## Letter Code System of Nomenclature for Puccinia graminis f. sp. avenae

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### **ABSTRACT**

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Current systems that describe the virulence phenotype in Puccinia graminis f. sp. avenae lack a systematic approach for the naming of races or to provide easily made comparisons of virulence among races. A new nomenclature system that simply and systematically characterizes virulence in P. graminis f. sp. avenae is described. The new system has the distinct advantage of providing easily seen relationships among races in contrast to previous nomenclature systems. This allows for easier interpretation of virulence relationships in the oat stem rust population and provides a large amount of virulence information with a minimum of written characters. This system uses single-gene differential lines with the resistance genes Pg1, Pg2, Pg3, Pg4, Pg6, Pg8, Pg9, Pg10, Pg12, Pg13, Pg15, and Pg16, grouped into three subsets of four lines in sequential Pg gene order. By grouping in sequential gene number order, the relationship of the new system to the "standard" system is easily seen. Each race is designated by a three-letter code, based on the seedling reaction (low or high) on 12 differential lines. The letter code nomenclature system is open ended and can be updated easily as new differential genes are identified. This system simply and precisely describes the virulence phenotypes of isolates of P. graminis f. sp. avenae, and allows for easily made comparisons of virulence of isolates collected over time and across geographical locations worldwide.

Several nomenclature systems to describe virulence in Puccinia graminis f. sp. avenae Erikss. & Henning have been developed; however, currently, there is no adequate system that simply describes the virulence phenotypes of isolates of P. graminis f. sp. avenae or provides easily made comparisons of virulence of isolates from different geographical areas. Physiologic specialization in P. graminis f. sp. avenae first was described by Stakman et al. in 1923 using the varieties Victory, White Tartar, and Monarch to describe four races of oat stem rust (20). In 1925, Bailey described five races of P. graminis f. sp. avenae, using White Tartar (=White Russian), Richland, and Jostrain (3), Levine and Smith added races 6 to 10 in 1937, and Newton and Johnson added races 11 to 13 in 1944, using the same three differential

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varieties (11,16). These first 13 races were used as the standard for describing physiologic specialization in *P. graminis* f. sp. *avenae*.

Subsequent to the description of the 13 "standard" races, additional genes conferring resistance to P. graminis f. sp. avenae were found and are described in Table 1. Genes originally were designated by using a letter (e.g., gene A) but currently are designated by number (e.g., Pg2) using a standardized gene nomenclature system (19). Deviations from standard races were described as subraces, adding letters to standard race numbers to identify additional races. In 1965, Green (7) devised a virulence formula nomenclature system for Canada (C-races) using genes from the differential varieties Richland, Rodney, Minrus, Jostrain, CI 4023, and CI 5844-1. In 1970, Stewart and Roberts published the first proposed international system for identifying races of P. graminis f. sp. avenae, using the same six differential varieties as Green (7) and the diploid oat line Saia to describe the "standard" races, numerous subraces, C-races, and exotic races from several countries, with a total of 97 races identified (21).

The development of backcross-derived lines (Rodney 0 background) with a single gene (*Pg* gene) for oat stem rust resistance led to the development of the North

American (NA) nomenclature system in 1979 (14), which currently is used to characterize virulence in P. graminis f. sp. avenae in Canada and the United States. În contrast, virulence in P. graminis f. sp. avenae in Australia currently is characterized using a modified Stewart and Roberts system (2). Because there is a lack of continuity across the various nomenclature systems and there appears to be no systematic approach in the naming of races (other than by order of discovery), we describe a new nomenclature similar to existing cereal rust nomenclature systems (4,12,17) that simply and precisely identifies the virulence phenotype (race) of isolates of P. graminis f. sp. avenae. This system will provide a method to easily and accurately compare virulence of isolates collected over time and across geographical areas.

