

Testing pleiotropy vs. separate QTL in multiparental populations

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Abstract

High-dimensional phenotypes in multiparental populations provide new opportunities for understanding complex trait architecture. New analysis tools are required. We developed a test of pleiotropy vs. separate QTL for multiparental populations. We applied it to find evidence for separate QTL for behavioral phenotypes in a 2-cM region in 261 Diversity Outbred mice. We share our methods in an R package, [qtl2pleio](https://github.com/BoehmLab/qtl2pleio) (<https://github.com/BoehmLab/qtl2pleio>).

Introduction

Experimentalists can now measure tens of thousands of phenotypes with high sequencing and mass spectrometry. Multiparent populations enable high-resolution QTL mapping. Together, high-dimensional phenotypes and multiparental populations can reform complex trait genetics. New analysis tools, such as our test of pleiotropy vs. separate QTL, are needed.

Background

Jiang and Zeng (1995) developed a pleiotropy vs. separate QTL test for two-parent crosses. Applied to two traits that map to a single genetic region. Pleiotropy is the null hypothesis. Separate QTL is the alternative hypothesis. Perform a two-dimensional QTL scan. Calculate likelihood ratio test statistic.

Challenges

Redundancy: Multivariate polynoms, random effects. Right transfer times, δ fixed effects. Test statistic calibration: Parametric bootstrap test.

Test procedure

Fit the model $Y = XB + G + E$ for each ordered pair of markers $G \sim \text{MN}(0, K, V_G)$ $E \sim \text{MN}(0, I, V_E)$ Y contains allele probabilities B contains allele effects Calculate likelihood for each model fit Test statistic: $\max_{\beta, \gamma} L_{\beta, \gamma}(B, S; \alpha)$ Parametric bootstrap to get p -value

Behavioral genetics

Logan et al. (2013) and Beale et al. (2014) genotyped and phenotyped 261 Diversity Outbred mice. Identified δ genes as the chromosomes δ gene affecting "Test plate latency" at 37 cM. Identified chromosome QTLs for "Percent time in light" and "Distance traveled in light" at 35 cM. Motivated us to ask if δ gene also affects "Percent time in light" and "Distance traveled in light".

3 pairwise tests

"Percent time in light" & "Test plate latency" $p = 0.119$ "Distance traveled in light" & "Test plate latency" $p = 0.198$ "Percent time in light" & "Distance traveled in light" $p = 0.071$

Conclusions

Evidence for two separate QTL affecting the 3 phenotypes. QTL effects both "Distance traveled in light" and "Percent time in light". Separate QTL contains δ gene and affects "Test plate latency".

Future directions

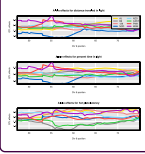
Expressive expression data from Keller et al. (2018). Expressive QTL, transfer direction. Statistical power studies.

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References

Jiang D, Zeng A (1995) Multiple trait analysis of genetic mapping for quantitative trait loci. *Genetics* 149: 1029–1042. <https://doi.org/10.1534/genetics.149.4.1029> Beale M, Logan M, Broman K, et al. (2014) Genetic mapping of behavioral traits in the Diversity Outbred mouse population. *Genetics* 196: 1029–1042. <https://doi.org/10.1534/genetics.149.4.1029> Logan M, Broman K, et al. (2013) Genetic mapping of behavioral traits in the Diversity Outbred mouse population. *Genetics* 196: 1029–1042. <https://doi.org/10.1534/genetics.149.4.1029> Keller M, et al. (2018) Expressive expression data from Keller et al. (2018). <https://doi.org/10.1534/genetics.149.4.1029>

Allele effects plots

Profile LOD plots for pairwise