluate Tajik wheat accessions for seedling resistance and postulate the presence of underlying *Sr*, *Yr* and *Lr* genes; (2) identify the presence of gene(s) conferring APR to the three rust diseases; and (3) verify the presence of resistance genes postulated by available molecular markers.

Materials and methods

Plant and pathogen materials

A total of twenty-nine wheat cultivars, seven advanced breeding lines, and five landraces were used in the present study and tested for response to the three rusts. These wheat accessions were provided by the national wheat breeding program in Tajikistan. The pedigree and origin of the materials are given in Table 1. In addition, differential wheat accessions with characterized resistance genes for stem rust (Jin et al. 2007), stripe rust (Hovmøller et al. 2017), and leaf rust (Kolmer and Hughes 2013) were also included to facilitate the gene postulations. Eleven *Pgt*, twelve *Pst* and nine *Pt* races with different virulence/avirulence combinations and geographic origins were used (Tables 2, 3 and 4).

Seedling rust resistance assays

Seedling resistance assays to stem rust and leaf rust were conducted at the United States Department of Agriculture-Agricultural Research Service-Cereal Disease Laboratory (USDA-ARS-CDL) and the University of Minnesota in St. Paul, USA. Five seeds of each wheat genotype were included for each rust assay. The seeds were planted in pots containing vermiculite (Sun Gro Horticulture), watered daily, and fertilized with 20–20–20 NPK soluble fertilizer (Spectrum Group, St. Louis). Stored urediniospores of the stem and leaf rust pathogens were removed from a – 80 °C freezer, heat-shocked at 45 °C for 15 min

and placed in a rehydration chamber for 2 to 4 h maintained at 80% relative humidity by a KOH solution, and then suspended in a lightweight mineral oil (Soltrol 170[®] Chevron Phillips Chemical Company LP, Woodlands, TX 77380) within gelatin capsules (size 00). Then, urediniospores were inoculated onto 8–10 day-old seedlings of the different accessions at the first leaf stage. Seedling resistance assays for stem rust were done according to the methods of Rouse et al. (2011) and those for leaf rust were done according to Oelke and Kolmer (2004). Infection types were scored 14 days after inoculation using a 0–4 scale (Stakman et al. 1962; Long and Kolmer 1989). Seedling resistance to *Pgt* race TKTTF (bulk collection from Turkey) was carried out at the Regional Cereal Rust Research Center (RCRRC), located at the Aegean Agricultural Research Institute, International Center for Agricultural Research in the Dry Areas (ICARDA) in Izmir, Turkey (Rahmatov et al. 2016). The methods used for this test were similar to those used for the other races, the exception being that fresh urediniospores collected from plants in the field were used instead of frozen urediniospores. Ten-day-old seedlings with the first leaves fully expanded were inoculated with race TKTTF according to Rahmatov et al. (2016).

All accessions were evaluated for seedling stripe rust resistance at the Global Rust Reference Center (GRRC) at Aarhus University in Flakkebjerg, Denmark and at the RCRRC. For these evaluations, ten seeds were sown in pots containing a mixture of peat moss and soil. Inoculations with races of *Pst* were carried out on 14-day-old seedlings when the second leaves were fully expanded. For inoculations completed at the GRRC and RCRRC, *Pst* urediniospores were suspended in Novec Fluid (3 M Novec[™] 7100 Engineered Fluid) and lightweight mineral oil, respectively (Rahmatov et al. 2017). After inoculation, plants were moved to a dark chamber at 100% RH at 10 °C for 24 h for the infection period. Thereafter, plants were incubated in a greenhouse at 18 °C for 18 h during the day and 12 °C for 6 h during the night, protocols routinely used at both the GRRC and RCRRC (Hovmøller et al. 2017; Rahmatov et al. 2017). After 16 days of incubation, stripe rust infection types were scored using a 0–9 scale as described by McNeal et al. (1971).

Table 1 List of wheat accessions evaluated in this study

#	Accession	Pedigree	Origin	Type	Accession status
1	Navruz	(S)MIRONOVSKAYA-YUBILEINAYA	Tajikistan	Facultative	Cultivar
2	Sarvar	CHEN/AEGILOPS SQUARROSA (TAUS)//BCN/3/BAV92	ESWYT25	Spring	Cultivar
3	Vahdat	VORONA/CNO79//KAUZ/3/MILAN	ESWYT25	Spring	Cultivar
4	Yusufi	SOROCA	ESWYT25	Spring	Cultivar
5	Isfara	SW89.5181//KAUZ	ESWYT25	Spring	Cultivar
6	Alex	PAYNE(PYN)/(BAU)BAGULA	1WWEERYT	Facultative	Cultivar
7	Oriyon	NORD-DESPREZ/VG-9144//KALYANSONA/BLUEBIRD/3/YACO/4/VEERY-5	N.A.	Facultative	Cultivar
8	Sadokat	JUPATECO-73/BLUEJAY//URES-81	Mexico	Spring	Cultivar
9	Ziroat-70	N.A.	N.A.	Facultative	Cultivar
10	Norman	OR-F-1-158/(FDL)FUNDULEA//(BLO)BOLILLO/3/SHI-4414/CROW	5FAWWON	Facultative	Cultivar
11	Somoni	N.A.	N.A.	Facultative	Cultivar
12	Tacicar	TR.AE/SPARROW//ZARAPITIN	5FAWWON	Facultative	Cultivar
13	Ormon	NWT/3/TAST/SPRW//TAW12399.75	8FAWWON	Facultative	Cultivar
14	Iqbol	RUSALKA,BGR/CA-8055//CHAM-6	N.A.	Facultative	Cultivar
15	Starshina	COLT/SPARTANKA	Russia	Winter	Cultivar
16	Shokiri	SHARK/F-4105-W-2-1	5WWEERYT	Facultative	Cultivar
17	Fayzbaksh	TAM200//KAUZ	6WWEERYT	Facultative	Cultivar
18	Nurbakhsh	PRINA/STAR	N.A.	Facultative	Cultivar
19	BASRIBEY-95	JUPATECO-73/(SIB)BLUEJAY//URES-81	Turkey	Facultative	Cultivar
20	Jagger	KS-82-W-418/STEPHENS	USA	Winter	Cultivar
21	Kaboi Panjakent	N.A.	Tajikistan	Facultative	Landrace
22	Surkhaki 5	N.A.	Tajikistan	Spring	Landrace
23	Zafar	N.A.	Tajikistan	Facultative	Cultivar
24	Steklovidnaya-24	BOGARNAYA-56/TEPLOKL YUCHENSKAYA-2//ROSTOVCHANKA	Kazakhstan	Winter	Cultivar
25	SIETE-CERROS-66	PENJAMO-62(SIB)/GABO-55	Mexico	Spring	Cultivar
26	Krasnodarskaya-99	LUTESCENS-2665-G-10233/ERYTHROSPER MUM-4695-h-449//LUTESCENS-2621-h-24-82	Russia	Winter	Cultivar
27	Jayhun	N.A.	Turkey	Facultative	Cultivar
28	IZ-80	KAUZ * 2/CHEN//BCN/3/MILAN		Facultative	Cultivar
29	AIKT-20	CBRD//KAUZ		Facultative	Cultivar
30	N.A.	OTUS TOBA 97		Facultative	Advanced line
31	N.A.	PASTOR/3/VORONA		Facultative	Advanced line
32	N.A.	CMN82A.1294/2*		Facultative	Advanced line
33	N.A.	HUAVUN INIA		Facultative	Advanced line
34	Trakua Hatti	N.A.	Turkey	Facultative	Advanced line
35	Murodi-2013	CHEN/AE.SQ//WEAVER/3/SSERI1	27ESWYT	Spring	Cultivar
36	N.A.	CHEN/AE.SQ//WEAVER/3/PASTOR	27ESWYT	Spring	Advanced line
37	Ganj	NAC/TH.AC//3 * PVN/3/MIRLO/BUC/4/2 *vPASTOR	27ESWYT	Spring	Cultivar
38	N.A.	NAC/TH.AC//3 * PVN/3/MIRLO/BUC/4/2 * PASTOR	27ESWYT	Spring	Advanced line

