LOCAL STEM RUST VIRULENCE IN PAKISTAN AND FUTURE BREEDING STRATEGY

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Abstract

Evolution and spread of stem rust race Ug-99 has created an alarming global situation. Majority of the CIMMYT germplasm protected by gene *Sr31* fell susceptible to this catastrophic strain. Like other parts of the world stem rust of wheat was very successfully controlled in Pakistan by the introduction of resistant germplasm during and after green revolution. In 2001 stem rust reappeared and hit many commercial varieties in the province of Sindh. Its sporadic infections were recorded in 2005 summer crop of Kaghan followed by 2005-06, 2006-07, 2007-08 and 2008-09 spring crops of Sindh. In a regional scenario, where by wheat crop of the whole CWANA region was threatened by the Ug-99, it was thought that the crop is attacked by Ug-99. To date data from the stem rust TRAP nurseries and analysis of the disease samples is presented here to clarify the situation about presence/absence of Ug-99 in Pakistan. Commercial cultivars resistant to local stem rust race and genes effective against local and Ug-99 races are identified and their implications to improve stem rust resistance of Pakistani cultivars are discussed.

Introduction

Stem rust of wheat caused by Puccinia graminis f. sp. tritici (Pgt), responsible for heavy yield losses in wheat crop, has successfully been controlled throughout the world by introduction of semi-dwarf resistant cultivars during 1960s and 1970s and there has been no report of significant loss since then. Resistance in majority of the wheat cultivars sown in this region is based on gene Sr31 which is introgressed into bread wheat from Secale cereale cv. Petkus (Rye) (McIntosh et al., 1995). These lines gave significant yield advantage in addition to powdery mildew, stem leaf and stripe rust resistance. Breakdown of stem rust resistance gene Sr31 with the introduction of pgt race Ug-99 in Uganda during 1999 (Pretorius, 2000) created an alarming situation throughout the world as the gene was protecting 80% of the developing country's wheat lines derived from the CIMMYT germplasm. Virulence to this gene was subsequently reported along Yr9 path following its first report from Uganda. On its way along the path it acquired virulences to the genes of importance and three variants were reported from Ethiopia (Jin et al., 2008; Jin et al., 2009). Khanzada (2008) reported stem rust infection in Pakistan on mega cultivars during 2000-01 wheat cropping cycle. Sporadic infection of stem rust were recorded in 2005 summer crop of Kaghan followed by 2005-06, 2006-07, 2007-08 and 2008-09 spring crops of Sindh. Appearance of stem rust on mega wheat cultivars of Pakistan was alarming under the prevailing scenario of the region where by Ug-99 was spawning new strains and was moving ahead towards Pakistan along Yr9 Path. Moreover all Pakistani wheat cultivars screened under Kenyan field conditions were susceptible to Ug-99 (Anon., 1995). Analysis of the situation was realized as prime factor in adaptation of breeding strategy to incorporate genes resistant to Ug-99 in Pakistani cultivars. Our analysis indicated that the virulence found in Pakistan did not match Ug-99 but still had the capability to attack genes resistant to Ug-99. This paper discusses virulence patterns of the local stem rust race found in Sindh province of Pakistan and its importance to adopted breeding strategy to combat Ug-99 in this region.

Material and Methods

Stem rust TRAP Nursery, commercial cultivars and disease sampling: A TRAP nursery consisting of 39 stem rust isogenic lines was planted at Thatta and Matli. Details of the lines in this nursery are given in Table 1. The nursery was planted in 1m row 30cm apart from each other. Morocco was added as susceptible check and spreader.

Seventy one commercial wheat cultivars, most of them with known susceptibility to Ug-99 were cultivated at Matli, Sakrand and Karachi. Third stem rust screening nursery $(3^{rd}$ SRSN) received from CIMMYT was planted at Thatta and Karachi for field screening. Morocco was added as susceptible check and spreader.

