Specific Virulence of *Puccinia recondita* f. sp. *tritici* in the United States from 1978 Through 1983

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ABSTRACT

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Puccinia recondita f. sp. tritici was identified from wheat leaf collections made by cooperators throughout the United States and from cereal rust field surveys of the Great Plains and Gulf Coast from 1978 through 1983. Testing of 1,928 isolates for virulence to 12 single-gene resistant tester lines showed 33 virulence combinations, which were categorized into 12 Unified Numeration (UN) races. Ninety-five percent of the isolates were classified into six UN races. No virulence was found to Lr 16 or 19.

Additional key words: wheat leaf rust

Wheat leaf rust, caused by Puccinia recondita Rob. ex. Desm. f. sp. tritici, is currently an important disease worldwide (11). For example, in 1982 in the southeastern United States, state average leaf rust loss estimates ranged from 3 to 15% (D. L. Long, unpublished). To select effective resistance genes in breeding cultivars for resistance to this pathogen, continual monitoring of virulence and combinations of virulence in the pathogen population is needed. Data from such surveys also provide an insight into the epidemiological patterns by monitoring the changes and movement of virulence combinations. The objective of this study was to characterize the virulence of the P. recondita population in the United States by regions. This information provides a data base needed by wheat breeders, epidemiologists, and modelers. Results are presented in a form to provide historical continuity with the Unified Numeration (UN) designation proposal of Johnston (4) and Basile (1).

MATERIALS AND METHODS

Leaf rust urediniospore collections were

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made by cooperators throughout the United States and by personnel of the Cereal Rust Laboratory during annual field surveys. These surveys were conducted over a 24,000-km route covering the Gulf Coast and the Great Plains of the United States (Fig. 1, areas 1, 4, 5, and 6). The surveys followed a predetermined, generally circular route through selected areas where small-grain cereals are important. Stops were made at commercial fields each 32 km or at the first field thereafter. Additional stops were made at nurseries and trap plots along the route. At least one rust collection was made in each locality visited during 1978-1981 and from each rusted field or nursery observed during 1982-1983. A collection consisted of a varying number of leaves bearing urediniospores from a single plant or cultivar.

When collections were received at the Cereal Rust Laboratory, urediniospores from them were used to inoculate 7-dayold seedlings of wheat (Triticum aestivum L. 'Thatcher,' CI 10003) treated with maleic hydrazide to enhance spore production. After 12-15 days, up to three leaves, each bearing a single uredinium or pruned to a single uredinium, were saved for inoculum. Six to 8 days later, sufficient urediniospores were collected from a single uredinium per collection in 1978 and 1979 and separately from two uredinia per collection in 1980 through 1983 to provide isolates to inoculate a differential host series. Spores suspended in a lightweight mineral oil were sprayed onto plants, which were placed in a dew chamber overnight at 18 C. Plants were

then placed in a greenhouse at 18-28 C. Observations were recorded 10-14 days later on a dichotomous high or low virulence basis, following the susceptible and resistant host classes described by Levine et al (5). Wheat single-gene isolines known to possess resistance genes Lr1, 2a, 2c, 3, 3ka, 9, 10, 16, 17, 18, and 19 were included in the host series to evaluate these isolates (9). Lr24 was included in



Fig. 1. Agroecological areas for *Puccinia recondita* f. sp. *tritici* in the United States: Area 1, mainly southern-adapted soft red winter wheats; 2 and 3, both planted to northern-adapted soft red winter wheat and white winter wheats, but appear to be epidemiologically separated by geographic features; 4, a mixture of wheat types but largely hard red winter; 5, hard red winter wheat; 6, mixed wheat types but largely hard red spring; 7, spring wheats planted in late fall; and 8, mixed wheats but largely soft white winter types.