Table 1 continued

#	Accession	Pedigree	Origin	Type	Accession status
39	Safedaki Pomir	N.A.	Tajikistan	Spring	Landrace
40	Safedaki Ishkoshim	N.A.	Tajikistan	Spring	Landrace
41	Babilo Pomir	N.A.	Tajikistan	Spring	Landrace

N.A. not available, *ESWYT* elite spring wheat yield trial, *FAWWON* facultative and winter wheat observation nursery, *WWEERYT* winter wheat eastern european regional yield trial

Assessment of field response to stem rust and stripe rust

Adult plant stem rust responses were evaluated under field conditions at the Kenyan Agricultural and Livestock Research Organization in Njoro (2010 and 2011), at the RCRRC in Izmir (2014) and at the University of Minnesota in St. Paul (2014). In Tajikistan, the wheat accessions were exposed to naturally occurring races of *Pst* during the growing season of 2010. The stripe rust-infected leaves were collected in Tajikistan and sent to the GRRC for race analysis (Hovmøller et al. 2017), and the race TJ01a/10 was detected and subsequently used at the seedling resistance test. To provide sufficient stripe rust infection in the nurseries at RCRRC, mixtures of susceptible wheat cultivars were used as spreader rows surrounding and between the plots (Rahmatov et al. 2017). In Njoro and Minnesota, urediniospores of *Pgt* (TTKSK + TTKST, and MCCFC) were needle-injected (i.e. injecting urediniospores directly into the stems of susceptible spreader plants) at the tillering, booting and heading stages. Additionally, direct foliar inoculations were made on the spreader rows using a urediniospore/oil suspension (Rahmatov et al. 2016). In Izmir, the spreader rows were inoculated five times at the tillering, booting and heading stages by dusting a mixture of fresh urediniospores of *Pgt* (TKTTF) and *Pst* (TK34/11) together with talcum powder. After inoculation, the nurseries in Njoro and Izmir were mist-irrigated three times per day (i.e. morning, afternoon and evening) to ensure a moist environment and thereby enhance stem and stripe rusts development. The adult plant response to stem and stripe rust were assessed between growth stages 50–90 (Zadoks et al. 1974). Disease severity was assessed using the modified Cobb scale (Peterson et al. 1948) and adult plant infection types were rated according to Roelfs

et al. (1992). The presence of the pseudo-black chaff (*PBC*) and leaf tip necrosis (*LTN*) phenotypes were assessed using 0–4 scale in all field trials (Juliana et al. 2015).

Molecular marker analysis

Total genomic DNA was isolated from the leaves of 10 day-old seedlings according to Edwards et al. (1991) with some slight modifications. The molecular markers *XcsSr2*, *Xgwm533* and *wMAS000005* for *Sr2/Yr30/Lr27* (Spielmeyer et al. 2003; Mago et al. 2011), *Xcfd43* for *Sr6* (Tsilo et al. 2009) *Xwmc364* for *Yr2* (Lin et al. 2005), *Xscm9* and *Xiag95* for *Sr31/Yr9/Lr26* (Saal and Wricke 1999; Mago et al. 2005), *csLV34* and *wMAS000003* for *Lr34/Yr18/Sr57* (Lagudah et al. 2006), and *VENTRIUP/LN2* for *Sr38/Yr17/Lr37* (Helguera et al. 2003) were assessed. The PCR assays were conducted according to Rahmatov et al. (2016).

Results

Stem rust seedling response assays

A majority of the wheat accessions showed seedling resistance towards the *Pgt* races of RKQQC, QTHJC, TPMKC, BCCBC, and MCCFC with infection types (ITs) ranging from 0 to 2+ (Table 5). A lower proportion of the wheat accessions showed seedling resistance towards the more widely virulent *Pgt* races of TTTTF, TTKSK, TTTSK, TTKST, TRTTF and TKTTF (Table 5). The resistance gene *Sr5* was postulated in Navruz and Steklovidnaya-24 based on its resistance reaction to race BCCBC (Table 5). *Sr6* and *Sr11* were postulated in Siete-Cerros-66 based on its resistance reactions to races RKQQC, TPMKC, TKTTF, MCCFC and BCCBC (Table 5). Resistance

Table 2 The origin and virulence phenotype of *Puccinia graminis* f. sp. *tritici* races used in this study

Race	Isolate	Origin	Virulence profile																			
			Sr5	Sr21	Sr9e	Sr7b	Sr11	Sr6	Sr8a	Sr9g	Sr36	Sr9b	Sr30	Sr17	Sr9a	Sr9d	Sr10	SrTmp	Sr24	Sr31	Sr38	SrMcN
RKQC	99KS76A-1	USA	5	21	-	7b	-	6	8a	9g	36	9b	-	-	-	9d	-	-	-	-	-	McN
QTHC	75ND717C	USA	5	21	-	-	11	6	8a	9g	-	9b	-	17	-	9d	10	-	-	-	-	McN
TPMKC	74MIN1409	USA	5	21	9e	7b	11	-	8a	9g	36	-	17	-	9d	10	Tmp	-	-	-	-	McN
TTTTF	02MIN84A-1-2	USA	5	21	9e	7b	11	6	8a	9g	36	9b	30	17	9a	9d	10	Tmp	-	-	38	McN
TTKSK	04KENI156/04	Kenya	5	21	9e	7b	11	6	8a	9g	-	9b	30	17	9a	9d	10	-	-	31	38	McN
TTTSK	07KEN24-4	Kenya	5	21	9e	7b	11	6	8a	9g	36	9b	30	17	9a	9d	10	-	-	31	38	McN
TTKST	06KENI9-V-3	Kenya	5	21	9e	7b	11	6	8a	9g	-	9b	30	17	9a	9d	10	-	24	31	38	McN
TRTTF	06YEM34-1	Yemen	5	21	9e	7b	11	6	-	9g	36	9b	30	17	9a	9d	10	Tmp	-	-	38	McN
TKTTF		Turkey	5	21	9e	7b	-	6	8a	9g	36	9b	30	17	9a	9d	10	-	-	-	-	McN
BCCBC	09CA115-2	USA	-	-	-	-	-	-	-	9g	-	-	-	17	-	-	-	-	-	-	-	McN
MCCFC	59KS19	USA	5	-	-	7b	-	-	-	9g	-	-	-	17	-	10	Tmp	-	-	-	-	McN

- Indication of avirulence

gene *Sr31* was postulated in Alex, Sadokat, Ziroat-70 and Otus Toba97 based on their susceptible reactions to races TTKSK, TTTSK and TTKST (Table 5); and *Sr38* in Jagger and IZ-80 based on their susceptible reactions to races TTTTF, TTKSK, TTTSK, TTKST and TRTTF (Table 5). The landraces of Kaboi Panjakent, Surkhaki-5, Jayhun, Safedaki Pomir, and Safedaki Ishkoshim were resistant to races TTKSK, TTTSK and TTKST (Table 5). Only Sarvar was highly resistant to all the tested races (Table 5). If any previously described resistance genes were present in this group of accessions, they could not be postulated because the resulting ITs did not match those of any differential accessions. Thus, these accessions either carry combinations of previously described genes or new resistance gene/s.