Survey of the wheat crop in districts Karachi, Thatta, Tando Muhammad Khan, Badin, Hyderabad, Mirpurkhas, Umerkot, Larkana, Sukkur, Nausheroferoze and Dadu was conducted from March 11-20, 2009 to see distribution of stem rust and collect disease samples. Disease samples were collected from farmer's fields and research stations. Some disease samples were also received from Wheat Research Institute Bahawalpur and Islamabad. Disease intensity on commercial cultivars was recorded at farmer field.

Virulence analysis and seedling screening: Virulence analysis of the disease samples was conducted under glasshouse conditions. Urediospores washed from disease samples in petroleum spirit were re-suspended in mixture of mineral oil and petroleum spirit (ratio 1: 3 v/v) before spraying them with fine atomizer on 10 day old seedlings of the susceptible cultivar Morocco grown in disposable pots. Inoculated plants were left in shade for two hours, to evaporate oil, before transferring them to growth room. The growth room was set at $18\pm1^{\circ}$ C temperature, 99% relative humidity and 20 hours dark followed by 4 hour florescent light. The plants were then moved to glasshouse set at 18- 20° C temperatures. On the appearance of pustules on 10^{th} day all the leaves except the ones with isolated pustule were clipped, plants were washed under running water and placed in isolation chambers to sporulate the pustules. On sporulation the single pustule inoculums were multiplied on the newly grown leaves of same plant using standard procedure (Knott, 1989). Single pustule inoculums were collected for each pustule separately and inoculated on ten day old seedlings of the tester host set planted in 38 x 46 x 10 cm tray. The tester set was composed of 20 North American stem rust differentials (Roelfs & Martens, 2007) and morocco as susceptible check. Differentials in trays were inoculated with spore suspension in mixture of mineral oil and petroleum sprit (1:3 v/v)by method mentioned above. Growth room and glasshouse incubation conditions were also same as mentioned above. Stem rust data for seedling infection types described by Stakman et al., (1962) was recorded on 10th day of inoculation when pustules were sporulating on susceptible cultivar morocco. Seedling screening of the selected SRSN lines was conducted using conditions and methods mentioned above.

	Č		2008-09		2005	2006
Sr. differential lines	Cene	Thatta	Matli	Kaghan	Kenya	Kenya
Morocco	check	80S	40S	I		I
PBW343	Sr31	40MSS	I	0	I	I
Cham-8	Sr31	10MSS	I	0	I	I
LcSr24Ag	Sr24	20MS	I	0	10MR	20R-MR
Sr24(Agent)/9*LMPG	Sr24	30MS	5MS	0		10R-MR
Sr36(CI 12632)/8*LMPG	Sr36	90S	40S		5MR	0
W2691SrTt-1	Sr36	30S	5S	0	I	I
Cook	Sr36	10MS	TS	0	I	I
Eagle	Sr26	20MS	I	0	40MR-MS	SR
Sr26/9*LMPG	Sr26	40S	I	I	20MR-MS	
Super Seri	Sr25	30S	0	ST	I	I
LcSr25Ars	Sr25	20MS	55	0	60MR-MS	20MR-MS
Coroong	Sr27	0	I	0	20MR	10R
Pollmer-2.1(Triticale)	Sr27	5 RMR	I	I	I	I
Hindhab	Sr2	40MSS	I	I	I	I
Pavon76	Sr2 complex	10MS	5MS	ST	I	I
Buck Buck	Sr2+Sr23	40MSS	5MS	0	I	I
ISr6-Ra	Sr6	30S	20MSS	0	80S	70S
W2691Sr9b	Sr9b	30S	20S		100S	70S

