

Inheritance of Phomopsis Stem Canker Resistance in Sunflower

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INTRODUCTION

Phomopsis stem canker (PSC) caused by *Phomopsis helianthi* Munt.-Cvet. (telemorph *Diporthe helianthi*)



First observed in former Yugoslavia in late 1970's

Yield loss as high as 50% in Europe

PSC is becoming wider in global distribution

Increasingly become damaging to the U.S. sunflower since 2010

Resistance to PSC is polygenic with additive gene action

PSC tolerance have been reported in USDA sunflower germplasm collection

GOAL OF THE PROJECT

Overall objective

- To improve Phomopsis resistance in cultivated sunflower

Specific objectives

- Investigate the inheritance of PSC resistance in sunflower
- Identify genes/QTL associated with PSC resistance
- Identify molecular markers associated with PSC resistance genes/QTL
- Design PCR based primers for use in marker-assisted PSC resistance breeding

MATERIALS & METHODS

Plant Materials

- **Parents**

HA-R3 is highly tolerant to PSC

HA 89 is susceptible to PSC

- **Mapping population**

- 164 F₆-derived RILs developed through Single Seed Descent method from the cross of HA 89/HA-R3

MATERIALS & METHODS

PSC evaluation

- **Environments**
 - 2016: Grandin, ND; Rothsay and Crookston, MN
 - 2017: Glyndon, Rothsay and Crookston, MN
- **Field design**
 - Randomized complete block with 3 replications
- **Field inoculation**
 - No artificial inoculum applied
- **Disease incidence (DI) scoring**
 - Percent of plants showing phomopsis symptom