Stripe rust seedling response assays

Postulations for *Yr* genes were conducted using 12 *Pst* races (Table 3). *Yr9* and *Yr17* were confirmed based on the stem rust gene postulations for *Sr31* and *Sr38* plus molecular markers because of their tight linkage with the respective genes within the 1BL.1RS wheat-rye and 2NS/2AS translocations. These assays confirmed the presence of *Yr9* in Alex, Sadokat, Ziroat-70, and Otus Toba97 and *Yr17* in Jagger and IZ-80 (Tables 6, 8). Because Alex, Sadokat, Ziroat-70, Otus Toba97, Jagger and IZ-80 were resistant to most of the *Pst* races used in this study, including those carrying virulence for *Yr9* and *Yr17*, thus it was not possible to postulate genes based on their ITs to the 12 *Pst* races used in this study (Table 6). The *Yr27* was confirmed in Isfara based on the *Yr27*-virulent isolates AF87/12 and TR34/11 conferring ITs of 7 on *Yr27* differential lines (Table 6). Sarvar, Fayzbakhsh, Otus Toba97, Vahdat, Oriyon, Sadokat and AIKT-20 were highly resistant (ITs 0-4) to all or nearly all races; thus, the genes they carry could not be postulated with the *Pst* races used in this study nor the molecular markers. These accessions carry combinations of previously described genes or new resistance gene/s (Table 6).

Leaf rust seedling response assays

For the leaf rust seedling evaluations, nine *Pt* races were used (Table 4). The number of resistant and susceptible accessions for each of the races is presented in Table 7. *Lr16* was postulated in Iqbol,

Table 3 The origin and virulence phenotype of *Puccinia striiformis* f. sp. *tritici* races used in this study

Race	Origin	Virulence profile																				
		Yr1	Yr2	Yr3	Yr4	Yr5	Yr6	Yr7	Yr8	Yr9	Yr9+	Yr10	Yr15	Yr17	Yr24	Yr25	Yr27	Yr32	YrSd	YrSu	YrSp	YrAvS
SE205/12	Sweden	1	2	3	-	-	6	7	8	9	9+	-	-	17	-	25	-	32	Sd	-	-	AvS
UK94/519	UK	1	2	3	-	-	-	-	-	9	9+	-	-	17	-	25	-	-	Sd	Su	-	AvS
DK66/02	Denmark	-	2	-	-	-	6	7	8	9	-	-	-	-	-	25	-	-	Sd	-	-	AvS
TJ01a/10	Tajikistan	1	2	3	4	-	6	-	-	9	9+	-	-	-	-	25	-	32	Sd	Su	-	AvS
ER02/03	Eritrea	-	2	-	-	-	6	7	8	9	-	10	-	-	24	25	-	-	-	-	-	AvS
DK11/09	Denmark	-	-	3	4	-	6	-	-	-	-	-	-	-	-	25	-	32	Sd	Su	Sp	AvS
DK71/93	Denmark	1	2	3	-	-	-	-	-	-	-	-	-	-	-	25	-	32	Sd	-	-	AvS
AF87/12	Afghanistan	-	2	-	-	-	6	-	8	-	-	-	17	-	-	-	27	32	-	-	-	AvS
DK09/11	Denmark	1	2	3	4	-	6	7	-	9	9+	-	-	17	-	25	-	32	Sd	Su	Sp	AvS
DK122/09	Denmark	1	2	3	4	-	6	-	-	9	9+	-	-	17	-	25	-	32	Sd	Su	-	AvS
SE100/09	Denmark	-	-	-	-	-	-	7	8	-	-	10	-	-	-	-	-	-	-	Su	-	AvS
TK34/11	Turkey	-	-	2	-	-	6	7	8	9	-	-	-	-	-	25	27	-	(Sd)	(Su)	-	AvS

- Indication of avirulence

OTUS TOBA97, and HUAVUN INIA based on their susceptible reactions (ITs of 33 +) to race MHDSB (Table 7). *Lr26* was postulated in Alex, Sadokat, and Ziroat-70 based on their susceptible reactions (ITs 33 +) to races KFBJS, MHDSB, and TCRKG and molecular markers (Tables 7, 8). OTUS TOBA97 was resistant to all *Pt* races, except MHDSB (Table 7); therefore, the presence of *Lr26* was confirmed based on the stem rust, stripe rust and molecular marker analysis (Tables 7, 8). Nine accessions (Sarvar, Vahdat, PRINA/STAR, Zafar, AIKT-20, PASTOR/3/VORONA, CMN82A.1294/2*, Murodi-2013, and Ganj) likely carry combinations of previously described *Lr* genes or new *Lr* gene/s.

Field stem rust responses

For all of the stem rust field evaluations in Kenya, Turkey, and USA, a high level of disease pressure was attained as severities were 100% in susceptible controls. Some accessions showing no discernible seedling resistance exhibited high levels of APR in the field evaluations (Table 5). Thus, despite susceptibility at the seedling stage for TTKSK and TTKST, accessions PASTOR/3/VORONA/CN079 (10MSS), CMN82A.1294/2* (50MR) and HUAVUN INIA (40MR) against the *Pgt* race TTKSK + TTKST were exhibited APR during 2010 and 2011 in Njoro (Table 5). Furthermore, the accessions Vahdat, Somoni, Iqbol, Fayzbaksh, Kaboi Panjakent, and Surkhaki-5 exhibited disease severities of 5 to 40% with R to MR infection types, whereas Murodi-2013, Ganj, Krasnodarskaya-99, and Babilo Pomir had severities of 20 to 40% with MR-MS or MS infection types against race TKTTF in Izmir (Table 5). To race MCCFC in the USA, Navruz, Starshina, Basirbey, Kaboi Panjakent, Surkhaki-5, Steklovodnaya-24, Jayhun, Safedaki Pomir, and Safedaki Ishkoshim exhibited severities of 5 to 40% with RMR to MRMS and MS infection types (Table 5). Four accessions exhibited all stage resistance against race TTKSK + TTKST, thirteen against race TKTTF, and 32 against race MCCFC (Table 5). Thereby, these lines carry seedling resistance genes that are effective into the adult plant stage and to diverse races at three different field sites (Table 5).

Table 4 The origin and virulence phenotype of *Puccinia triticina* races used in this study

Race	Origin	Virulence profile											
		<i>Lr1</i>	<i>Lr2a</i>	<i>Lr2c</i>	<i>Lr3</i>	<i>Lr9</i>	<i>Lr16</i>	<i>Lr24</i>	<i>Lr26</i>	<i>Lr3ka</i>	<i>Lr11</i>	<i>Lr17</i>	<i>Lr30</i>
TDBJG	USA	1	2a	2c	3	–	–	24	–	–	–	–	–
TFBJQ	USA	1	2a	2c	3	–	–	24	26	–	–	–	–
TNRJJ	USA	1	2a	2c	3	9	–	24	–	3 ka	11	–	30
MLDSD	USA	1	–	–	3	9	–	–	–	–	–	17	–
MBDSB	USA	1	–	–	3	–	–	–	–	–	–	17	–
TBBGG	USA	1	2a	2c	3	–	–	–	–	–	–	–	–
KFBJG	USA	–	2a	2c	3	–	–	24	26	–	–	–	–
MHDSB	USA	1	–	–	3	–	16	–	26	–	–	17	–
TCRKG	USA	1	2a	2c	3	–	–	–	26	3 ka	11	–	30