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			2008-09		2005	2006
эг. ишегелиагшеэ	Celle	Thatta	Matli	Kaghan	Kenya	Kenya
Vernstein	Sr9e	20S	10MS	0		I
St464Sr13	Sr13	TS	5MS	0	40MR-MS	40MS
Combination VII	Sr17(+13)	40S	80S	I	40MR-MS	40MS
Sr22TB	Sr22	10MS	TMS	0	10R	20R-MR
Pusa 4/Etoile de Choisy	Sr29	$\mathbf{S0L}$	20MSS	I	I	Ι
BtSr30Wst	Sr30	70S	40S	I	809	S-SM09
CnsSr32As	Sr32	40MS	20MS		30R	10R
RL5405	Sr33	40MSS	20MSS	I	30MR	30MR-MS
Mq(2)5*G2919	Sr35	10MS	70S	I	10MR-MS	IR
W2691SrTt2	Sr37	40S	80S			5R/60S
RL6081	Sr38	30MS	20MS	I	50S	40S
RL6082	Sr39	20MS	20MS	I	5R/70S	5R/10MS
RL6088	Sr40	40S	30MSS		SR	10R
Taf-2	Sr44	80S	40S		10R-MR	10R
Goldenball dervi	Srdp-2	40S	10S	I	I	I
W2691SrGtGt	SrGt	$\mathbf{S06}$	40MSS		I	I
CnSSrTmp	SrTmp	202	30MSS	0	40MS	30MR-MS
Bt/Wld	SrWld-1	202	809	I	80S	S09
Chris	Sr7a, Sr12, Sr6	20MS	5MS	I	I	I
Line A Seln	SrI4	5MS	20MS	5MS	I	I
W2691 Sr28 kt	Sr28	70S	30MSS	TS	40MS-S	40MR-MS

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Results

Stem rust TRAP Nursery and response of commercial cultivars: Results of the TRAP nursery are given in Table 1. Susceptible filed response was recorded for genes Sr13, 17, 25, 36, 37 and 44 in Matli and Thatta while genes Sr22, 32, 33, and 39 were moderately susceptible at these locations. Gene Sr35 was highly susceptible at Matli and moderately susceptible at Thatta while genes Sr28 and Tmp were moderately susceptible at Matli and susceptible at Thatta. Virulence for Sr6, 9b, 30, 38 and Wld-1 was absent in Kaghan.

Field screening results of the commercial cultivars are given in Table 4. Commercial cultivars Faisalabad 85, Kohinoor83, LU-26, Pasban-90, Shaheen 94, Shahkar 95, Bahawalpur 97, Abadgar 93 and Tatara which are highly susceptible to Ug-99 were immune at Karachi, Tandojam, Sakrand and Matli indicating absence of Ug-99 at these locations. Many other cultivars including Bakhtawar, Inqilab 91, Blue silver, Chakwal 86, Sindh 81, Zargoon and Faisalabad83, which are highly susceptible to Ug-99 were either immune or moderately susceptible at these locations.

Survey and sample collection: Maximum stem rust disease intensity of 40S-80S was recorded on wheat varieties Kiran, Sarsabz, unidentified cultivars named as Hira (named by seed companies), and other mixed material in Kunri, Jhuddo and neighboring fields. Disease intensity at different locations is given in Table 2. Barley infested with stem rust was observed in Districts Badin and Tando Muhammad Khan. Some pustules were observed on local variety Thori in Rattu Dero and neighboring fields of District Larkana. Disease sample were collected from Thatta, Matli and Mirpur Bathoro for further analysis.

Virulence analysis of the disease samples: Two virulence/avirulence combinations were identified in 24 stem rust isolates. These combinations differed from each other in response to *Sr9e*, *Sr24* and *SrTmp* genes and their virulence/avirulence formulae are given in Table 3.

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Location	Intensity	Variety
District Umerkot: Taluka Kunri	40S-80S	Kiran, Sarsabz and companies material Hira,
		and other mixture
Jhuddo and neighboring fields	40S-80S	Kiran, Sarsabz and other mixture
Distt. Thatta:	5S-60S	Companies seed Hira, Rustam and barley
Gharo, Sijawal, Mirpur Bathoro		
and Jhoke Sharif.		
Distt. Badin:	5S-60S	Companies Rustam seed, other local varieties
Matli, Talhar, Tando Bago and		and barley
neighbours field		
Distt. Tando Muhammad Khan:	5S-60S	Local wheat varieties and barley
Bulri Shah Karim and neighbours		90S Baley
field		-
District Hyderabad i.e. Kisano	10 S	Kiran
Mori		
Distt. Larkana i.e. Rattu Dero and	Trace	local variety Thori
neighbouring fields		-
Distt. Nawabshah and	Trace	-
Nausheroferoze		

Table 2. Intensity of stem rust disease in different districts of Sindh.

