

# Genotype Conditional Association Test Vignette

Wei Hao, Minsun Song, John D. Storey

November 25, 2024

## 1 Introduction

The **gcatest** package provides an implementation of the Genotype Conditional Association Test (GCAT) [2]. GCAT is a test for genetic association that is powered by Logistic Factor Analysis (LFA) [1]. LFA is a method of modeling population structure in a genome wide association study. GCAT performs a test for association between each SNP and a trait (either quantitative or binary). We have shown that GCAT is robust to confounding from population structure.

## 2 Sample usage

We include a sample dataset with the package. **sim\_geno** is a simulated genotype matrix. **sim\_trait** is a simulated trait. There are 10,000 SNPs and 1,000 individuals. The first five SNPs are associated with the trait. This simulations were done under the Pritchard-Stephens-Donnelly model with  $K = 3$ , with Dirichlet parameter  $\alpha = 0.1$  and variance allotment in the trait corresponding to 5% genetic, 5% environmental, and 90% noise. This dataset is simulated under identical parameters as the PSD simulation in Figure 2 of the paper [2], except that we have adjusted the size of the simulation to be appropriate for a small demo.

```
library(lfa)
library(gcatest)
dim(sim_geno)

## [1] 10000 1000

length(sim_trait)

## [1] 1000
```

### 2.1 gcat

The first step of **gcat** is to estimate the logistic factors:

```
LF <- lfa(sim_geno, 3)
dim(LF)

## [1] 1000 3
```

Then, we call the **gcat** function, which returns a vector of p-values:

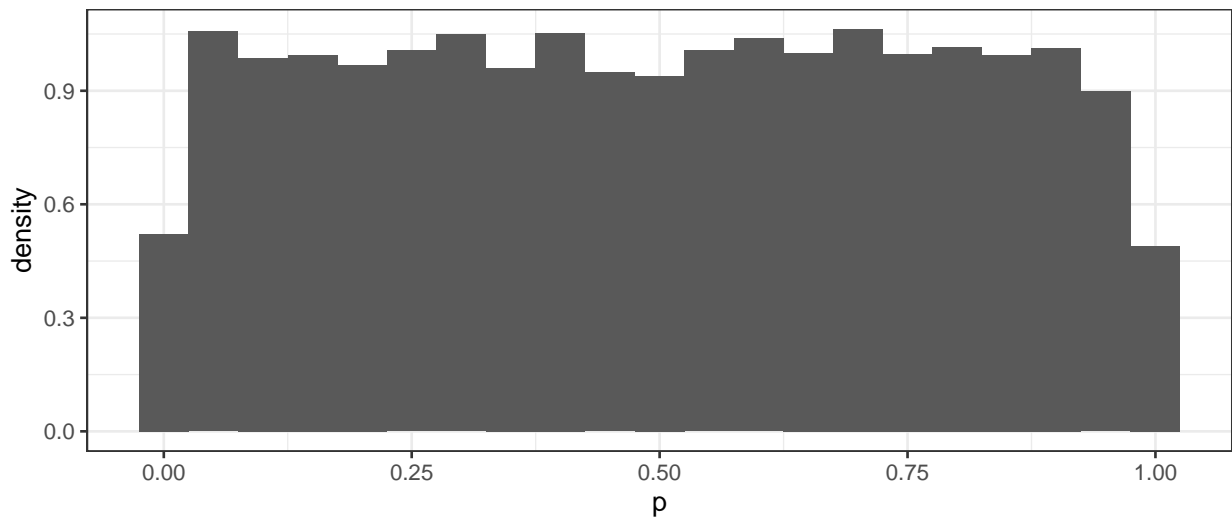
```
gcat_p <- gcat(sim_geno, LF, sim_trait)
```

We can look at the p-values for the associated SNPs:

```
gcat_p[1:5]
## [1] 5.873675e-01 1.570672e-01 1.240494e-03 4.799294e-01 8.607748e-11
```

And also plot the histogram of the unassociated SNPs:

```
library(ggplot2)
dat <- data.frame(p = gcat_p[6:10000])
ggplot(dat, aes(p, after_stat(density))) + geom_histogram(binwidth=1/20) + theme_bw()
```



### 3 Data Input

The **genio** package provides the function `read_plink` for parsing PLINK binary genotypes (extension: `.bed`) into an R object of the format needed for the `gcat` function. A `BEDMatrix` object (from the eponymous function and package) is also supported, and can result in reduced memory usage (at a small runtime penalty).

### References

- [1] Wei Hao, Minsun Song, and John D. Storey. Probabilistic models of genetic variation in structured populations applied to global human studies. *Bioinformatics*, 32(5):713–721, March 2016.
- [2] Minsun Song, Wei Hao, and John D. Storey. Testing for genetic associations in arbitrarily structured populations. *Nat. Genet.*, 47(5):550–554, May 2015.