Race	Origin	Virulence profile											
		<i>LrB</i>	<i>Lr10</i>	<i>Lr14a</i>	<i>Lr18</i>	<i>Lr21</i>	<i>Lr28</i>	<i>Lr39</i>	<i>Lr42</i>	<i>Lr3bg</i>	<i>Lr14b</i>	<i>Lr20</i>	<i>Lr23</i>
TDBJG	USA	–	10	14a	–	–	28	–	–	–	14b	–	–
TFBJQ	USA	–	10	14a	–	21	28	–	–	–	14b	20	23
TNRJJ	USA	–	10	14a	–	–	28	39	–	–	14b	20	–
MLDSD	USA	B	10	14a	–	–	–	39	–	3bg	14b	20	23
MBDSB	USA	B	10	14a	–	–	–	–	–	3bg	14b	20	–
TBBGG	USA	–	10	–	–	–	28	–	–	3bg	14b	20	23
KFBJG	USA	–	10	14a	–	–	28	–	–	–	14b	20	23
MHDSB	USA	B	10	14a	–	–	–	–	–	3bg	14b	20	–
TCRKG	USA	–	10	14a	18	–	28	–	–	3bg	14b	20	–

– Indication of avirulence

Field stripe rust responses

Stripe rust APR was detected in the seedling-susceptible accessions of Vahdat, Isfara, and Ormon (severities of 10 to 20% with infection types of R to MR) and also in Tacikar and CMN82A.1294/2* (severities of 40 to 50% with MR-MS infection types) against *Pst* race TK34/11 (Table 6). Somoni and Tacikar also possess some APR as they exhibited a stripe rust severity of 40% with MS infection types against race TJ01a/10 in Tajikistan. A total of twenty-one and eighteen accessions had all-stage resistance as they were highly resistant at both the seedling and adult plant stages to *Pst* races TK34/11 and TJ01a/10 in Turkey and Tajikistan, respectively (Table 6). The rest of the wheat accessions were susceptible at the seedling and adult plant stages (Table 6).

Phenotypic assessments of *PBC* and *LTN* in the field

The presence of the *PBC* and *LTN* phenotypes were associated with the pleiotropic *Sr2/Yr30/Lr27* and *Lr34/Yr18/Sr57* APR genes. The *PBC* phenotype (score of 2–3) was observed in 11 accessions, and the *LTN* phenotype (score of 2–3) was observed in 13 accessions in the field (Table 8).

Molecular marker analysis

The molecular markers *Xscm9* (220 bp), *Xiag95* (1100 bp) and *Xrems1303* (309 bp) indicated the presence of the *Sr31/Yr9/Lr26* resistance genes in Alex, Sadokat, Ziroat-70, and OTUS TOBA 97. Marker *Xcfd43* (215 bp) indicated the presence of *Sr6* in SIETE-CERROS-66, and marker *VENTRIUP/LN2* (262 bp) indicated the presence of the *Sr38/Yr17/Lr37* genes in Jagger and IZ-80. Marker *Xgwm533*

Table 5 Seedling infection types and field responses to stem rust, and molecular marker analysis for *Sr6*, *Sr31*, and *Sr38*

Accession	<i>Sr</i> seedling resistance									
	RKQQC	QTHJC	TPMKC	TTTTF	TTKSK		TTTSK		Rep. 1	Rep. 2
					Rep. 1	Rep. 2	Rep. 1	Rep. 2		
Navruz	1	0;/1	3+	3+	4	4	4	4	4	4
Sarvar	0;	0;	;1-	0;	22+	0;	22+	0;	22+	0;
Vahdat	0;	11+	1 + 2	0;	4	4	4	4	4	3+
Yusufi	1 + 2	1 + 2	1+	3+	4	4	4	3+	4	3+
Isfara	0;	1+	1;	3+	4	4	4	3+	4	3+
Alex	1;	11+	0;1	1;	4	4	4	3+	4	3+
Oriyon	0;	11+	11+	3+	4	4	4	3+	4	3+
Sadokat	0;	11+	1+	1+	4	4	4	3+	4	3+
Ziroat-70	0;	1 + 2	1	0;	4	4	4	3+	4	3+
Norman	4	1 + 2	0;1	3+	4	4	4	3+	4	3+
Somoni	1;1+	3+	3+	3+	4	4	4	3+	4	3+
Tacicar	4	3+	3+	3+	4	4	4	3+	4	3+
Ormon	3+	4	3+	3+	4	4	4	3+	4	3+
Iqbol	3+	3+	3+	1;	4	4	4	3+	4	3+
Starshina	22+	22+	4	3+	3+	3+	3+	3+	3+	3+
Shokiri	3+	3+	1;1+	3+	4	4	4	3+	4	3+
Fayzbaksh	0;	1+	2-	3+	4	4	4	3+	4	3+
PRINA/STAR	4	3+	4	3+	4	4	4	4	4	3+
BASRIBEY-95	2+	22+	3+	3+	4	4	4	3+	4	3+
Jagger	0;	0;	0;	3+	4	4	4	33+	4	3+
Kaboi Panjakent	1+	4	3+	4	2	2	2	1;	2	2
Surkhaki-5	2	4	3+	4	22+	1;	22+	1;	2	2
Zafar	3+	2 + 3-	3+	3+	4+	4	4+	4	4+	3+
Steklovidnaya-24	3+	3+	3+	3+	4	4	4	4	4	3+
SIETE-CERROS-66	1;	3+	;0	33+	4	4	4	3+	4	3+
Krasnodarskaya-99	1;	3+	33+	3+	4	4	4	3+	4	3+
Jayhun	2+	4	3+	3+	22+	2	22+	2	2	2
IZ-80	1;	0;	1;1+	3+	4	4	4	4	4	4
AIKT-20	0;	11+	1	3+	4	4	4	4	4	3+
OTUS TOBA 97	;	11+	1	0;1-	4	4	4	3+	4	3+

Table 5 continued

Accession	Sr seedling resistance									
	Sr Adult Plant Resistance									
	RKQOC	QTHJC	TPMKC	TTTTF	TTKSK	TTTSK	Rep. 1	Rep. 2	Rep. 1	Rep. 2
PASTOR3/VORONA/CN079	0;	11+	1 + 2	1 + 2	4	4	4	4	3+	3+
CMN82A.1294/2*	1;	11+	1;1+	0;	4	4	4	4	3+	3+
HUAVUN INIA	;1	11+	1;1+	3+	4	4	4	4	3+	3+
Trakua Hatti	0;	11+	11+	3+	4	4	4	4	3+	3+
Murodi-2013	22+	1+	1;2-	3+	4	4	4	4	3+	3+
CHEN/AE.SQ//WEAVER/3	1 + 2	1 + 2/2+	22+	3+	4	4	4	4	3+	3+
Ganj	33-	2+	3-	3+	4	4	4	4	3+	3+
NAC/TH.AC//3 * PVN/3/MIR	3	2 + 3	2 + 3	3+	4	4	4	4	3+	3+
Safedaki Pomir	4	4	2+	3+	4	4	4	4	0;-1	2
Safedaki Ishkoshim	3+	4	3+	3+	4	4	4	4	1;	2-
Babilo Pomir	4	22+	3+	3+	4	4	4	4	4	4
Accession	Sr seedling resistance					Sr gene postulation based on seedling and molecular marker				
	TTKST	TRTTF	TKTTF	BCCBC	MCCFC	TTKSK + TTKST	TTKS + TTKST	TKTTF	TKTTF	MCCFC
Rep. 1	Rep. 2	Rep. 1	Rep. 2	Rep. 1	Rep. 2	2010	2011			
Navruz	4	3+	4	0;	4	50S	60MSS	60MSS	5RMR	Sr5,+
Sarvar	0;	22+	11+	0;	0;	20MR	20MR	5R	TR	
Vahdat	4	4	11+	0;	0;	60S	70S	5R	TR	
Yusufi	3+	3+	1+	0;	0;	50S	60S	80S	TR	
Isfara	3+	3+	1+	0;	0;	60S	70S	80S	TR	
Alex	4	3+	2-	0;	0;	40S	50MSS	10RMR	TR	Sr31
Oriyon	3+	3+	3+	0;	0;	50S	60S	90S	5RMR	
Sadokat	4	3+	1+	0;	0;	40S	30S	10RMR	10RMR	Sr31
Ziroat-70	3+	3+	11+	0;	0;	40S	60S	20RMR	TR	Sr31
Norman	3+	3+	11+	0;	0;	40S	40S	70MSS	TR	
Somoni	3+	3+	1+	0;	0;	60S	60S	5RMR	TR	
Tacicar	3+	3+	1 + 2	0;	0;	60S	40S	70MSS	TR	
Ormon	4	3+	4	0;	0;	60S	80S	70MSS	TR	
Iqbol	3+	3+	1;	0;	0;	50S	80S	5RMR	TR	