# THE LETTER CODE RACE NOMENCLATURE SYSTEM

Currently, there are 17 numbered Pg genes identified, plus the Pg-a complex. Genes Pg5 and Pg7 are repetitive of genes Pg4 and Pg6, respectively, and are of no use as differential lines (13). Genes Pg11 and Pg17 are expressed only at the adult plant stage and would not be useful as differential lines at the seedling stage (10,15). Gene Pg14 is ineffective to numerous North American and exotic isolates tested thus far (T. Fetch, unpublished data) and is not noted for its resistance (13); thus, it is not included in the differential set. All other numbered genes are useful in characterizing virulence and are included in the differential sets.

The differential lines for the new nomenclature system are listed in Table 2. Most (Pg1, Pg2, Pg3, Pg8, Pg9, Pg13, Pg15, and Pg16) are in a Rodney 0 (CI 9317, no known Pg resistance gene) background as described by Martens et al. (14). Comparisons of infection type (IT) between original sources of resistance to their Rodney 0-derived lines were made with several races using previously reported methods (5), and the results are presented in Table 3. IT of infected seedlings was determined using a 0-to-4 scale, where ITs of 0, ;, 1, and 2, or combinations thereof, are classified as low ITs and indicative of resistance, whereas ITs of 3 or

4 are high ITs and indicative of susceptibility (3). Because expression of several Pg genes is affected by incubation temperature (6,14), all comparisons were conducted in a growth chamber at 19°C with a 16-h photoperiod and light intensity of 180 to 220 µmol photon m<sup>2</sup> s<sup>-1</sup>. Under these incubation conditions, all genes will express their low ITs to avirulent races of P. graminis f. sp. avenae, which is easily discernable from a susceptible (IT 34 or 4) reaction. For a given Pg gene, the observed ITs between the original source and the corresponding Rodney line were comparable (Table 3). However, it was found that several Pg gene sources and their Rodney 0 derivatives acquired from the Unites Department of Agriculture-Agricultural Research Service (USDA-ARS) National Small Grains Collection (NSGC) in Aberdeen, ID were mixed for ITs. Therefore, differential lines in the Rodney 0 background were reselected for the correct phenotype and increased for seed that is available for race identification of P. graminis f. sp. avenae.

The letter code nomenclature utilizes 12 single-gene Pg differential lines, using three subsets of four lines organized into a hexadecimal system that has 16 possible combinations of low (L) or high (H) reaction for each letter (Table 4). This is adapted from the existing nomenclature systems used for other cereal rusts (4,12,17). The advantage of the letter code system is a simple, short, and accurate description of virulence phenotype across most of the described resistance genes, which enables utilization of this system regardless of when and where isolates have been collected. In contrast, comparisons of virulence of isolates identified using the "standard", C number, and NA formula system were difficult and imprecise (8), due to the lack of a systematic approach in numbering of races and to addition of differential lines that occurred over time. Because the order of genes in the new system is sequential, it is relatively easy to make comparisons to the "standard" system originally described (e.g., race 1 = B\_ or  $C_{-}$  races, race  $2 = D_{-}$  races, race 6group = TG\_ or TD\_ races, race 7 group =  $N_{\rm or} P_{\rm races}$ , race  $8 = J_{\rm races}$ , and race 11 group = G\_ or H\_ races). Additionally, virulence of isolates collected worldwide cannot currently be compared, because differential sets are not uniform and different nomenclature systems are utilized. The letter code system is flexible in that, as new Pg genes are characterized, they can be added as additional subsets and letters appended accordingly. Data on supplemental lines can be added to the letter code using a plus (+) sign (e.g., race TGL+ Pga). In addition, the letter code system is suitable for international use because it incorporates almost all Pg genes to characterize virulence in P. graminis f. sp. avenae. Virulence information previously reported in North America is presented in Table 5 using the new letter code, NA number designation (14), and avirulence or virulence formula systems. Each NA race reported previously corresponded to a unique letter-code race designation following this system.

