



## The Effect of Artificial Selection on Phenotypic Plasticity: The Genotype by Environment Interaction Project in Maize

de Leon, N., D. Jarquin, M.C. Romay, J. Gage, S.M. Kaeppler, E.S. Buckler, A.J. Lorenz, G2F Consortium

PAG CSSA: Translational Genomics Workshop  
January 9<sup>th</sup>, 2016

[www.Genomes2Fields.org](http://www.Genomes2Fields.org)

### Premise:

- ✦ Phenotypic plasticity (or G X E) → ability of a single genotype to produce different phenotypes in response to different environments.
- ✦ Plasticity has a genetic foundation and therefore is affected by selection
- ✦ This adaptation capacity is directly related to the extent and nature of standing genetic variation
- ✦ Has selection for high productivity decrease available genetic variation that controls plasticity in crop species?
- ✦ Also understanding the types of genetic architectures and modulation mechanisms controlling G X E will provide opportunities to enhance prediction ability on variable environments
- ✦ Using the platform of the **Maize G X E project** that is part of **Genomes to Fields Initiative** to test these hypotheses

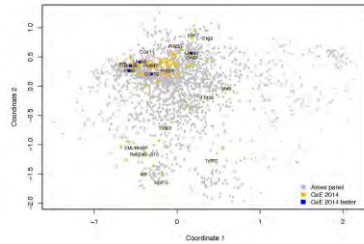


### Experimental Description - Genotypes:

- ✦ 853 hybrids from 374 inbreds crossed by 4 testers and evaluated across 22 environments (243 hybrids per location)
- ✦ GBS data collected for all lines
- ✦ Synthetic genotypes of hybrids created from inbred parent genotypes



Cintia Romay  
Cornell Univ



### Experimental Description – Environments in 2014:



Figure generated by Darwin Campbell

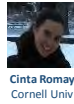
### Experimental Description - Phenotypes:

- ✦ Plant Height
- ✦ Days to Pollen
- ✦ Days to Silk
- ✦ Yield (bushels/acre)

### Hypothesis 1:

- ✦ Selection tends to reduce variation in genomic regions that are relevant for that trait, therefore regions that underwent reduction in variability due to selection should explain less of the G X E variability observed compared to regions under no selection





### Analysis – Definition of Differential $F_{ST}$ Regions:

- Categories were defined by comparing  $F_{ST}$  on 20 SNP windows for 60 inbred lines from the HapMap (30 on each extreme tropical vs temperate) based on MDS distance (confirmed by Jeff Ross-Ibarra)
- Contrasting  $F_{ST}$  regions: Low ( $F_{ST} < 0.15$ ) and High ( $F_{ST} > 0.5$ ) SNPs in the genome across all hybrids and environments
- Set of 9,194 SNPs with high  $F_{ST}$  values and random samples of 9,194 SNPs from low  $F_{ST}$



### Analysis – Variance Components Analysis:

- Use variance components approach in Gusev et al., 2014 (Am. J. of Hum Gen, 95:535-552)
- Method calculates  $r^2$  between the phenotype and the prediction for SNPs from specific functional category (in our case differential  $F_{ST}$ )
- We analyzed variance components to evaluate if interaction between the studied regions (i.e.: differentially selected regions) and environment show significant differences in the amount of phenotypic variability explained



### Analysis – Model Details:

Random effects model:

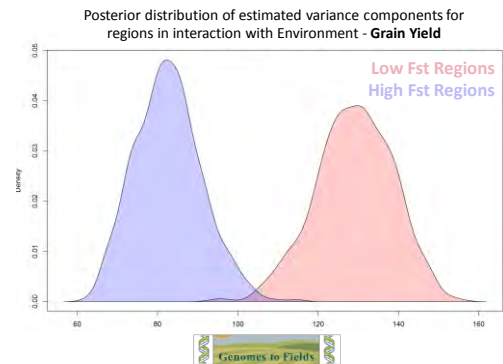
$$y = L + E + g + g_L \times E + g_H \times E + e$$

where:

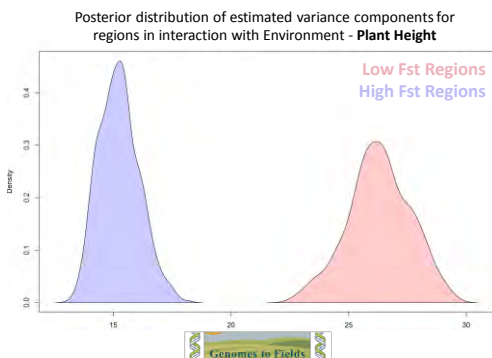
- $y$  is the vector of phenotypes
- $L$  &  $E$  = vectors of hybrid and environmental effects, such that  $L \sim N(0, I\sigma_L^2)$  and  $E \sim N(0, I\sigma_E^2)$
- $g$  = the vector of genomic values following a multivariate normal distribution such that  $g \sim N(0, G\sigma_g^2)$ ,  $G$  being the Genomic Relationship Matrix computed using all the available markers
- The interaction terms were included through covariance structures (following Jarquin et al., 2014). Thus  $g_L \times E \sim N(0, Z_L G_L Z_L' g \# Z_E Z_E')$  and  $g_H \times E \sim N(0, Z_H G_H Z_H' g \# Z_E Z_E')$
- $G_H$  computed using the 9,194 SNPs with high  $F_{ST}$  values and  $G_L$  computed taking random samples of 9,194 SNPs from the low  $F_{ST}$
- analysis repeated 1,000 time