Table 5 continued

Accession	Sr seedling resistance				Sr Adult Plant Resistance				Sr gene postulation based on seedling and molecular marker		
	TTKST		TRTTF		TTKSK + TTKST		TTKS + TTKST			MCCFC	
	Rep. 1	Rep. 2	TRTTF	TKTTF	TTKSK + TTKST	TTKS + TTKST	TKTTF	MCCFC			
Starshina	1 + 2	22+	4	2	0;	4	60MSS	70S	10RMR	TR	
Shokiri	3+	33+	3 + 4	0;	0;	0;	40MSS	60S	10RMR	5RMR	
Fayzbaksh	4+	3+	11+	3+	0;	0;	70S	90S	5RMR	TR	
PRINA/STAR	4	3 +/ 2	4	4	0;	0;	40MSS	70S	40MSS	10MRMS	
BASRIBEY-95	3+	3+	4	4	0;	4	80S	70S	80S	20MS	
Jagger	3+	3+	4	1	0;	0;	40MS	50MS	5R	TR	Sr38
Kaboi Panjakent	22+	2	4	4	0;	4	30RMR	20RMR	40MR	30MR	
Surkhaki-5	22+	2	3+	4	0;	4	20MR	30RMR	40MR	TR	
Zafar	4	3+	4	4	0;	1-	60S	80S	80S	10RMR	
Steklovidnaya-24	3+	3+	3+	4	0;	4	60S	60S	80S	30MS	Sr5
SIETE-CERROS-66	4	3+	3+	1	0;	0;	50MSS	40MSS	10RMR	TR	Sr6, Sr11
Krasnodarskaya-99	3+	3+	3+	3+	0;	0;	50S	80S	20MRMS	TR	
Jayhun	22+	22+	4	4	0;	4	30MR	30MR	70MSS	40MR	
IZ-80	4	4	11+	1-	0;	0;	50S	60S	20RMR	TR	38, +
AIKT-20	3+	3+	1+	;	0;	0;	40MS	50MS	20MR	TR	
OTUS TOBA-97	3+	3+	11+	0;	0;	0;	60S	60MSS	10RMR	TR	Sr31
PASTOR3/VORONA/CN079	3+	3+	11+	0;	0;	0;	20MSS	10MSS	20RMR	10RMR	
CMN82A.1294/2*	3+	3+	11+	1-	0;	0;	40MRMS	50MR	30MR	5RMR	
HUAVUN INIA	3+	3+	11+	3+	0;	0;	30MR	40MR	30MSS	TR	
Trakua Hatti	3+	3+	3+	3+	0;	1	40S	100S	50MSS	5R	
Murodi-2013	3+	3+	4	3+	0;	1+	60S	60MSS	30MS	5RMR	
CHEN/AE.SQ//WEAVER/3	3+	3+	4	3+	0;	1+	50S	50S	40MS	5RMR	

Table 5 continued

Accession	Sr seedling resistance				Sr Adult Plant Resistance				Sr gene postulation based on seedling and molecular marker	
	TTKST		TRTTF TKTTF		TTKSK + TTKST		TTKS + TTKST			MCCFC
	Rep. 1	Rep. 2	0	3+	0	3+	2010	2011		
Ganj	3+	3+	4	3+	0;	1 +/2+	40S	50S	40MR	5RMR
NAC/TH.AC// 3 * PVN/3/ MIR	3+	3+	4	3+	0;	1+	40S	60MSS	50MR	10RMR
Safedaki Pomir	1 + 2	22+	3+	3+	0;	4	-	-	60MSS	10RMR
Safedaki Ishkoshim	2	;2-	3+	3+	0;	4	-	-	60MSS	20MR
Babilo Pomir	4	;13	3+	3+	0;	;0	-	-	40MRMS	5R

Molecular marker analysis for *Sr6*, *Sr31* and *Sr38* are presented in Table 8.

Accessions exhibiting infection types of; 0 to 2 + were classified as resistant and those exhibiting infection types of 3–4 were classified as susceptible at the seedling stage. Adult plant rust severity was scored on a 0 to 100% basis and the infection responses on the size and type of uredinia where R (resistant), RMR (resistant to moderately resistant), MR (moderately resistant), MR-MS moderately resistant to moderately susceptible, MS (moderately susceptible), MS-S (moderately susceptible to susceptible) and S (susceptible). APR was tested only against races TTKSK + TTKST, MCCFC, and TKTTF

Table 6 Seedling infection types and field responses to stripe rust, and molecular marker analysis for *Yr9* and *Yr17*

Accession	Yr seedling resistance										Yr adult plant resistance		Yr gene postulation based on seedling and molecular marker		
	SE205/12	UK94/519	DK66/02	TJ01a/10	ER02/03	DK11/09	DK71/93	AF87/12	DK09/11	DK122/09	SE100/09	TK34/11		TJ01a/10	
	7	7	7	7	7	7	7	7	7	7	6	7		7	
Navruz	7	7	7	7	7	7	7	7	7	7	6	7	70MS	100S	
Sarvar	1	1	0	1	4	2	2	2	2	1	0	2	20MR	5R	
Vahdat	1	0	0	1	1	0	0	0	1	1	0	7	10MR	10R	
Yusufi	4	7	7	0	5	1	7	7	6	7	0	3	20MR	10R	
Isfara	0	0	0	1	0	0	1	7	1	0	0	7	20MR	10R	<i>Yr27,+</i>
Alex	7	7	7	0	5	1	1	0	6	4	0	3	40MR	40MS	<i>Yr9,+</i>
Oriyon	4	2	0	0	4	0	2	7	0	4	3	2	30MR	20MR	
Sadokat	3	0	7	2	4	0	1	0	1	2	0	3	30MR	10R	<i>Yr9,+</i>
Ziroat-70	2	7	0	7	7	0	0	0	5	5	0	7	60MS	60S	<i>Yr9,+</i>
Norman	5	7	0	7	7	0	3	0	7	0	0	7	50MS	100S	
Somoni	2	7	0	6	1	0	1	6	6	5	0	2	60MR	40MS	
Tacikar	5	0	0	7	6	0	1	0	6	6	0	7	50MRMS	40MS	
Ormon	4	0	7	4	1	0	1	6	1	0	0	7	10MR	10R	
Iqbol	6	0	0	7	0	0	1	0	6	5	0	2	40MRMS	100S	
Starshina	5	7	7	0	0	0	6	6	6	7	0	4	30MR	10R	
Shokiri	3	7	7	7	1	0	0	7	0	7	3	3	40MRMS	80S	
Fayzbaksh	4	0	1	1	3	0	1	1	0	4	0	4	40MR	20MR	
PRINA/STAR	7	7	7	7	4	6	5	0	6	7	0	1	40MR	60S	
BASRIBEY-95	7	7	7	7	1	0	7	7	6	7	0	7	70S	60S	
Jagger	6	7	0	1	0	0	1	0	6	7	0	4	5R	10R	<i>17,+</i>
Kaboi Panjakent	2	7	2	0	0	2	3	1	0	4	0	7	100S	20MR	
Surkhaki-5	2	7	3	0	0	0	2	0	0	4	0	7	100S	40MR	
Safedaki	7	7	7	7	7	6	7	6	6	7	0	3	20MR	70S	
Ishkoshimi															
Safedaki Pomir	7	7	7	7	4	2	7	6	6	5	2	3	20MR	70S	
Babilo Pomir	7	7	7	4	6	1	7	0	6	2	0	4	30MR	60MR	
Krasnodarskaya-99	7	0	7	7	1	0	7	1	6	7	0	3	30MRMS	50S	
Jeyhun	7	0	7	7	1	0	7	6	6	7	3	4	40MRMS	50S	
IZ-80	7	0	0	0	0	1	0	6	0	1	0	2	20MR	10R	<i>17,+</i>

