**Brief Communication**

**Burden tests can be used to map causal genes for a simple metabolic trait in an exome-sequenced polyploid mutant population**

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Forward genetic screens are an excellent tool to assign gene function, but it is often necessary to employ map-based cloning to identify the causal genes. This can be laborious and represents a bottleneck in plant fundamental and applied research. With advances in DNA technology, it is becoming increasingly affordable to sequence large populations. Krasileva et al. (2017) exome sequenced tetraploid and hexaploid wheat ethyl methanesulphonate (EMS) mutagenized populations, primarily to facilitate reverse genetic screens. Gene redundancy allows a very high mutant load of 35–40 mutations per kilobase, and the populations of ~1500 and ~1200 lines each harbour ~22–23 missense or truncation mutations per gene. Here, we show that burden tests, a simple form of rare-variant association analysis developed for human disease genetics (Lee et al., 2014), can be used to identify causal genes in the hexaploid wheat (*Triticum aestivum*) cv. Cadenza mutant population, without the need for map-based cloning.

The statistical power to detect association with rare variants is very limited (Lee et al., 2014), and most mutations in the Cadenza EMS population are singletons (Krasileva et al., 2017). Burden tests work by collapsing multiple variants within a gene (or other functional groups) into a single test score, thereby increasing frequency and providing greater power (Lee et al., 2014). However, this power relies on the selected variants mostly being causal and having the same direction and magnitude of effect (Lee et al., 2014). Such assumptions likely hold for mutant populations where causal variants are most frequently deleterious (Meinke, 2013), and their severity can be predicted from sequence analysis (Kumar et al., 2009). The absence of genetic structure in mutant populations should simplify association studies and collapsing homoeologous groups, that lack functional divergence in ‘recent’ polyploids like wheat (Krasileva et al., 2017), should also improve power.

To investigate whether burden tests can be applied to the Cadenza population, we measured the fatty acid composition of lipids in individual *M₂* grains (caryopses) from 1188 exome-sequenced lines using gas chromatography and calculated the proportion of unsaturated fatty acids that are polyunsaturated (ω-6 desaturation efficiency or ω-6-DE), which is a simple adaptive metabolic trait (Menard et al., 2017) and a determinant of edible oil quality (Hajiahmadi et al., 2020). As summarized in Figure 1a, we extracted a list of putative deleterious mutations in the M₂ population (Krasileva et al., 2017) using BioMart within EnsemblPlants (https://plants.ensembl.org/biomart/martview) and collapsed them by gene and by homoeologous group (tripid) (Ramírez-González et al., 2018). These mutations were given equal weight and include stop codon gained, start codon lost, splice donor and acceptor variants and non-synonymous mutation with a SIFT (sorting intolerance from tolerance) score <0.05 (Kumar et al., 2009). We then performed gene and triad-based burden tests using a single-locus linear model (CMLM) implemented in GAPIT (genomics association and prediction integrated tool) (Lipka et al., 2012).

We identified three genes and two triads that are significantly (*P* < 0.05) associated with ω-6DE, after applying Bonferroni correction (Figures 1b,c and S1). The three genes *TraesCS6A02G280000*, *TraesCS6B02G309400* and *TraesCS6D02G260200* form one triad and are predicted to encode homologues of FATTY ACID DESATURASE 2 (FAD2) (Hajiahmadi et al., 2020). FAD2 is a microsomal ω-6 fatty acid desaturase that is known to control ω-6DE in *Arabidopsis thaliana* seeds (Menard et al., 2017; Okuley et al., 1994). Hexaploid wheat contains eleven putative FAD2 genes (Hajiahmadi et al., 2020), and *TraesCS6A02G280000* (*TaFAD2.1*), *TraesCS6B02G309400* (*TaFAD2.6*) and *TraesCS6D02G260200* (*TaFAD2.8*) are the most strongly expressed in developing grains of cv. Azhurnaya (Figure 1d; Ramírez-González et al., 2018). The second triad (*TraesCS7A02G378300*, *TraesCS7B02G280100* and *TraesCS7D02G375100*) encode putative homologues of REDUCED OLEATE DESATURATION 1 (ROD1), which supplies FAD2 with substrate (Lu et al., 2009).