RESULTS

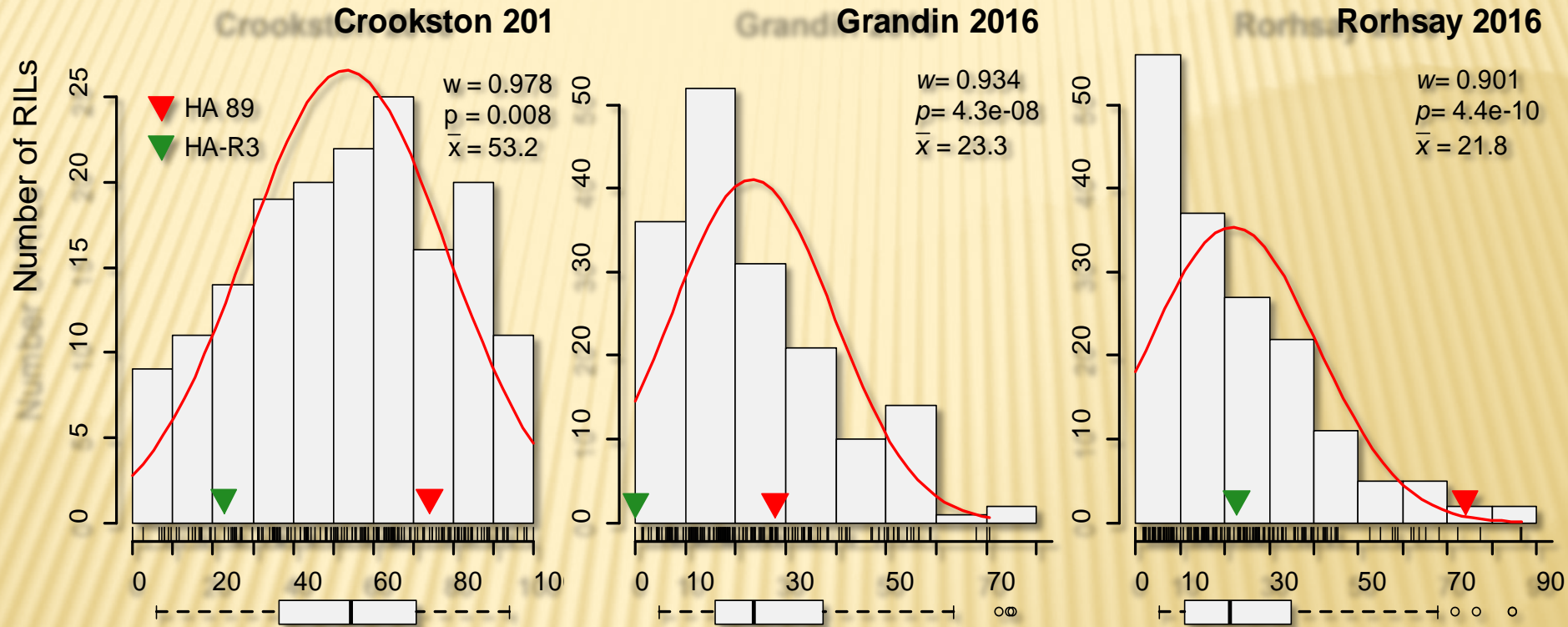


Fig 1. Phenotypic frequency distributions of PSC disease incidence (DI) for HA 