Table 6 continued

Accession	Yr seedling resistance										Yr adult plant resistance		Yr gene postulation based on seedling and molecular marker		
	SE205/12	UK94/519	DK66/02	TJ01a/10	ER02/03	DK11/09	DK71/93	AF87/12	DK09/11	DK122/09	SE100/09	TK34/11		TJ01a/10	
AIKT-20	2	0	7	0	0	1	1	1	0	1	0	2	40MR	10R	
OTUS TOBA ₉₇	4	0	0	1	2	0	0	0	0	2	0	4	30MR	10R	Yr9, +
PASTOR/3/ VORONA/ CN079	3	0	0	7	0	0	0	0	6	6	0	3	20MRMS	100S	
CMN82A.1294/ _{2*}	2	0	7	2	0	0	0	0	1	2	0	7	40MRMS	20RMR	
HUAVUN INIA	4	0	7	2	0	0	0	0	1	2	0	7	60MSS	10RMR	

Molecular marker analysis for Yr9 and Yr17 are presented in Table 8

Accessions exhibiting infection types of 0–4 were classified as resistant, those exhibiting 5–6 as moderately susceptible, and those exhibiting 7–9 as susceptible at the seedling stage. Adult plant rust severity was scored on a 0 to 100% basis and the infection responses on the size and type of uredinia where R (resistant), RMR (resistant to moderately resistant), MR (moderately resistant), MR-MS moderately resistant to moderately susceptible, MS (moderately susceptible), MS-S (moderately susceptible to susceptible) and S (susceptible). APR was tested only against races TJ01a/10 and TR34/11

Table 7 Seedling infection types to leaf rust, and molecular marker analysis for *Lr26*

Accession	<i>Lr</i> seedling resistance									<i>Lr</i> gene postulation based on seedling and molecular marker
	TDBGG	TFBJQ	TNRJJ	MLDSD	MBDSB	TBBGJ	KFBJG	MHDSB	TCRKG	
Navruz	;1-	3	3+	3	33+	33+	3+	3+	33+	
Sarvar	0;	0;	;	3	;	;	;	;	;	
Vahdat	;	;	;	;	;	;1-	32+	11+	;1-	
Yusufi	;	;12	;1	33+	;	;1	3+	;	;1	
Isfara	;1-	33+	3	3	;	;11+	33+	1 + 2	12	
Alex	0;	;1	;	11+	;	;1-	33+	33+	33+	<i>Lr26</i>
Oriyon	;1-	;1	3	11+	;1+	;12-	3	3+	33+	
Sadokat	;	;1-	;	1+	1-	;1-	2 + 3	3	;1	<i>Lr26</i>
Ziroat-70	;	;12	;	22+	11+	;1-	2;3	3	;1	<i>Lr26</i>
Norman	;1-	3	3+	33+	3	;1	3+	33+	3+	
Somoni	2+	33+	3	22+	11+	33+	22+	2+	33+	
Tacicar	;1-	;12	3	33+	;1-	;11+	33+	22+	33+	
Ormon	;	;1/ 22 +	33+	3	12	3	3	33+	;11+	
Iqbol	;12	2	;12	11+	;1	11+	11+	33+	11 + 2-	<i>Lr16</i>
Starshina	;1-	3	3	3	22+	3	3+	3+	33+	
Shokiri	;	;1	;	22+	1+	3	22+	33+	;1	
Fayzbaksh	33+	22 + ;	3+	;1	;	;	;	3+	;1	
PRINA/STAR	;1-	2 + 3	1	;1	11+	1 + 2	;1-	3+	;11+	
BASRIBEY-95	;1-	33+	33+	2+	1+	;11+	3+	;1-	11+	
Jagger	3+	3 + ;	2+	2+	11+	;1	33+	33+	1 + 2	
Kaboi Panjakent	3+	3+	3+	3 + 4	3+	3 + 4	3 + 4	3 + 4	3+	
Surkhaki 5	3+	33+	3+	3+	3 + 4	3 + 4	3 + 4	3 + 4	3+	
Zafar	;1-	;1+	;	11+	;1	;1-	;1	;1	11+	
Steklovidnaya-24	;2	3 +	3+	12-	1 + 2	33+	3	3	12	
SIETE-CERROS-66	;12	3+	3+	3+	33+	3+	3+	3+	3+	
Krasnodarskaya-99	2 + 3	3+	3+	33+	2 + 3	3+	3+	3+	3+	
Jayhun	3+	33+	3+	3+	;	3+	3+	3+	3+	
IZ-80	33+	3+	;1	33+	3	3	;	3+	11+	
AIKT-20	0;	0;	3	2	;	;	;	;	;	
OTUS TOBA 97	;	;12-	;	;1	;1-	;	;	33+	;1-	<i>Lr16, Lr26</i>
PASTOR/3/VORONA	;	;12	;	;1	;1-	;1-	3	;	11+	
CMN82A.1294/2*	;	;12	;	;1+	;1-	;11+	3	33+	11+	
HUAVUN INIA	;	;12	0;	;12	;	;	11+	2 + 3	;1-	<i>Lr16</i>
Trakua Hatti	;2-	2	3+	3	2 + 3	2 + 3	33+	3+	11 + 2-	
Murodi-2013	;	;1+	;12	12	12+	12+	33+	;	1 + 2	

Table 7 continued

Accession	<i>Lr</i> seedling resistance									<i>Lr</i> gene postulation based on seedling and molecular marker
	TDBGG	TFBJQ	TNRJJ	MLDSD	MBDSB	TBBGJ	KFBJG	MHDSB	TCRKG	
CHEN/AE.SQ// WEAVER/3/ PASTOR	;1	22+	3	3	33+	33+	33+	33+	12	
Ganj	;	22+	;1	11+	;1	;1	;	;	;1–	
NAC/TH.AC// 3*PVN/3/ MIRL	;	22+	;	11+	;1	;1	;	;	;1–	
Safedaki Pomir	3	2+		;1			3+		3+	
Safedaki Ishkoshim				1+			3+	3+		
Babilo Pomir										Not tested