Table 1. UN race designation of *Lr* virulence/avirulence character of *Puccinia recondita* f. sp. *tritici* isolates

UN race	Virulence (V) or avirulence (A) to <i>Lr</i> gene								
number	1	2a	2c	3					
1	Α	Α	A	Α					
2	Α	Α	Α	V					
10	Α	Α	V	Α					
3	Α	Α	V	V					
12	Α	V	V	Α					
17	Α	V	V	V					
11	V	Α	Α	Α					
5	V	Α	Α	V					
14	V	Α	V	Α					
6	V	Α	V	V					
9	V	V	V	Α					
13	V	V	V	V					

the cultivar Agent (3).

Isolates were grouped by eight agroecological geographic source areas (Fig. 1) on the basis of the locations of collections: area 1, mainly southernadapted soft red winter wheats; areas 2 and 3, both planted to northern-adapted

soft red winter and white winter wheats but appear to be epidemiologically separated by geographic features; area 4, a mixture of wheat types but largely hard red winter; area 5, hard red winter wheat; area 6, a mixture of wheat types but largely hard red spring wheat; area 7,

spring wheats planted in late fall; and area 8, a mixture of wheats but largely soft white winter types.

RESULTS AND DISCUSSION

The data are grouped by modified UN numbers based on the reactions of Lrl, 2a,

Table 2. Virulence combinations of isolates of Puccinia recondita f. sp. tritici from collections made in the United States from 1978 through 1983 as determined by the

							Num	ber of is	olates p	er viru	ence form	ula gro	uped b	y UN ra	aces ^a						
								١	UN 5 (1,	3)		UN 1	3 (1,2a,	2c,3)	U	N 3 (2c,	3)		UN 6 (1	,2c,3)	
			UN 2 (3) 3ka)	10	UN 17 (2a,2c,3)		10	3ka	10	10 17		17	10 17	3ka	3ka	3ka 9	3ka	3ka	10	1
ar	0	10	9	24	24	10	10	17	17	24	18	0	18	18	9	18	10	9	10	17	1
8		2	3			,	1		<i>-</i>	2	Area 1 ^{b,c}										
9		2 1	3		1					2			1				1 11				
Ó	23	4	6		2	1	•••				4	1					34	4	4		
1	•••	19	19		2	6	1				4	10		•••	3		15				
2	16	28	9	•••	•••	5	2	4		2	15	7		4	59	6	43	•••			
3	27	18	2	•••	•••	4	18	•••	•••		1	10		7	64		•••	11	4		
											Area 2										
8	2		•••	•••		•••		•••	•••		•••	•••						•••			
9	1			•••				•••	•••		3	•••		•••	•••	•••		•••	•••	•••	
)		7 1				4 					7						8 7				
1 2	1	1					1			1	2				17					2	
3		4				•••	6	7	4		2			1	4			15	5		
											Area 3										
3	2	3																			
•			6									•••		•••							
)	•••	2	6			5					3	•••				•••	20	•••			
ı	•••	•••	1				•••	•••	•••	•••	•••	•••		•••	•••	•••	1	•••	•••	•••	
2	1	4			1	2 2	1	2				1			13			 1		1	
,						2	1											1			
3		3				1	1			4	Area 4	3									
,	1	1				10				5		2	4								
)		2				17				5	1						8				
ı		26	1		2	23				1		3		1			6				
2	7	47	6		4	54	23	2		8	13	•••		3	27	•••	•••	•••	•••		
3	5	13	3	•••	1	30	67			11	4	•••	•••	2	2	•••		•••			
											Area 5										
3	2	4	•••	•••	•••	•••	•••	•••	•••		***	•••	•••	•••	•••	•••		•••			
)	3	1	•••		•••		•••	•••		•••		•••		•••		•••	1	•••	•••		
) 		22	2			19 25				1	2						1			•••	
2	3	11				25 25	30	1		1 2	1	1		2	2		1 			· · ·	
3		2				3	22											•••			
											Area 6										
3	7	10												•••		•••					
)	•••	•••	2	•••	•••	22	•••	•••	•••	1		•••		•••		•••		•••			
)	•••	3		•••	•••	29	3	•••	•••	1	7	•••	•••	•••	•••	•••	5			•••	
	1	12			1	38	4		•••	4	6	•••		1			6	•••	•••		
2	1 	4 2		2	2	51 29	53 15	6		6 	3				27 	2				2	
3		1									Area 7				•••						
)	•••					•••	•••	•••	•••						•••				•••	•••	
)	•••	•••		•••		•••	•••	•••	•••				•••	•••		•••	•••		•••	•••	
1	•••	•••	•••			•••		••••	•••	•••				•••	•••	•••	•••			•••	
2		4				 7	3					2		 1							
,		4				,	3			•••				1	•••	•••		•••	•••		
;											Area 8										
)	•••																				
)	•••																				
	•••							•••							•••						
2	•••	•••		•••			2	•••	•••						•••	•••					
3	•••	4					1	•••	•••		1		•••	•••	•••	•••		•••	•••	•••	