### Results – Assessment of Effect of Selection:



### Results – Assessment of Effect of Selection:



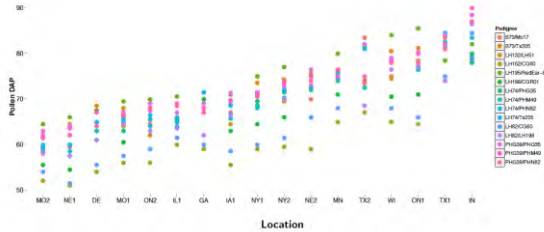
### Hypothesis 2:

- Is  $G \times E$  modulated by the genetic architecture of the trait or are there alternative sources of control?
  - If  $G \times E$  is mostly due to the genetic architecture of the trait (combination of alleles, epistasis, etc) → current models that focus on assigning values to polymorphisms for prediction would work
  - If  $G \times E$  is due to more complex regulatory mechanisms → information needs to be included in the model to account for  $G \times E$
  - Important not only to make decisions of what to select, but also of what type of variation is important to keep in the germplasm for long term breeding gains



### Analysis – Quantifying Stability:

- ✦ 12 checks evaluated at 21 environments or more
- ✦ Average check at each location → reference value for location

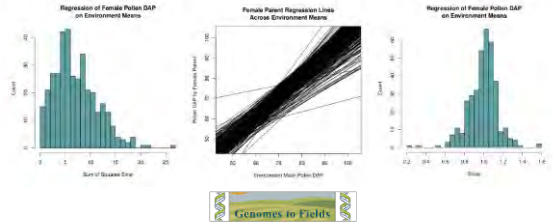


### Analysis – Definition of Parameters:



Joe Gage  
Univ of WI

- ✦ For each unique hybrid at 3 locations or more:
  - ✦ Plot hybrid value at each location against the location's reference value (average check performance)
  - ✦ Fit a line
  - ✦ Extract regression parameters: slope and SSE



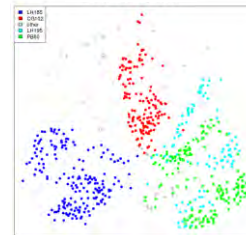
### Relationship between Regression Parameters and Stability:

- ✦ Type I/II Stability: Environmental response is parallel to mean response → quantified by slope
- ✦ Type III Stability: Small among-environment variance → quantified by SSE
- ✦ Each regression parameter was evaluated for each trait, resulting in eight parameter-trait combinations to be mapped separately



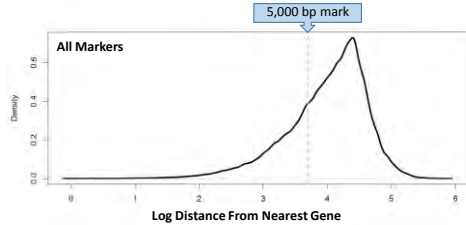
### Analysis - GWAS:

- ✦ Each phenotype and parameter-trait combination mapped separately using TASSEL 5.0
- ✦ Evaluated both additive and dominance effects
- ✦ Kinship, as well as 3 principal components covariates used to compensate for strong population structure (four males)

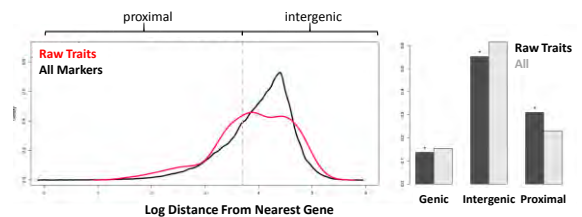


### Results - Patterns of Functional Variation:

- ✦ Classified SNPs based on their distance to the nearest gene model (following Wallace et al., 2014 – Plos Genetics):
  - ✦ Genic: Inside a gene
  - ✦ Proximal: 1-5,000 base pairs from closest gene
  - ✦ Intergenic: >5,000 base pairs from closest gene
  - ✦ Use all markers (~420k) to form a null distribution



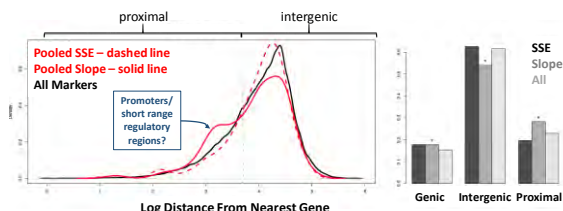
### Results – Patterns of Functional Variation:



→ 100 SNPs with the lowest p-values for each trait per se were classified as intergenic, gene-proximal or genic