The new system uses three seedling Pg genes (Pg6, Pg10, and Pg12) that have not been used previously to characterize P. graminis f. sp. avenae. Gene Pg6 (from CI 6956) and the resistance gene in the diploid line Saia (which is used in the Stewart and Roberts nomenclature) appear to be identical (18). Although Pg6 has not yet been transferred into a hexaploid background, it has been shown to be useful in differentiating isolates of stem rust at the Winnipeg Rust laboratory. In testing of 23 common North American races stored at

Table 2. Differential lines used for characterizing virulence in Puccinia graminis f. sp. avenaea

Pg gene	CI no.	CN no.	RL no.	IT at 20°Cb	Source of resistance
Pg1	9318	58243	899	11+	White Russian, CI 551
Pg2	9319	58244	815	;1	Hajira, CI 1001
Pg3	2660	58245		;1 to X-	Jostrain, CI 2660
Pg4	6661	56534	2123	;1-	Hajira, CI 1001
Pg6	6956	56818		0;1-	CD 3820, CI 6956
Pg8	9321	58246	903	12-	Hajira, CI 8111
Pg9	9322	58247	879	11+	CI 6792
Pg10	8457	58055		23n	Illinois Hulless, CI 2824
Pg12	8250	64102		;1	Kyto, CI 8250
Pg13	9212	28959	618	11+	Avena sterilis, PI 324798
Pg15	9351	58276	997	11+	A. sterilis, CAV 1830
Pg16	9352	58277	882	12-	A. barbata, D203
Pg-a		1947	996	0;1-	Omega, CI 9139

<sup>&</sup>lt;sup>a</sup> CI is the Cereal Introduction number (United States Department of Agriculture–Agricultural Research Service, Aberdeen, ID), CN is the Canadian Number (Plant Genetic Resources Collection, Saskatoon, SK, Canada), and RL is the Rust Lab accession number (Cereal Rust Lab, Winnipeg, MB, Canada)

Table 1. Background information for resistance genes to Puccinia graminis f. sp. avenae

Pg gene	Lettera	Year	Origin	Source	Avena sp.	CI number	Temperature <sup>b</sup>
1	D	1925	Russia	White Russian	sativa	551	None
1	D	1938	Russia	Minrus	sativa	2144	None
2	A	1925	Russia	Richland	sativa	787	None
3	E	1925	USA	Sevnothree	sativa	3251	21°C
3	E	1925	France	Jostrain	sativa	2660	21°C
1	В	1954	South Africa	Hajira	sativa	1001	26°C
5	C	1954	South Africa	Hajira	sativa	4019	
5		1956	Uruguay	CD 3820	strigosa	6956	None
7		1956	Uruguay	CD 3820	strigosa	6956	
3	F	1959	North Africa	Hajira	sativa	8111	27°C
)	Н	1965	Argentina	Sante Fe selection	sativa	5844-1	25°C
10	G	1965	USA	Illinois Hulless	nuda	2824	None
11		1968	Rhodesia	Burt	sativa	3034	Adult <sup>c</sup>
12		1968	Yugoslavia	Kyto	sativa	8250	25°C
13	M	1970	Tunisia	CW490-2	sterilis	PI 324798	27°C
14	N	1972	Wales	Milford	sativa	5039	
5		1980	Turkey	CAV 1830	sterilis	9351	26°C
6	R	1979	Israel	D-203	barbata	9125	25°C
7		1990	Spain	IB 3056	sterilis		Adult
Pg-a	X	1981	USA	Omega	hybrid	9139	26°C

<sup>&</sup>lt;sup>a</sup> Genes originally were designated by letter, then changed later to a Pg gene number (17).

<sup>&</sup>lt;sup>b</sup> Infection types (ITs) are based on a 0-to-4 scale (3).

b Incubation of lines at or above this temperature renders ineffective the expression of resistance (6). For genes Pg1, Pg2, Pg6, and Pg10, there is no effect of incubation temperature up to 30°C.

<sup>&</sup>lt;sup>c</sup> CI 3034 also has been reported to possess the seedling gene *Pg1* (11).

the Winnipeg Rust Laboratory, only NA1 and NA70 have been found to be virulent on Pg6 (T. Fetch and Zegeye, *unpublished data*); thus, most North American races were presumed to be avirulent to Pg6 in Table 5. In contrast, virulence to Saia (= Pg6) is observed frequently in isolates of P. graminis f. sp. avenae in Australia (2).