S. No.	Location	Virulence/avirulence combinations
1.	Thatta	Sr5,6,7b,9a,9b,9d,9g,10,11,17,21,30,36,Tmp,McN/Sr8a,9e, 24, 31
2.	Matli	Sr5,6,7b,9a,9b,9d,9g,10,11,17,21,30,36,Tmp,McN/Sr8a, 9e, 24, 31
		Sr5,6,7b,9a,9b,9d,9e,9g,10,11,17,21,24,30,36,Tmp,McN/Sr8a, 31
3.	Mirpur mathilio	Sr5,6,7b,9a,9b,9d,9g,10,11,17,21,30,36,Tmp,McN/Sr8a, 9e, 24, 31
4.	Bahawalpur	Sr5,6,7b,9a,9b,9d,9g,10,11,17,21,30,36,Tmp,McN/Sr8a, 9e, 24, 31
5.	Islamabad	Sr5,6,7b,9a,9b,9d,9g,10,11,17,21,30,36,Tmp,McN/Sr8a, 9e, 24, 31

Table 3. Virulence/avirulence combinations identified among isolates from Pakistan.

Discussion

Stem rust infection found earlier in limited coastal area of Sindh spread to Distt. Umarkot, Taluka, Jhuddo, Kunri, Thattta, Sajawal, Gharo, Mirpur Bathoro, Jhoke Sharif, Matli, Talhar, Tando Bhago, Tando Muhammad Khan, Larkana, Nawabshah in Sinsh, Bahawalpur in Punjab and in Islamabad. Most popular cultivar of Sindh are susceptible to the virulence found in these areas. Field resistances in numerous Ug-99 susceptible commercial cultivars at Karachi, Tandojam, Sakarand and Matli indicate absence of Ug-99 in these locations. Moreover susceptible filed response of Sr13, 25, 36, 37 and 44 in Matli and Thatta indicate a virulence pattern different from Ug-99. Glasshouse virulence analysis revealed that stem rust samples from all locations lacked Sr31 virulence. Presence of Ug-99 is also nullified by data from Kaghan where virulence for Sr31 was lacking. Among other stem rust resistant genes Sr24, for which virulence is detected in new strains of Ug-99, showed MS type of reaction under field conditions in Thatta and Matli. Virulence analysis confirmed Sr24 virulent isolates in Matli which may be attacking this gene in Thatta as well. This gene in combination with other genes is still effective against local and Ug-99 races as lines of third stem rust screening nursery (SRSN) carrying this gene in combination with SrTmp and Sr36 are showing very good field resistance in Thatta, seedling resistance against Sr24 virulent isolate from Matli and isolates from Bahawalpur (Table 5). Gene Sr25 which is closely linked with Lr19 was thought to become more susceptible under high temperatures (Gough & Markle 1971). Genetic stocks with 7D-Ag translocations carrying this genes has never been cultivated in these areas in past but virulence to this gene was detected in Nilgiri Hills of India in 2007 (Singh et al., 2008) and coastal areas of Sindh in 2008 (Mirza et al., 2009). Sr25 gave susceptible field response at Thatta and Matli. Promising level of field resistance at Thatta and their seedling resistance with isolates from Matli show that SRSN lines postulated for Sr25 have additional genes in them (Table 5).