89/HA-R3 RIL population evaluated in 2016

RESULTS

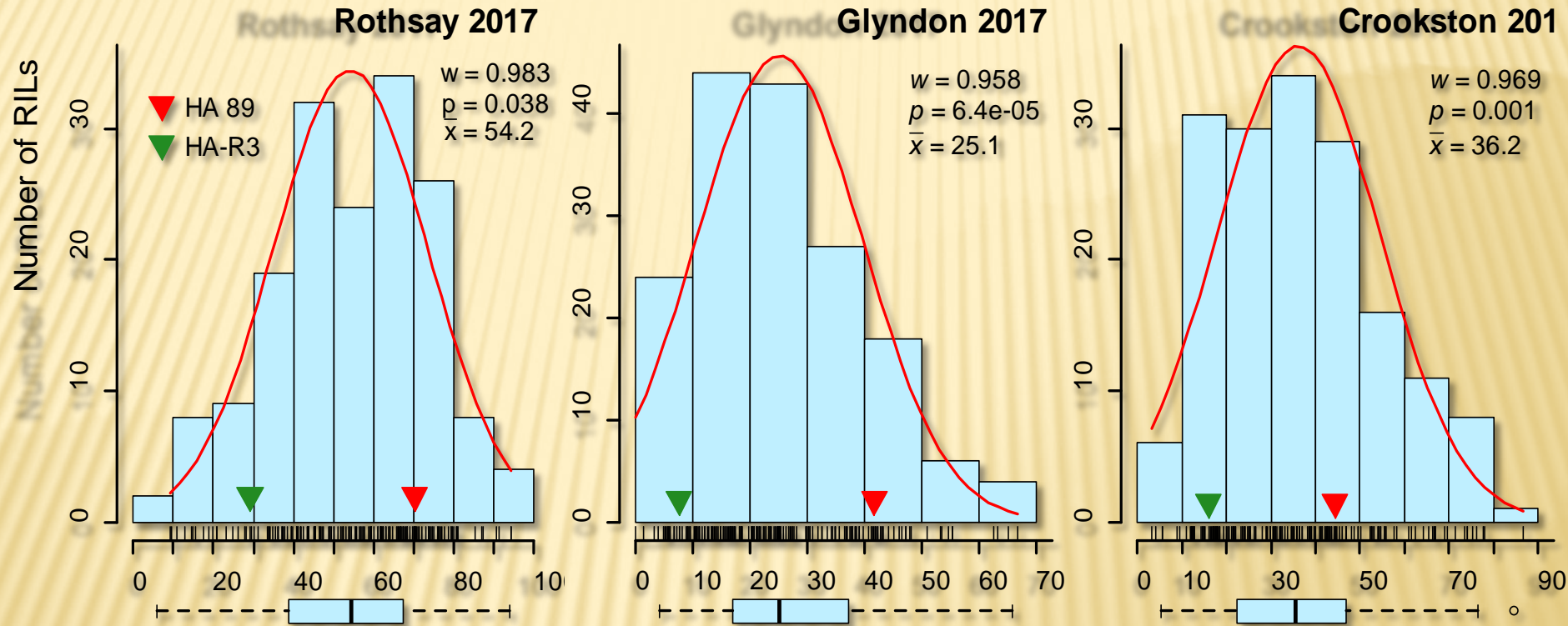
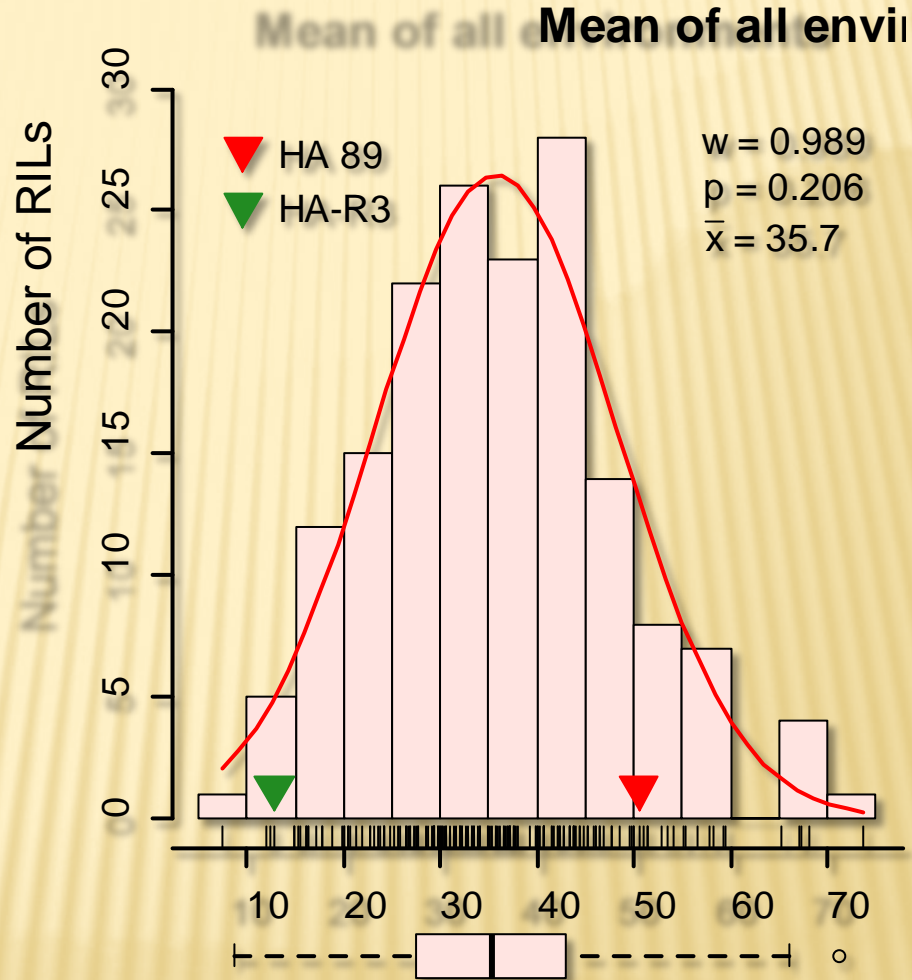