Gene Pg10 (from CI 2824) has not been used in previous nomenclature systems due to temperature sensitivity (14). Studies at Winnipeg indicate that the reaction of Pg10 is intermediate (IT = 23N) at low (15 to 21°C) temperatures, but becomes more resistant (12+N) at elevated (>24°C) temperatures (6). Although Pg10 exhibits an intermediate resistance reaction (IT = 23n), the distinct necrosis associated with its resistance response is unique to this gene (9) and is easily discernable from a susceptible (IT = 3 or 4) reaction. Virulence to Pg10 has not been observed in North America among several thousand isolates tested at Winnipeg and St. Paul rust laboratories, but virulence to Pg10 has been found from an isolate obtained from New Zealand (T. Fetch, unpublished data).

Gene Pg12 has not been used previously in nomenclature systems, but the related Pg-a complex was used by Martens et al. (14) in the NA nomenclature. Gene Pg12 is included in the new differential set in lieu of the Pg-a complex because (i) it is a single gene, (ii) no differential response (high versus low) between Pg12 and Pg-a has been found among several thousand isolates tested simultaneously on the two lines, and (iii) allelism tests of  $F_2$  progeny from crosses between the Pg12 source

'Kyto' (CI 8250) and Pg-a source 'Omega' (CI 9139) found no susceptible segregants among 4495  $F_2$  plants (T. Fetch, unpublished data). Thus, the low seedling IT observed on the Pg-a complex apparently is conditioned by the gene Pg12 and may be modified by an additional unknown recessive gene. This is in contrast to previously published data that reported that Pg12 is not involved in the Pg-a response (1). The Pg-a complex was included in Table 2 as a supplementary line because of historical use, but is not intended as a differential line in this nomenclature system.

The letter code nomenclature system presented here is simple, precise, and describes the virulence spectrum of races of *P. graminis* f. sp. *avenae* on most known seedling resistance genes in oat. This system will be useful in enabling rust workers to uniformly characterize *P. graminis* f. sp. *avenae* isolates for discussion on virulence dynamics and population studies. Limited seed for all differential lines is available from the authors, and also from seed depositories in Plant Gene Resources Canada (PGRC, Saskatoon, SK) and USDA-ARS NSGC in Aberdeen, ID.

**Table 4.** Letter code designations for races of *Puccinia graminis* f. sp. *avenae* using 12 differential single  $P_g$ -gene lines in three ordered subsets of four lines each

	Subset	Classification of infection types (ITs) <sup>a</sup>						
	1	Pg1	Pg2	Pg3	Pg4			
	2	Pg6	Pg8	Pg9	Pg10			
Letter code	3	Pg12	Pg13	Pg15	Pg16			
В		L	L	L	L			
C		L	L	L	Н			
D		L	L	Н	L			
F		L	L	Н	Н			
G		L	Н	L	L			
Н		L	Н	L	Н			
J		L	Н	Н	L			
K		L	Н	Н	Н			
L		Н	L	L	L			
M		Н	L	L	Н			
N		Н	L	Н	L			
P		Н	L	Н	Н			
Q		Н	H	L	L			
Ř		Н	H	L	Н			
S		Н	Н	Н	L			
T		H	H	Н	H			

<sup>&</sup>lt;sup>a</sup> Classification of infection types: L = low/resistant (ITs of 0, ;, 1, and 2, or combination thereof) and H = high/susceptible (ITs of 3, 4, or combination thereof) (17).