*TaFAD2* and *TaROD1* transcripts are average length for wheat (~1.6 and ~1.5 kb), encoding proteins of ~390 and ~300 amino acid residues, respectively. The 1188 *M₂* lines that we screened contained 22–24 putative deleterious mutations in each *TaFAD2* gene, and 6–9 in each *TaROD1* gene, when the *M₂* generation was exome sequenced (Krasileva et al., 2017). To confirm that disruption of the *TaFAD2* genes causes a reduction in ω-6DE, we selected two independent lines with mutations in each gene that had lowω-6DE in our screen (Figure 1e). We backcrossed them to wildtype and identified five homozygous and five wildtype segregant BC₁F₂ plants using KASP (kompetitive allele specific PCR) assays and further confirmed their genotype by DNA sequencing (Krasileva et al., 2017). We then analysed the fatty acid composition of their BC₁F₂ grains and found that ω-6DE is significantly (*P* < 0.05) lower in all the homozygous *TaFAD2* mutants (M) versus wildtype (WT) segregants (Figure 1e). The decrease in ω-6DE is small (~9%), but owing to the high broad-sense heritability of the trait (*H²~0.9*), the effect size is very large (Cohen’s *d* > 0.8).
In conclusion, we show that gene and homoeologous group-based burden tests can identify causal genes for a simple metabolic trait in an exome-sequenced polyploid mutant population. Many rare-variant association analysis methods have been developed and may be applicable, including burden tests with more sophisticated weighting, variance-component and combined tests (Lee et al., 2014). We have collapsed point mutations in the Cadenza population, but deletions are also present.

Figure 1 Applying burden tests to the Cadenza exome-sequenced EMS population to identify genes that control grain \(\omega-6\) fatty acid desaturation efficiency (\(\omega-6\)DE). (a) Workflow diagram. White boxes show resources created by Krasileva et al. (2017). Manhattan plots showing trait associations with (b) 82,950 genes and (c) 17,616 triads. Collapsed variant frequency threshold \(= 0.002\). Dotted line marks significance threshold after Bonferroni correction for \(\alpha = 0.05\). Putative \(TaFAD2\) and \(TaROD1\) genes are highlighted. Quantile–quantile plots shown on right. (d) \(TaFAD2\) expression in grains at hard dough stage (mean \(\pm\) SE, \(n = 3\)). tpm is transcripts per kilobase million. RNA-seq data from Ramírez-González et al. (2018). (e) Box plots for \(\omega-6\)DE in M4 grain from all mutant lines containing putative deleterious (D) and non-deleterious (ND) variants in each \(TaFAD2\) gene (\(n = 22–1166\)) and from two independent BC1F2 homozygous mutants (M) and their wild type segregants (WT) (\(n = 5\)). Asterisks denote significant differences (\(P < 0.05\), unpaired Student’s t-test). Cadenza line numbers and \(TaFAD2\) mutations leading to amino acid substitutions or premature stop codons* are 0277 (W107*), 0290 (P31S), 1569 (W107*), 1235 (L347F), 1366 (Q167*) and 1423 (W92*).
(Krasileva et al., 2017) and could be included. The gene redundancy that exists in polyploid mutant populations likely provides a trade-off between power and effect size when applying burden tests. Redundancy allows polyploids to tolerate high mutant loads (Krasileva et al., 2017), providing smaller populations with more collapsible variants per gene (and homoeologous group). However, redundancy also hides the phenotypic effects of variants (Krasileva et al., 2017). It is intuitive that more heritable traits that are controlled by fewer (and larger) genes will likely be more amenable to genetic dissection using burden tests. Mutant populations of tetraploid wheat (Krasileva et al., 2017) and many other polyploid crops such oilseed rape (Brassica napus) and false flax (Camelina sativa) might also be amenable to burden tests.

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Conflict of interest
The authors declare no conflicts of interest.

Author contributions
P.J.E. conceived the idea and wrote the manuscript. G.M. and P.J.E. conducted the experiments.

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Supporting information
Additional supporting information may be found online in the Supporting Information section at the end of the article.

Figure S1 Complete repeat of the burden test experiment shown in Figure 1b,c.