^{*}Specific virulence of designated UN race to Lr genes is in parentheses. Additional virulence to other single-gene Lr lines is indicated in columns below.

Areas are based on host types and geographic isolation (Fig. 1). Total isolates per area per year are listed in Table 3.

2c, and 3, historical differential host materials. Basile (1) proposed 27 UN races based on five differential cultivars: Malakof, Webster, Loros, Mediterranean, and Democrat. Soliman et al (12) subsequently showed that in North America, the resistance provided by both Mediterranean and Democrat is con-

ditioned by a single gene, Lr3; thus for North American race identification purposes, one may be dropped. The differential cultivars of Basile (1) were replaced by Samborski (8) with Thatcher isolines of Lr1, 2a, 2c, and 3. In North America in recent years, Lr2c has been susceptible whenever Lr2a is susceptible

(3). These four differential lines thus distinguish only 12 races in North America, generally corresponding to 12 of Basile's UN races. To facilitate comparisons, the UN numbers of Basile (1) for which Mediterranean and Democrat provide a similar response are used in the present report. These are

Table 3. Virulence combinations of isolates of *Puccinia recondita* f. sp. tritici from collections made in the United States from 1978 through 1983 as determined by the reactions of 12 wheat lines containing single genes for resistance (infrequently found UN races)

	Number of isolates per virulence formula grouped by UN races ^a												
					UN 11	10 17	UN 9 ((1,2a,2c)	UN	10 (2c)	UN 1	T 1	
ear	UN 1	9	UN 12 (2a,2c)		17			10		10		10	Total per are
аг	0		17	17	18	18	17	17	18	17	10	18	per yea
78							Area 1 ^c						
79	1	•••					•••						10
90			•••		•••		•••						18 83
1	•••		•••										79
2	•••						•••				17		218
3	1	•••	•••		•••	•••	•••				16	•••	192
							Area 2						
8	•••		•••	•••	•••		•••			•••			2
9	•••		•••	•••	•••	•••		•••		•••	•••	•••	6
0			•••	•••	•••	•••	•••		•••	•••		•••	28
1	2		•••		•••			•••		•••		•••	12
3				3	 17	2	2 		2		5		36 85
				-							J		03
8	2				•••		Area 3	•••					7
9			•••								•••		6
0					•••			•••			•••		36
1	•••	2		•••	•••		•••	•••					4
2	1		***			1	•••		•••				30
3	•••		***			•••	•••	•••	•••	•••	•••		9
							Area 4						
8	•••		***				•••	•••	•••				12
9	•••	•••	•••	•••		•••	•••		•••	•••	•••	•••	23
0	•••	•••	•••	•••		•••	•••	•••	•••	•••		•••	33
1	•••		•••	•••	•••		•••		•••				64
2								2		1			196 139
										•			139
8							Area 5			•••			4
9			•••								•••		6 5
0	•••		•••			3	•••						25
I			•••	•••	•••	•••	•••			•••			54
2	•••	•••	•••	•••	•••	•••	•••	•••	•••	•••	•••		77
3	•••			•••	•••	•••	•••	•••			•••		27
							Area 6						
8		•••		•••	•••			•••	•••	•••			17
9			•••		•••		•••	•••		•••	•••		25
0 1					•••	2		•••		•••	•••	•••	50
2			1								•••		74
3		•••											159 46
							A 7						7.0
8			•••				Area 7	•••					2
9			•••	•••	•••		•••						0
0	•••	•••		•••			•••		•••				0
1	•••		•••		•••	•••			•••				0
2		•••		•••	•••	•••	•••		•••				2
3	•••	•••		•••	***	•••	•••		•••	1		•••	16
0							Area 8						
8			•••	•••	•••		•••	•••	•••				0
9 0									•••				0
1											•••	•••	0
2	•••						•••						0
3					•••		•••					4	5 10