**Table 3.** Infection type (IT) comparison of oat lines with original sources of Pg gene resistance and equivalent Rodney 0-derived lines to races of Puccinia graminis f. sp. avenae

		 Genetic background	IT at 20°Ca					
Pg gene	CI no.b		NA1	NA8	NA16	NA20	NA55	
Pg1	551	White Russian	2 <sup>±</sup>	12-	3+4	3+4	34n	
	9318	Rodney 0	12-	$1^{\pm}$	3+4	4	3+n	
Pg2	787	Richland	;1	;1-	;1	4	34n	
	9319	Rodney 0	;1	;1	;1	4	34n	
Pg3	2660	Jostrain	$X^{-}$	X=	4	0;	X=	
	9320	Rodney 0	$X^{-}$	X-	4	0;	X=	
Pg4	1001	Hajira	;1	33 <sup>+</sup>	;1	4	34n	
	6661	Rodney	;1-	33 <sup>+</sup>	;1	4	34n	
Pg8	8111	Hajira	;1	;1+	34	11+	;1	
	9321	Rodney 0	1-	$1^{\pm}$	3+4	12-	;1	
<sup>2</sup> g9	5844-1	Sante Fe select	11+	4	;1	4	34n	
	9322	Rodney 0	12	4	;1+	4	3+4	
Pg10	2824	Illinois Hulless	$3^{\pm}N$	23+N	24N	23+N	13-N	
	8457	X-1588	13N	12+N	23N	12+N	22+n	
Pg13	2647	A. sterilis	;1	34	;1	;1	;1	
~	9212	Rodney 0	;1-	34	;1-	;1	;1	
Pg15	1830	A. sterilis	34	34	;1	34	34	
~	9351	Rodney 0	34	34	;1	34	23+	
Pg16	D203	A. barbata	1-	;1-	1-	;1-	34	
~	9352	Rodney 0	11-	;1	11-	;1	34	

<sup>&</sup>lt;sup>a</sup> ITs are based on a 0-to-4 scale where ITs of 0, ;, 1, and 2 are indicative of a resistant (low) response and ITs of 3 or 4 of a susceptible (high) response (3). Symbols + and – indicate slightly larger and smaller pustule sizes, respectively. North American (NA) races use the nomenclature of Martens et al. (14).

<sup>&</sup>lt;sup>b</sup> CI is the Cereal Introduction number (United States Department of Agriculture–Agricultural Research Service, Aberdeen, ID).

Table 5. Key to races of *Puccinia graminis* f. sp. avenae using the letter code nomenclature, North American (NA) nomenclature, and avirulence/virulence formula system<sup>a</sup>

