Integrating Multi-omics Data to Fine-map Wheat **Grain Weight and** Morphology Genes

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Grain width weight and are significantly associated with two nearby regions on chromosome 5A.

Yield is a balancing act



Recent advancement of the wheat reference genome assembly and genome editing tools can help facilitate the characterization of genes underlying quantitative trait loci (QTL) for yield components. Grain weight and morphology are valuable traits to consider when releasing a new wheat variety because they can impact the number of kernels it takes to fill a bushel and milling quality.



A QTL for thousand grain weight (TGW) and grain width was mapped using R/qtI (Broman et al., 2003) to a 100 Mbp region on chromosome 5AL in the W7984 Synthetic x Opata M85 spring bread wheat doubled haploid population (SynOp DH, 145-line subset of 215, Sorrells et al., 2011). Mixed models were fitted to extract BLUPs across 6 site-year combinations.





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HIFs enhance genetic resolution of QTL

Fine-mapping a causal genetic variant is **limited by crossovers** that disrupt linked markers. We developed three heterogeneous inbred families (HIFs) $F_{6.6}$ to increase genetic resolution of the 5A QTL. HIFs were selected from two SynOp recombinant inbred lines (SynOp RIL 2,039 F6 lines, Sorrells et al., 2011) based on heterozygosity across the 5A QTL. The individual progeny were phenotyped and genotyped in order to track recombination events across the 5A QTL. The resulting HIFs have highly homogenous background genomes and distinct crossovers across the QTL that can be confidently associated with the line's phenotype (Tuinstra et al., 1997).

> 7,000 individual plants genotyped phenotyped six across and generations in the field and greenhouse were narrowed down to **128 lines** with crossovers that limit the QTL region of significance. Mixed models were fitted to extract BLUPs across 2019 replicated headrows.

HIFs



Comparing HIF crossovers further explains two QTL peaks

2019 HIFs grown in headrows and genotyped with KASP markers across the 5A QTL were analyzed with R/qtl, narrowing the significant sequence to a 9.9 and 10.5 Mbp region.

Tests for epistasis between the most significant markers were inconclusive because a simple linear interaction







grain development time sequence analysis

model cannot estimate an interaction from the present genotype frequencies (341.5 : 380.8 Mbp, 73 o:o, 54 w:w, 1 o:w, 0 w:o). The two QTL peaks could be due to multiple causal variants, or linkage. **HIF I is** the only line with a recombination event resulting in o.w genotype for the most significant markers and will be very useful for comparison and understanding the two-peak phenomenon in a gene expression study.



ns: p > 0.05, *: $p \le 0.05$, **: $p \le 0.01$, ***: $p \le 0.001$, ***: $p \le 0.001$, ****: $p \le 0.0001$

grain morphology for 10 HIF genotypes with positive or negative 5A QTL causal variant alleles were tracked during the 2019 field season (Days post anthesis, DPA). grain width, fresh weight and dry weight were measured with 2 replicates / genotype, 10 spikes / time point and 10 kernels / spike. Significant difference in grain morphology is measurable 10-16 DPA for 5A QTL HIFs grown in Ithaca, NY. The difference in phenotypes for the positive and negative causal variant genotypes is likely due to a gene expression event prior to 10 DPA, providing us with a high confidence window for timing RNA extraction.

Next steps: RNA-seq

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Relying on large populations over many generations to detect crossovers and capture finer resolution of the QTL is resource limiting. Our attention has now turned to gene expression to facilitate characterization of the underlying causal grain weight and morphology gene, in a greenhouse environment.

Opata vs W7984 QTL haplotype, 8 DPA

Differential expression genome-wide indicates 5AS missing, potential epistasis/compensation on 1A. Further genome assembly and differential expression analysis required to understand relationship to TGW/GW QTL.

High confidence gene annotation (Alaux et al., 2018) and gene expression reports from wheat-expression.com indicate that 24 out of the 113 candidate genes have zero gene expression in the kernels prior to 15 DPA.

36 gene expression studies 118 grain tissue samples 65 different varieties of wheat Grey text: 0.0 tpm before 15 DPA

Borrill et al., 2016; Ramírez-González et al., 2019 Figure code credit Dr. Shantel A. Martinez





Alaux et al., 2018 doi: 10.1186/s13059-018-1491-4 Borrill et al., 2016 doi: 10.1104/pp.15.01667 Broeman et al., 2003 doi: 10.1093/bioinformatics/btg112

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