Gene *Sr27* showed field resistance in Thatta. This is another gene of potential importance for which Ug-99 lacks virulence. Incorporation of this gene in Pakistani cultivars resistant to local race can enhance their resistance to Ug-99. *Sr6*, *9b*, *30*, *38* and *Wld* are the genes for which Ug-99 carries virulence. All isolates analyzed under glasshouse conditions from Sindh and Punjab were virulent to *Sr6*, *Sr9b*, *Sr30* and *Wld*. The use of these genes should not be encouraged in Pakistani cultivars. Stem rust resistant genes *Sr13*, *Sr14* from *Triticum turgidum*, *Sr28*, *Sr29* from *T. aestivum*, *Sr33* from *T. tauschii*, *Sr36*, *Sr37* from *T. tempoheevi* and *Sr35* from *T. monococum* were of special interest to breeders after the evolution of Ug-99 (Singh *et al.*, 2006). *Sr14* has field resistance at Matli, Thatta and Kaghan while *Sr28* was susceptible at these locations. Virulence for *Sr13* & *Sr37* was found in past among isolate collections from

Ent #.	Line		Stem rust	response in	
сш #.	Line	Karachi	Sakrand	Matli	Kenya (2005
1.	Bakhtawar 92	0	0	5MSS	20S
2.	Blue silver	0	0	5MSS	20S
3.	Chakwal 86	0	0	5MSS	40S
4.	Sind-81	0	0	5MSS	40S
5.	Zarghoon	5MSS	0	5MSS	60S
6.	Faisalabad 83	0	0	20S	30S
7.	Faisalabad 85	0	0	0	40S
8.	Inqilab 91	0	5RMR	10MSS	50S
9.	Kaghan 93	0	5RMR	10MSS	30S
10.	Morocco	0	20MSS	30S	
11.	Kirin 95	0	0	10MSS	30S
12.	Kohinoor 83	0	0	0	40S
13.	LU-26	0	0	0	40S
14.	Nowshera 96	0	0	TMS	30S
15.	Parwaz 94	0	0	5MSS	10S
16.	Pasban 90	0	0	0	20S
17.	Mexipak 65	0	5MS	5MSS	
18.	Punjab 96	0	0	5MSS	30S
19.	Sariab-92	0	0	TMS	20S
20.	Morocco	0	0	30S	
21.	Sarsabz	0	0	5MSS	40S
22.	Shaheen 94	0	0	0	30S
23.	Shahkar 95	0	0	0	10S
24.	Soughat 90	0	0	5MSS	20S
25.	Tandojam 83	0	5MS	5MSS	20S
26.	SH-2002	0	0	20MSS	20S
27.	Pak 81	0	0	5MSS	20S
28.	Bahaw-97	0	0	0	40S
29.	MH-97	0	0	70S	30S
30.	Morocco	0	0	10MSS	_
31.	Kohistan 97	0	0	20MS	40S
32.	Kohsar 95	0	0	5MSS	_
33.	Rohtas 90	0	0	0	_
34.	Suleman 96	0	0	10MSS	_
35.	WL 711	0	0	5MSS	20S
36.	Zardana	0	0	5MSS	30S
37.	Abadgar 93	0	0	0	10S
38.	Anmol-91	0	0	5MSS	20S
39.	Bahawal-2000	0	0	5RMR	40S

		Table 4 (Co	nt'd.).		
Ent #.	Line		Stem rust	response in	
EIII #.	Line	Karachi	Sakrand	Matli	Kenya (2005)
40.	Morocco	0	58	40S	_
41.	Bahkhar-2002	0	0	10MSS	308
42.	Fakhr-e-Sarhad	70S	10MSS	10MSS	208
43.	Marvi-2000	0	0	TMS	208
44.	Mehran-89	0	5MS	TMS	_
45.	Soorab-96 (Barley)	0	Missing	TMS	208
46.	Tatara	0	0	0	208
47.	Takbeer	0	10MS	60S	208
48.	AS-2002	0	5MS	5MSS	10S
49.	Iqbal 2000	0	0	5MSS	10S
50.	Morocco	0	5MSS	30S	
51.	Auqab-2000	0	0	30S	208
52.	Chakwal-97	0	0	0	_
53.	Durum-97	0	0	TMS	40S
54.	Watan 94	0	0	10MSS	50S
55.	Moomal 2002	0	0	10S	30S
56.	Zarlashata	0	0	TMS	40S
57.	GA-2002	0	0	5MSS	208
58.	Wafaq-01	0	0	5MSS	40S
59.	Margalla-99	0	0	5MS	208
60.	Morocco	0	0	20S	_
61.	Manthar-3	0	0	0	308
62.	Saleem 2000	0	0	5MS	_
63.	Khyber 87	0	0	0	_
64.	Pirsabak 2004	0	0	0	_
65.	Pirsabak 2005	0	0	TMS	_
66.	Punjnad-1	0	0	0	_
67.	Darawar-97	0	0	0	_
68.	V-87094 (Wattan)	0	0	TS	_
69.	Shafaq 2006	0	0	5MSS	_
70.	Morocco	0	0	20S	_
71.	Sehar 2006	0	0	30MSS	_
72.	Bhittai	0	0	5MSS	_
73.	Chakwal-50	0	0	0	_
74.	Saussi	0	0	5MSS	_
75.	Lasani-08	5MSS	0	10MSS	_
76.	Meraj-08	0	0	5MSS	_
77.	Fareed-06	0	0	0	_
78.	Faisalabad-08	208	0	20MSS	_