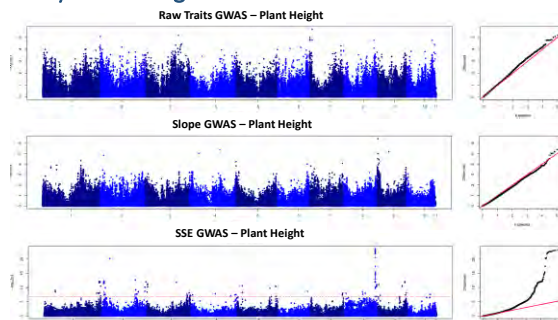
### Results – Patterns of Functional Variation:



→ 100 SNPs with the lowest p-values for each trait-parameter combination were classified as intergenic, gene-proximal or genic



### Results – Are Traits Per Se and Stability Controlled by Similar Regions?



### Take Home Messages:

- ✦ Proportion of the phenotypic variance explained for G X E differs for regions that have undergone differential selection in this dataset
- ✦ Control of stability for the traits evaluated appears to involve variants located mostly in intergenic regions
- ✦ No obvious co-localizations of stability and phenotype *per se* associations
- ✦ Need further analysis of the ontology of potential regions associated with control of plasticity



### GxE Consortium: Data Usage Disclaimer

*This presentation includes data analysis and interpretation conducted by the presenter and does not necessarily reflect the observations and conclusions of the GxE Consortium.*



### G X E Cooperators

Researchers who grew GxE trials in 2014 and 2015

- |                             |                            |                              |
|-----------------------------|----------------------------|------------------------------|
| ✦ Martin Bohn (UIUC)        | ✦ Judith Kolkman (Cornell) | ✦ Torbert Rocheford (Purdue) |
| ✦ Ed Buckler (ARS)          | ✦ Greg Kruger (UNL)        | ✦ Oscar Rodriguez (UNL)      |
| ✦ Ignacio Ciampitti (KSU)   | ✦ Nick Lauter (ARS)        | ✦ Cinta Romay (Cornell)      |
| ✦ Jode Edwards (ARS)        | ✦ Aaron Lorenz (UMN)       | ✦ James Schnable (ISU)       |
| ✦ Sherry Flint-Garcia (ARS) | ✦ Liz Lee (Guelph)         | ✦ Brian Scully (ARS)         |
| ✦ Christopher Graham (SDSU) | ✦ Natalia de Leon (IOW)    | ✦ Margaret Smith (Cornell)   |
| ✦ Candy Hirsch (UMN)        | ✦ Jonathan Lynch (PSU)     | ✦ Nathan Springer (UMN)      |
| ✦ Jim Holland (ARS)         | ✦ Steve Moose (UIUC)       | ✦ Peter Thomson (OSU)        |
| ✦ David Hooker (Guelph)     | ✦ Seth Murray (TAMU)       | ✦ Mitch Tuinstra (Purdue)    |
| ✦ Shawn Kaeppeler (UW)      | ✦ Rebecca Nelson (Cornell) | ✦ Randy Wisser (UDel)        |
| ✦ Joe Knoll (ARS)           |                            | ✦ Wenwei Xu (TAMU)           |

### Genomes To Fields Sponsors

### Genomes To Fields Collaborators

*Project planning, data management, phenotyping initiatives, and Executive Committee members*

- ◇ Martin Bohn (IUIUC)\*
- ◇ Ed Buckler (ARS)
- ◇ Darwin Campbell (ISU)\*
- ◇ James Clohessy (Cornell)
- ◇ Michael Coen (UW)
- ◇ Carolyn Lawrence Dill (ISU)\*
- ◇ Liang Dong (ISU)
- ◇ Jode Edwards (ARS)\*
- ◇ David Ertl (IA Corn)\*
- ◇ Sherry Flint-Garcia (ARS)\*
- ◇ Joseph Gage (UW)
- ◇ Jack Gardiner (ISU)\*
- ◇ Byron Good (Guelph)
- ◇ Mike Gore (Cornell)
- ◇ Patricio Grassini (UNL)
- ◇ Jerry Hatfield (ARS)
- ◇ Diego Jarquin (UNL)\*
- ◇ Shawn Kaeppler (UW)
- ◇ Liz Lee (Guelph)\*
- ◇ Natalia de Leon (UW)\*\*
- ◇ Zhizhai Liu (TAMU)
- ◇ Aaron Lorenz (UMN)\*
- ◇ Jonathan Lynch (PSU)
- ◇ Nathan Miller (UW)
- ◇ Jane Petzoldt (UW)\*
- ◇ Seth Murray (TAMU)\*
- ◇ Cinta Romay (Cornell)\*
- ◇ James Schnable (UNL)
- ◇ Pat Schnable (ISU)\*
- ◇ Nathan Springer (UMN)
- ◇ Edgar Spalding (UW)
- ◇ Srikanth Srinivasan\*
- ◇ Kelly Thorp (ARS)
- ◇ Rod Williamson (IA Corn)
- ◇ Randy Wisser (UDEL)\*
- ◇ Jianming Yu (ISU)

*GxE Coordinating Groups
†G2F co-lead
G2F Executive Committee members



### Madison-Wisconsin, June 12-17, 2016



<http://www.icqg5.org>

Follow @ICQG2016

### Questions?