^aSpecific virulence of designated UN race to Lr genes is in parentheses. Additional virulence to other single-gene Lr lines is indicated in columns below.

^bFor Tables 2 and 3.

^cAreas are based on host types and geographic isolation (Fig. 1).

Table 4. Percentage of isolates of *Puccinia recondita* f. sp. tritici virulent to the single-gene differential lines used in the surveys from 1978 through 1983 by areas

/00F	Lr1	Lr 2a	Lr2c	Lr3	Lr3ka	Lr9	es (%) ^a <i>Lr</i> 10	Lr17	Lr 18	Lr2
'ear	LII	LF 48	L1 40	шэ		217	2710	2/1 1/	21.10	L'
978	30	10	20	100	Area 1 ^b 40	40	70	0	0	20
978 979		6	67	94	78	78	72	6	6	6
	6				58	53	59	5	5	2
980	16	2	53	100						3
981	19	20	43	100	47	47	59	5	5	9
982	24	7	65	92	54	51	56	11	12	9
983	40	12	69	96	45	42	42	4	9	0
otal	27	10	60	94	50	48	52	7	9	2
978	0	0	0	100	Area 2	0	0	0	0	0
979	83	ő	33	100	Ö	Õ	83	83	83	0
			50		29	29	100	32	32	0
980	32	14		100			100	33	33	0
981	33	0	75 75	100	58	58				3
982	36	5	75	78	47	47	39	31	19	
983	91	1	49	71	33	22	49	54	38	0
otal	64	4	56	80	36	30	60	44	34	1
					Area 3					
978	0	0	0	71	0	0	43	0	0	0
979	0	0	0	100	100	100	0	0	0	0
980	8	14	69	100	72	72	69	8	8	0
981	Ö	0	25	50	50	50	75	50	50	0
982	27	10	63	93	43	43	40	20	10	3
								56	56	0
983	78	22	89	100	11	11	89			
otal	20	11	58	93	52	54	53	15	12	1
070	47	22	22	100	Area 4	Λ	75	0	0	33
978	67	33	33	100	0	0				
979	48	70	70	100	0	0	70	17	17	22
980	18	52	76	100	24	24	100	3	3	15
981	9	42	53	100	11	11	94	3	3	5
982	26	30	44	99	17	17	80	10	2	6
983	60	23	25	99	4	4	93	5	4	9
otal	36	33	43	99	11	11	86	7	3	ģ
Otai	30	33	43	"			00	,	3	ŕ
					Area 5					
1978	0	0	0	100	0	0	67	0	0	0
1979	0	0	20	100	20	20	40	0	0	0
980	20	76	80	88	4	4	100	20	20	C
1981	7	46	52	100	6	6	50	6	6	4
					3	3	92	4	3	3
982	47	36	39	100						
983	81	11	11	100	0	0	100	0	0	0
Total	35	39	42	98	4	4	80	6	5	2
					Area 6					
978	0	0	0	100	0	0	59	0	0	0
979	4	88	88	100	0	0	92	0	0	12
980	26	58	68	96	10	10	100	24	18	2
981	20	54	62	99	8	8	97	11	9	7
982	44	32	52	100	17	18	80	7	3	ϵ
983	33	63	63	100	0	0	100	0	0	C
Fotal	31	46	57	99	11	10	88	8	6	5
					A					
978	50	0	0	100	Area 7 0	0	100	50	50	(
1979	0	0	0	0	0	0	0	0	0	(
1980	0	0	ő	Ö	Õ	0	0	0	0	(
1981	0	0	0	0	0	0	0	ŏ	Õ	(
				100	0	0	0	ő	0	(
1982	100	100	100							
1983	25	50	56	94	0	0	100	12	6	(
[otal	35	50	55	95	0	0	90	15	10	(
					Area 8					
1978	0	0	0	0	0	0	0	0	0	(
1979	0	0	0	0	0	0	0	0	0	(
1980	ő	ő	0	0	0	0	0	0	0	(
1981	100	0	60	100	0	0	100	60	60	(
					0	0	100	10	10	ì
1982	60	0	40	60						
1983 Total	0 73	0	0 47	0 73	0	0	0 100	0 27	0 27	
ividi	13	U	4/	13	J	U	100	21		