Table 4 (Cont'd.).

Entry #	Parentage/Cross name	*Postulated genes on the basis of Kenyan	*2008 filed screening at	Stem ru screer Pakis	Stem rust field screening in Pakistan at	Gl screen with in	Glasshouse screening Seedling with inoculum from
		data 2000-07 & 2007	Nenya	Thatta	Karachi	Matli	Bahawalpur
6021	PFAU/MILAN/3/SKAUZ/KS94U215//SKAUZ	Sr-Unknown	20 RMR	5MSS	0	•••	3C ⁼
6036	WHEAR/VIVITSI//WHEAR	Sr25	20 RMR	5MS	0		3C ⁼
6037	WHEAR/VIVITSI//WHEAR	Sr25	15 RMR	10MS	0		3C ⁼
6044	WHEAR/KIRITATI/3/C80.1/3*BATAVIA//2*WBLL1	Sr25	30 MR	10MS	0		·
6045	WHEAR/KIRITATI/3/C80.1/3*BATAVIA//2*WBLL1	Sr25	30 RMR	10MS	0		,
6046	WHEAR/VIVITSI/3/C80.1/3*BATAVIA//2*WBLL1	Sr25	20 MR	10MS	0	0	,
6047	WHEAR/VIVITSI/3/C80.1/3*BATAVIA//2*WBLL1	Sr25	20 MR	5MS	0	•••	
6056	HUW234+LR34/PRINIA//PFAU/WEAVER	SrHuw234	40 M	5MS	0	•••	
6909	SUNCO//TNMU/TUI	Sr24+Sr36	1 R	5MS	0	0	
6070	SUNCO//TNMU/TUI	Sr24+Sr36	1 R	5MS	0	0	
6071	SUNCO//TNMU/TUI	Sr24+Sr36	1 R	TMR	0	0	
6072	SUNCO//TNMU/TUI	Sr24+Sr36	1 R	0	0	0	
6084	SHARP/3/PRL/SARA//TSI/VEE#5/5/VEE/LIRA//BOW/3/BCN/4/KAUZ	Sr-Sharp	30 MR	10MS	0	3C	
6085	SHARP/3/PRL/SARA//TSI/VEE#5/5/VEE/LIRA//BOW/3/BCN/4/KAUZ	Sr-Sharp	40 MR	10MS	0	••	
6105	PFAU/WEAVER*2//TRANSFER#12,P88.272.2	Sr24+TmP	90 M	5MS	0	••	ı
6108	PYN/BAU/3/MON/IMU//ALD/PVN/4/VEE#5/SARA//DUCULA	Sr24+Sr-Unknown	20 RMR	5MS	0	0	ı
6109	YANG87-142//SHA4/CHIL/3/TNMU	SrSha7	40 RMR	10RMR	0	ŝ	ı
6010	WBLL1*2/KIRITATI	MR	60 MSS	5MS	0	0	3C
6011	WBLL1*2/KIRITATI	MR	50 M	5MS	0	3;	3.C
6012	PFAU/WEAVER*2//CHAPIO	MR	30 M	10MS	0	;;	3-C
6014	KIRITATI//PBW65/2*SERI.1B	R-MR	40 M	TMS	0	б	4
	PFAU/MILAN/5/CHEN/AEGILOPS SQUARROSA (TAUS)// BCN/3/						
6019	VEE#7/BOW /4/PASTOR	R-MR	50 MSS	5MS	0	;0	3C
6021		Sr-Unknown	20 RMR	5MSS	0	••	13 =
6030		MR	30 M	5MS	0	;1	0
6031	KIRITATI/4/2*SERI.1B*2/3/KAUZ*2/BOW//KAUZ	MR	30 M	5MS	0	; 0	••
6033	KIRITATI//2*SERI/RAYON	MR	30 MSS	5MS	0	С	4
6050	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*WBLL1	MR	60 MSS	5MS	0	-	
6051	WHEAR/KUKUNA/3/C80.1/3*BATAVIA//2*WBLL1	MR	60 MSS	10MS	0	-	
6061	INQALAB 91*2/KUKUNA//KIRITATI	R-MR	30 MR	10MS	0	•••	
6110	BETTY/3/CHEN/AE.SQ//2*OPATA	MR	80 MSS	10MSS	0	0	
				000	0		-