Molecular marker analysis for *Lr26* is presented in Table 8

Accessions exhibiting infection types of; 0 to 2 + were classified as resistant and those exhibiting 3–4 were classified as susceptible at the seedling stage

(120 bp), which is linked to the *Sr2/Yr30/Lr27* pleiotropic resistance gene, was amplified in 25 accessions with the *PBC* phenotype (score 1–3) (Table 8). Markers *XcsSr2* (172 bp) and *wMAS000005* did not detect the presence of *Sr2/Yr30/Lr27* in any accessions, while marker *Xgwm533* detected its presence in all accessions with the *PBC* phenotype (score 1–3) (Table 8). Initially, all accessions with and without the *LTN* phenotype (score 0–3) were screened with the *csLV34* (150 bp) marker. In thirteen cases, this marker indicated the presence of the *Lr34/Yr18/Sr57* APR resistance genes, which were subsequently validated by the *wMAS000003* Kompetitive Allele Specific PCR (KASP) marker (Table 8). Use of KASP marker *wMAS000005* positively detected the presence of *Sr2/Yr30/Lr27* in Hope and CS-Hope DS 3B, but failed to do so in the Tajik accessions; thus, this KASP marker is located in the “Hope and CS Hope DS 3B” allele. The *Xwmc364* (207 bp) marker was used on all accessions to detect the presence of *Yr2*, but all of them amplified a 201 bp marker allele, indicating the absence of *Yr2*.

Discussion

In this study, we identified the presence of major-effect (seedling) and pleiotropic APR genes conferring resistance against three important rust diseases, i.e. stem rust, stripe rust and leaf rust pathogens in wheat cultivars, landraces and advanced breeding lines that are widely cultivated and used in the national wheat breeding program in Tajikistan. The major-effect resistance genes identified by seedling and adult plant responses, and molecular marker analysis were *Sr5*, *Sr6*, *Sr11*, *Sr31/Yr9/Lr26*, *Sr38/Yr17/Lr37*, *Yr27*, and *Lr16*. Additionally, the pleiotropic APR genes of *Sr2/Yr30/Lr27* and *Lr34/Yr18/Sr57* were also identified based on the *PBC* and *LTN* phenotypes in the field and confirmed with linked molecular markers. The APR gene *Lr37* was detected by a molecular marker (*VENTRIUP/LN2*), which is completely linked with the *Sr38/Yr17* genes. In addition, pedigree information (<http://wheatpedigree.net/>) also was used to augment gene postulation data. A number of the wheat accessions showed resistance to all races of the three rusts used in this study, and their infection type pattern did not correspond to the avirulence/virulence profiles of the races as identified on the differential accessions. Therefore, the resistance genes present in these accessions could not be postulated. We conclude that

Table 8 Molecular marker analysis to determine the presence and absence of postulated *Sr*, *Yr* and *Lr* seedling and APR genes

Accession	PBC	LTN	<i>Sr2Yr30/Lr27</i>		<i>Yr2</i>	<i>Sr6</i>	<i>Sr31Yr9/Lr26</i>		<i>Sr38/Yr17/Lr37</i>	<i>Lr34Yr18/Sr57</i>		<i>Sr</i> , <i>Yr</i> and <i>Lr</i> gene	
			<i>Xgwm533</i>	<i>XcsSr2</i>			<i>wMAS000005</i>	<i>Xwmc364</i>		<i>Xcfd43</i>	<i>Xscm9</i>		<i>Xtag95</i>
Navruz	0	0	-	-	-	-	-	-	-	-	-	-	<i>Sr2Yr30/Lr27</i>
Sarvar	3	0	+	-	-	-	-	-	-	-	-	-	<i>Sr2Yr30/Lr27</i> , <i>Lr34/Yr18/Sr57</i>
Vahdat	2	3	+	-	-	-	-	*	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Lr34/Yr18/Sr57</i>
Yusufi	3	1	+	-	-	-	-	*	-	-	-	-	<i>Sr2Yr30/Lr27</i>
Isfara	1	3	+	-	-	-	-	*	-	+	+	+	<i>Lr34/Yr18/Sr57</i>
Alex	2	3	+	-	-	-	+	+	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Sr31/Yr9/Lr26</i> , <i>Lr34/Yr18/Sr57</i>
Sadokat	1	3	+	-	-	-	+	+	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Sr31/Yr9/Lr26</i> , <i>Lr34/Yr18/Sr57</i>
Ziroat-70	2	3	+	-	-	-	+	+	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Sr31/Yr9/Lr26</i> , <i>Lr34/Yr18/Sr57</i>
Iqbol	1	3	+	-	-	-	-	-	-	+	+	+	<i>Lr34/Yr18/Sr57</i>
Shokiri	1	3	+	-	-	-	-	*	-	+	+	+	<i>Lr34/Yr18/Sr57</i>
PRINA/STAR	1	3	+	-	-	-	-	*	-	+	+	+	<i>Lr34/Yr18/Sr57</i>
Jagger	0	0	-	*	-	-	-	-	+	-	-	-	<i>Sr38/Yr17/Lr37</i>
SIETE-CERROS-66	1	1	+	-	-	-	-	*	-	-	-	-	<i>Sr6</i>
IZ-80	1	1	+	-	-	-	-	*	+	-	-	-	<i>Sr38/Yr17/Lr37</i>
OTUS TOBA 97	2	3	+	-	-	-	+	+	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Sr31/Yr9/Lr26</i> , <i>Lr34/Yr18/Sr57</i>
PASTOR/3/VORONA/CN079	2	1	+	-	-	-	-	*	-	-	-	-	<i>Sr2Yr30/Lr27</i>
HUAVUN INIA	1	3	+	-	-	-	-	*	-	+	+	+	<i>Lr34/Yr18/Sr57</i>
Murodi-2013	2	2	+	-	-	-	-	*	-	+	+	+	<i>Sr2Yr30/Lr27</i> , <i>Lr34/Yr18/Sr57</i>

Table 8 continued

Accession	PBC	LTN	Sr2/Yr30/Lr27		Yr2	Sr6	Sr31/Yr9/Lr26		Sr38/ Yr17/Lr37	Lr34/Yr18/Sr57		Sr, Yr and Lr gene
			Xgwm533	XcsSr2			Xwmc 364	Xcfd43		Xscm9	Xiag95	
CHEN/AE.SQ// WEAVER/3	2	2	+	-	-	-	-	-	-	+	+	Sr2/Yr30/Lr27, Lr34/ Yr18/Sr57
Ganj	2	1	+	-	-	-	-	-	-	-	-	Sr2/Yr30/Lr27
NAC/TH.AC// 3 * PVN/3/ MIRLO	2	1	+	-	-	-	-	-	-	-	-	Sr2/Yr30/Lr27
Safedaki Ishkoshim	0	2	-	*	-	-	-	-	-	+	+	Lr34/Yr18/Sr57

+ Presence – or absence of the genes according to corresponding marker alleles, and * indicates molecular markers not applied

wMAS000005 and wMAS000003 KASP markers were obtained from the MAS protocols: <https://maswheat.ucdavis.edu/protocols/index.htm>

The presence of PBC (Pseudo-Black Chaff) and LTN (Leaf Tip Necrosis) phenotypes were assessed using 0–4 scale in the field, and in addition were confirmed with linked molecular markers

these accessions carry previously described gene(s) in combinations or new genes. To elucidate the genetic basis of resistance in these widely resistant accessions, biparental crosses, allelism tests and/or additional phenotyping tests with a wider array of rust races should be implemented (Li et al. 2015; Randhawa et al. 2015). The resistance gene *Sr5* in Navruz and *Sr6* and *Sr11* in Siete-Cerros-66 were identified in this investigation. Navruz is commonly used as a control in all wheat breeding nurseries and official trials (Husevov et al., 2015), and Siete-Cerros-66 has been cultivated by Tajik farmers since 1970 (Muminjanov et al. 2008). *Sr5*, *Sr6*, and *Sr11* have been effective and valuable stem rust resistance genes; however, *Pgt* races with virulence for these genes are spreading in many wheat growing regions worldwide (Singh et al. 2015). Combinations of seedling and APR genes (i.e. *Sr2/Yr30/Lr27*, *Sr31/Yr9/Lr26*, *Lr34/Yr18/Sr57*, *Lr16* etc.) were also present in some of the accessions (Table 8), thus being promising sources for improved resistance to rusts in Tajik breeding programs. Gene pyramiding using the pleiotropic APR genes of *Sr2/Yr30/Lr27* and *Lr34/Yr18/Sr57* in a combination with seedling resistance genes in several wheat breeding programs has provided durable rust resistance (Ellis et al. 2014).