^a No virulence was observed for *Lr*16 or 19.

designated in Table 1.

The virulence formulas for the 1,928 isolates, based on the 12 lines each possessing a single gene for resistance, are shown in Tables 2 and 3. The arrangement of the 12 UN race categories places populations together that have apparent developmental or geographic relationships.

Virulence to either Lr16 or 19 was not identified during this period of years in the U.S. survey. These results are similar to data of Samborski (10) for Canada and Statler et al (15) for North Dakota. Virulence to Lr16, resistance that is present in Exchange and Selkirk, was previously common in Canada (7) and North Dakota (14) when Selkirk and related cultivars were grown and has been found in the Pacific Northwest (6). Lr16 is not currently present in common cultivars in the United States (2,3), and the corresponding virulence characteristic has become rare in recent years (10,15). Rare reports of virulence to Lr19 have not been verified by subsequent retained cultures. This resistance probably has not been used commercially.

In contrast, virulences to Lr3 (95%) and to Lr10 (73%) were widespread in the U.S. population of P. recondita f. sp. tritici (Table 4). These two genes for resistance are carried by numerous widely grown cultivars (3,13).

Because of their patterns of occurrence. virulences to the other differential lines are considered in the context of the UN races. The six UN races with virulence to Lr3 (UN 2, 3, 5, 6, 13, 17), constituting 95% of the isolates identified, are the major populations of these surveys. UN 3 and 6 were more prevalent in the south and east (primarily areas 1 and 2 but also 3 and 4) (Table 2), although a large sample of UN 3 (27 isolates) was collected in area 6 in 1982. UN 3 and 6 are characterized by virulence to Lr2c and avirulence to Lr2a. Much of the virulence to Lr3ka and Lr9 occurs in these populations, but these virulences were also found to a lesser extent in a small portion of the UN 2 population, otherwise only virulent to Lr3 (largely in area 1), in which these two virulence characters again occur together (69 isolates, Table 2). Lr9 was incorporated into soft red winter wheat cultivars (3), which were widely grown in the South and East (2). Usage of Lr3ka is not known. UN 6, which includes virulence to Lr1 compared with UN 3, increased considerably in 1983 (Table 2) and was found mainly in areas 1 and 2.

UN 2, 17, and 5, the most numerous elements of the pathogen population, were most prevalent in the south and central regions (areas 1, 4, 5, and 6) (Table 2), in comparison to the more easterly orientation of UN 3 and 6. UN 17 and 5 appear similar to UN 2 with the addition of virulence to Lr2a and 2c in UN 17 and virulence to Lr1 in UN 5. They were proportionately more prevalent in

^bArea description in text and Figure 1.

^{&#}x27;Percentage of total isolates from the area for the 6-yr period.