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India and Pakistan (Luig, 1983) and for Sr35 was found in collection from Ethiopia, Kenya, Nepal (Huerta-Espino, 1992; Admassu & Fekadu, 2005). Virulence for Sr36 is present in collections from Ethiopia since 1987 (Huerta Espino, 1992, Admmassu & Fekadu 2005) and among new variants of Ug-99 (Jin et al., 2009). Field susceptibility of genes Sr36 and 37 at Thatta, Matli and seedling virulence for Sr36 in all isolates analyzed discourage their use for our breeding program. Similarly field virulence for Sr35 at Matli, Sr28, Sr29 & Sr13 at Thatta, renders them useless if used alone to protect wheat against local and Ug-99 races. These genes can however be incorporated in commercial cultivars, resistant to local races, to protect them against both Ug-99 and local races. Virulence for Sr29 was detected in isolates from Western Asia (Huertta-Espino, 1992) and three races from Ethiopia (Admassu and Fekadu, 2005). Confirmation of the presence of high virulence at seedling stage for genes Sr28, 29 & 13 and studies on their effectiveness when used in combination with other genes need to be conducted prior to their use in our lines. Virulence for Sr33 which conferred moderate level of resistance to Ug-99 in past (Jin et al., 2007) is reported in seven races from Ethiopia (Admassu & Fekadu, 2005). This gave MS-MSS reactions under field conditions in Thatta and Matli this year. Sr39 resistant to Ug-99 have effective field resistance at Thatta and Matli. This gene can be deployed in Pakistani wheat cultivars to enhance their resistance. Sr40, 44, dp-2, Gt and Tmp appear to have virulence in Thatta. SrTmp virulent isolates were found in samples analyzed under glasshouse conditions. The gene is showing effective level of field resistance in combination with Sr24 in 3rd SRSN lines (Table 5).

Wide spread of local stem rust race/s up to Bahawalpur in southern Punjab and susceptibility of mega cultivars planted in Sindh emphasize replacement of these cultivars and adaptation of a breeding strategy which fulfils our requirement to meet upcoming challenge of Ug-99 and local race/s. Out of 17 commercial cultivars, identified as immune to local stem rust, Faisalabad85, Kohinoor83, LU-26, Pasban90, Shahen94, Shahkar95, Bahawalpur97, Rohtas90 and Abadgar, are highly susceptible to Ug-99 and resistant to local race/s. These lines should be included in breeding program to improve their resistance by incorporating genes resistant to Ug-99 in them. Genes susceptible both to local and Ug-99 races should be avoided in our breeding program. Stem rust resistant genes Sr13, 25, 36, 37 and 44 should very carefully be used in combination with genes resistant to local race/s. Lines resistant to local and Ug-99 races identified in 3^{rd} stem rust screening nursery should also be included in our breeding program and be focused for direct adaptation and release.

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