Fig 2. Phenotypic frequency distributions of PSC disease incidence (DI) for HA 89/HA-R3 RIL population evaluated in 2017

RESULTS

Fig 3. Mean phenotypic frequency distributions of PSC DI for the RIL population across six environments



RESULTS

Table 1. Analysis of variance (ANOVA) of PSC disease incidence (DI) for HA 89/HA-R3 RIL population evaluated in six environments

Component	df	Variance estimate	Confidence limit (0.05)		F/Z value	Pr > F/Z
			lower	upper		
Env	5	-	-	-	39.19	<.0001
Rep (Env)	12	$\sigma^2_r = 12.66$	6.08	40.63	2.17	0.0148
Genotype	165	$\sigma^2_g = 116.41$	88.66	159.65	6.70	<.0001
Genotype x Env	825	$\sigma^2_{ge} = 152.94$	131.49	180.13	12.47	<.0001
Error	1979	$\sigma^2_e = 264.41$	248.63	281.76		

Analysis was performed using PROC MIXED of SAS version 9.4. All factors were treated as random effects except environment

RESULTS

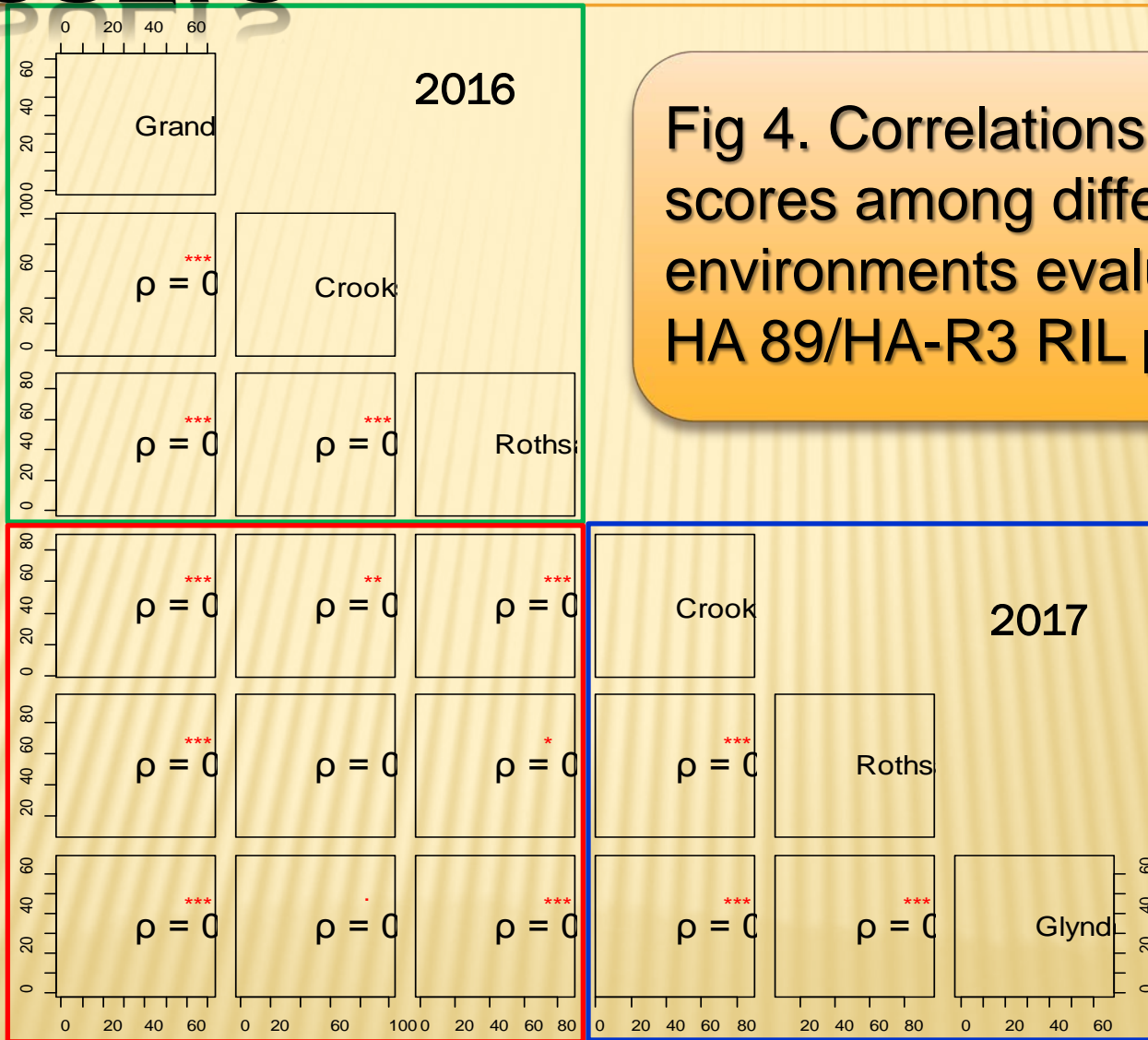


Fig 4. Correlations of PSC DI scores among different environments evaluated for HA 89/HA-R3 RIL population

FUTURE PLAN

Genotype-by-sequencing (GBS) technology will be used for genotyping of HA 89/HA-R3 RIL population with large numbers of SNP markers

Complete linkage map construction and QTL mapping in the HA 89/HA-R3 RIL population

Design PCR based primers for SNPs flanking important QTL to use in marker-assisted PSC resistance sunflower breeding

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