Four wheat accessions (Alex, Sadokat, Ziroat-70 and Otus Toba 97) were identified as carrying the *Sr31/Yr9/Lr26* resistance genes. Accessions possessing this gene complex are known to have the 1BL.1RS wheat-rye translocation, originating from Petkus rye (Friebe et al. 1996). The *Sr31/Yr9/Lr26* complex is very common in wheat accession due to the wide utilization of the wheat cultivars Kavkaz and Aurora in CIMMYT breeding programs worldwide (Rajaram et al. 1983). The individual genes in this complex have been overcome by virulent races of *Pgt*, *Pst* and *Pt*, respectively, in various wheat growing regions (Pretorius et al. 2000; Chen et al. 2010; Huerta-Espino et al. 2011; Wellings 2011). *Sr31* has provided durable resistance to stem rust for more than 30 years, and still remains an effective source of resistance to many *Pgt* races with the exception of those in the Ug99 race group. Races of *Pst* with virulence for *Yr9* have been reported from all major wheat production areas in Tajikistan based on trap nurseries and race surveys (<http://wheatrust.org>). Additionally, virulence against the leaf rust resistance gene *Lr26* is common in Tajikistan (Kolmer and Ordoñez 2007) and many

other parts of the world. Thus, although the resistance genes on the 1BL.1RS translocation do not confer a high degree of resistance towards new races of the rust pathogens, wheat accessions carrying this translocation are cultivated throughout the country. Two wheat cultivars widely cultivated in Tajikistan (Jagger and IZ-80) were identified as possessing the *Sr38/Yr17/Lr37* gene complex. Previous studies have characterized the *Sr38/Yr17/Lr37* locus as a translocation of chromosome 2NS from *Triticum ventricosum* replacing the homoeologous region of 2AS in *Triticum aestivum*; thus, this translocation confers resistance against a range of races of *Pgt*, *Pst* and *Pt* (Helguera et al. 2003). The presence of *Yr27* in Isfara and *Lr16* in Iqbol, OTUS TOBA 97 and Murodi-2013 were postulated. *Lr16* is known as an effective source of leaf rust resistance in wheat (Kolmer and Hughes 2013) and should provide stable resistance when pyramided with *Lr27*, *Lr34*, and *Lr37* in the Tajik breeding program for the developing resistant wheat cultivar.

Both phenotyping (using 0–4 scale for the *PBC* phenotype) and genotyping (using the *Xgwm533*, *XcsSr2*, and *wMAS000005* markers) were applied for detection of the *Sr2/Yr30/Lr27* APR genes; thus, only 25 accessions with the *Xgwm533* marker (score 1–3 for the *PBC* phenotype) were identified. However, only eleven accessions were considered to truly possess the *Sr2/Yr30/Lr27* APR genes based on the *Xgwm533* marker and *PBC* phenotype, i.e. score of 2 for medium pigmentation and 3 for high pigmentation (Table 8). The *PBC* phenotype is known to be associated with the *Sr2/Yr30/Lr27* gene complex, although its expression is sometimes variable due to both the genotype and environment (McFadden 1930). In addition, the *PBC* phenotype is genetically associated with several quantitative trait loci (QTL) on the chromosome arms 2DS, 3BS, 4AL, and 7DS (Juliana et al. 2015). With respect to the molecular markers in the present study, *XcsSr2* and *wMAS000005* were only able to identify the *Sr2/Yr30/Lr27* APR genes in the Hope and CS-Hope DS 3B lines, while the *Xgwm533* marker positively detected the gene complex in 25 accessions. These results corroborate previous investigations that showed no perfect match between amplification of the *XcsSr2/wMAS000005* and *Xgwm533* markers in various accessions (Mago et al. 2011; Pretorius et al. 2012). Thus, in the present study, the *PBC* phenotype, with scores of 1–3 in 25 accessions, showed a high degree of correlation with

amplification of the *Xgwm533* marker. However, the *Xgwm533* marker may also positively amplify even when *Sr2/Yr30/Lr27* is not present in certain wheat accessions (Spielmeyer et al. 2003; Mago et al. 2011). Hope and CS-Hope DS 3B (172 bp) have been shown to carry *Sr2/Yr30/Lr27* based on studies using the *XcsSr2* marker (Mago et al. 2011). Initially, *Sr2* was reported to be linked with the *PBC* phenotype in the cultivar Hope; thus, this phenotypic trait has become a valuable selection trait for wheat breeders in the field (McFadden 1930). The KASP marker *wMAS000005* identified the allele in Hope and CS-Hope DS 3B, thereby identifying the presence of *Sr2/Yr30/Lr27*. However, this marker failed to amplify any signal in the Tajik wheat, thus indicating the absence of *Sr2/Yr30/Lr27*. Molecular markers *csLV34* and *wMAS000003* successfully identified the presence of the *Lr34/Yr18/Sr57* APR genes in 13 accessions; therefore, these markers can be reliably used with *LTN* phenotype for assessing APR genes. In addition to gene postulation, the *Xwmc364* marker can be used to confirm the presence or absence of *Yr2* gene. This *Xwmc364* (207 bp) marker positively confirmed *Yr2* in the Kalyansona and Heines VII differential accessions, but amplified the 201 bp or null allele in all of the Tajik accessions, suggesting the absence of *Yr2*.

In this study, we demonstrated that some of the evaluated accessions carry seedling and pleiotropic APR resistance genes against all the used rust races. Thereby, our results show that some of the wheat accessions may be used as a diverse source of rust resistance. The gene postulation, together with the use of molecular markers, successfully applied to detect the presence of known seedling and APR genes in some of the evaluated accessions. Moreover, the genetic basis of resistance in some accessions should be characterized through other genetic analyses because gene postulation and molecular markers failed to do so in this study. In the meantime, these accessions can be used by the national wheat breeding program in Tajikistan as crossing parents to develop new varieties with durable rust resistance.

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Author's contribution MR designed the study and was responsible for conducting all experiments. MO and HM developed some of the wheat accessions and provided seeds. MNR and BJS supervised the stem rust seedling and adult plant tests in the University of Minnesota and USDA-ARS-CDL in USA. MSH supervised the stripe rust seedling test in the GRRC in Denmark. KN supervised the stem rust and stripe rust seedling and adult plant tests in the RCRR in Turkey. MNR, BJS, and EJ supervised the overall study. MR wrote the manuscript and all authors contributed to writing and editing the manuscript.

Compliance with ethical standards

Conflict of interest All authors have no conflict of interest.

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