^dPercentage of total isolates from the United States for the 6-yr period.

the north than UN 2. Lr2a is possessed by several commonly grown hard red spring wheat cultivars (3,13) in area 6 (2), and Lr1, by Newton (3), a recent extensively grown hard red winter cultivar in area 5. UN 17 has a unique virulence uniformity and was found mainly in areas 4, 5, and 6 (very little in area 1). Virulence to Lr24 occurred in very low frequency only in UN 2 and 5, and in this survey sample, never in combination with virulence to Lr2a, 2c, 3ka, or 9. It was most prevalent in area 4. Several cultivars possessing Lr24 (3) were grown in limited amounts in the Southeast (area 1) and the hard red winter wheat region (areas 4 and 5) (2).

UN 13 occurred primarily in the South (mostly in area 1) (Table 2). This contrasts with the UN 3 and 6 populations and the UN 2 and 5 populations, which were found not only in the South but also farther north in the eastern and central areas, respectively.

Virulences to Lr17 and 18 appeared to

occur randomly at low frequencies in most elements of the pathogen population as otherwise evaluated and described but were not found in widely distributed UN 2 and 17. Neither virulence was found in combination with virulence to *Lr*9 or 24. *Lr*17 and 18 have not been used in any widely grown North American commercial cultivars (2,3,13).

LITERATURE CITED

- Basile, R. 1957. A diagnostic key for the identification of physiologic races of *Puccinia rubigo*vera tritici grouped according to a unified numeration scheme. Plant Dis. Rep. 41:508-511.
- Briggle, L. W., Straus, S. L., Hamilton, D. E., and Howse, G. H. 1982. Distribution of the varieties and classes of wheat in the United States in 1979. U.S. Dep. Agric. Agric. Res. Serv. Stat. Bull. 676. 107 pp.
- Browder, L. E. 1980. A compendium of information about named genes for low reaction to Puccinia recondita in wheat. Crop Sci. 20:775-779.
- Johnston, C. O. 1956. Unified number for races of *Puccinia triticina*. Robigo 1:2-3.
- Levine, M. N., Ausemus, E. R., and Stakman, E. C. 1951. Wheat leaf rust studies at Saint Paul, Minnesota. Plant Dis. Rep. Suppl. 199. 17 pp.

- Milus, E. A., and Line, R. F. 1980. Virulence of Puccinia recondita in the Pacific Northwest. Plant Dis. 64:78-80.
- Samborski, D. J. 1967. Leaf rust of wheat in Canada in 1966. Can. Plant Dis. Surv. 47:3-4.
- Samborski, D. J. 1968. Leaf rust of wheat in Canada in 1967. Can. Plant Dis. Surv. 48:6-8.
- Samborski, D. J. 1976. Leaf rust of wheat in Canada in 1976. Can. Plant Dis. Surv. 56:123-125.
- Samborski, D. J. 1983. Occurrence and virulence of *Puccinia recondita* in Canada in 1982. Can. J. Plant Pathol. 5:194-196.
- Samborski, D. J. 1985. Leaf rust of wheat. In: The Cereal Rusts. Vol. 2. Diseases, Distribution, Epidemiology and Control. A. P. Roelfs and W. R. Bushnell, eds. Academic Press, New York. In press.
- Soliman, A. S., Heyne, E. G., and Johnston, C. O. 1964. Genetic analysis of leaf rust resistance in the eight differential varieties of wheat. Crop Sci. 4:246-248.
- Statler, G. D. 1984. Probable genes for leaf rust resistance in several Hard Red Spring wheats. Crop Sci. 24:883-886.
- Statler, G. D., and MacArthur, R. A. 1975. Wheat leaf rust in North Dakota in 1973 and 1974. Plant Dis. Rep. 59:323-326.
- Statler, G. D., Miller, J. D., and Leben, S. 1982. Wheat leaf rust in North Dakota during 1979-1981. Plant Dis. 66:1174-1176.