Code	Race	Eff/ineff Pg genes	Code	Race	Eff/ineff Pg genes	Code	Race	Eff/ineff Pg genes
BDB	NA38	1,2,3,4,8,13,15,16,a/9	KBD	NA11	1,8,9,13,16,a/2,3,4,15	RBD	NA19	3,8,9,13,16,a/1,2,4,15
BDD	NA2	1,2,3,4,8,13,16,a/9,15	KDD	NA12	1,8,13,16,a/2,3,4,9,15	RDD	NA20	3,8,13,16,a/1,2,4,9,15
BDJ	NA3	1,2,3,4,8,16,a/9,13,15	KDJ	NA32	1,8,16,a/2,3,4,9,13,15	RDF	NA58	3,8,13,a/1,2,4,9,15,16
BLD	NA1	1,2,3,4,8,9,13,16,a/15	KJD	NA13	1,13,16,a/2,3,4,8,9,15	RDJ	NA54	3,8,16,a/1,2,4,9,13,15
CDJ	NA4	1,2,3,8,16,a/4,9,13,15	LDD	NA65	2,3,4,8,13,16,a/1,9,15	RGB	NA21	3,9,13,15,16,a/1,2,4,8
CLD	NA70	1,2,3,8,9,13,16,a/4,15	LGB	NA14	2,3,4,9,13,15,16,a/1,8	RGD	NA77	3,9,13,16,a/1,2,4,8,15
DBB	NA39	1,2,4,8,9,13,15,16,a/3	MDD	NA71	2,3,8,13,16,a/1,4,9,15	RJB	NA51	3,13,15,16,a/1,2,4,8,9
DBD	NA5	1,2,4,8,9,13,16,a/3,15	MGB	NA44	2,3,9,13,15,16,a/1,4,8	RJD	NA72	3,13,16,a/1,2,4,8,9,15
DDD	NA6	1,2,4,8,13,16,a/3,9,15	NBB	NA15	2,4,8,9,13,15,16,a/1,3	RJJ	NA75	3,16,a/1,2,4,8,9,13,15
DDJ	NA7	1,2,4,8,16,a/3,9,13,15	NDB	NA45	2,4,8,13,15,16,a/1,3,9	SBD	NA22	4,8,9,13,16,a/1,2,3,15
DJD	NA53	1,2,4,13,16,a/3,8,9,15	NDD	NA35	2,4,8,13,16,a/1,3,9,15	SDD	NA61	4,8,13,16,a/1,2,3,9,15
FBD	NA56	1,2,8,9,13,16,a/3,4,15	NDJ	NA47	2,4,8,16,a/1,3,9,13,15	SGB	NA23	4,9,13,15,16,a/1,2,3,8
FDD	NA74	1,2,8,13,16,a/3,4,9,15	NDL	NA46	2,4,8,13,15,16/1,3,9,a	SGD	NA63	4,9,13,16,a/1,2,3,8,15
FDJ	NA8	1,2,8,16,a/3,4,9,13,15	NGB	NA16	2,4,9,13,15,16,a/1,3,8	TBD	NA24	8,9,13,16,a/1,2,3,4,15
GBD	NA59	1,3,4,8,9,13,16,a/2,15	NGD	NA18	2,4,9,13,16,a/1,3,8,15	TDB	NA52	8,13,15,16,a/1,2,3,4,9
GDD	NA64	1,3,4,8,13,16,a/2,9,15	NGL	NA17	2,4,9,13,15,16/1,3,8,a	TDD	NA25	8,13,16,a/1,2,3,4,9,15
GDJ	NA34	1,3,4,8,16,a/2,9,13,15	NJB	NA48	2,4,13,15,16,a/1,3,8,9	TDF	NA55	8,13,a/1,2,3,4,9,15,16
HBD	NA69	1,3,8,9,13,16,a/2,4,15	PBD	NA57	2,8,9,13,16,a/1,3,4,15	TDJ	NA26	8,16,a/1,2,3,4,9,13,15
HDB	NA40	1,3,8,13,15,16,a/2,4,9	PBG	NA66	2,8,9,15,16,a/1,3,4,13	TGB	NA27	9,13,15,16,a/1,2,3,4,8
HDD	NA9	1,3,8,13,16,a/2,4,9,15	PDB	NA49	2,8,13,15,16,a/1,3,4,9	TGD	NA29	9,13,16,a/1,2,3,4,8,15
HDJ	NA31	1,3,8,16,a/2,4,9,13,15	PDD	NA36	2,8,13,16,a/1,3,4,9,15	TGL	NA28	9,13,15,16/1,2,3,4,8,a
JBD	NA10	1,4,8,9,13,16,a/2,3,15	PDJ	NA37	2,8,16,a/1,3,4,9,13,15	TJB	NA68	13,15,16,a/1,2,3,4,8,9
JDB	NA41	1,4,8,13,15,16,a/2,3,9	QBB	NA73	3,4,8,9,13,15,16,a/1,2	TJD	NA30	13,16,a/1,2,3,4,8,9,15
JDD	NA33	1,4,8,13,16,a/2,3,9,15	QBD	NA60	3,4,8,9,13,16,a/1,2,15	TJG	NA76	15,16,a/1,2,3,4,8,9,13
JDJ	NA42	1,4,8,16,a/2,3,9,13,15	QDD	NA62	3,4,8,13,16,a/1,2,9,15	TJJ	NA67	16,a/1,2,3,4,8,9,13,15
KBB	NA43	1,8,9,13,15,16,a/2,3,4	QGB	NA50	3,4,9,13,15,16,a/1,2,8			

<sup>&</sup>lt;sup>a</sup> Code = letter code, Race = NA race, and Eff/ineff = effective/ineffective. The letter code system infers that reaction to Pg12 is identical to Pg-a, all NA races are avirulent to Pg10, and only races NA1 and NA70 are virulent to Pg6.

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