

Summer Institute in Statistical Genetics Module 6: Computing for Statistical Genetics

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6. Permutation Tests

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Overview

- Permutation tests
- A mean
- Smallest p-value across multiple models
- Cautionary notes

Testing

In testing a null hypothesis we need a test statistic that will have different values under the null hypothesis and the alternatives we care about (eg a relative risk of diabetes)

We then need to compute the sampling distribution of the test statistic when the null hypothesis is true. For some test statistics and some null hypotheses this can be done analytically. The p-value for the is the probability that the test statistic would be at least as extreme as we observed, if the null hypothesis is true.

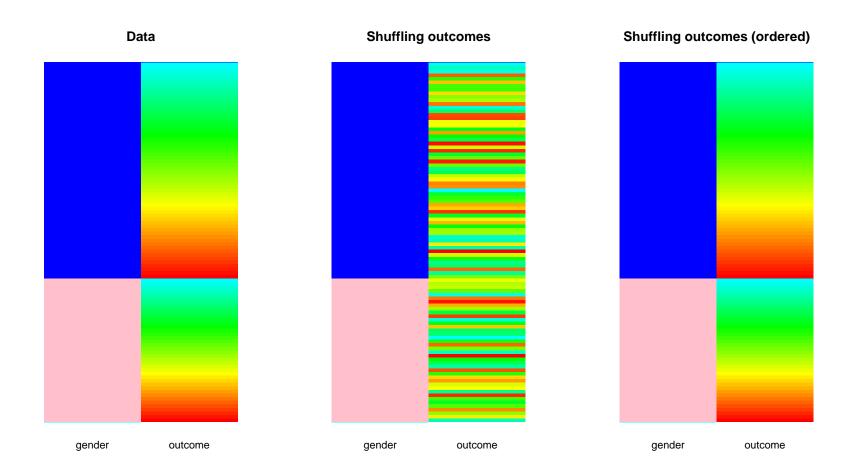
A permutation test gives a simple way to compute the sampling distribution for any test statistic, under the strong null hypothesis that a set of genetic variants has absolutely no effect on the outcome. To estimate the sampling distribution of the test statistic we need many samples generated under the strong null hypothesis.

If the null hypothesis is true, changing the exposure would have no effect on the outcome. By randomly shuffling the exposures we can make up as many data sets as we like.

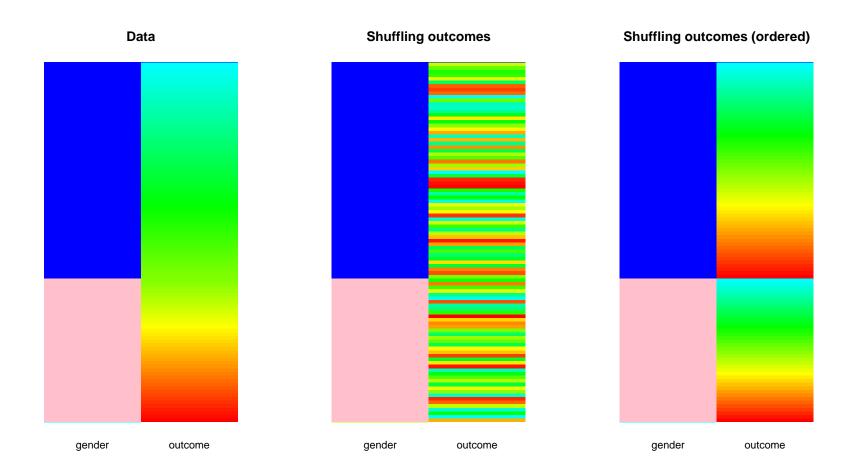
If the null hypothesis is true the shuffled data sets should look like the real data, otherwise they should look different from the real data.

The ranking of the real test statistic among the shuffled test statistics gives a p-value

Example: null is true



Example: null is false



Our first example is a difference in mean outcome in a dominant model for a single SNP

```
## make up some 'true' data
carrier<-rep(c(0,1), c(100,200))
null.y<-rnorm(300)
alt.y<-rnorm(300, mean=carrier/2)</pre>
```

In this case we know from theory the distribution of a difference in means and we could just do a t-test.

We can compare the *t*-test results to the results of a permutation test on the mean difference.

Means: *t*-test

```
> t.test(null.y<sup>~</sup>carrier, var.equal=TRUE)
Two Sample t-test
```

```
data: null.y by carrier
t = 0.5348, df = 298, p-value = 0.5932
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-0.1740603 0.3039669
sample estimates:
mean in group 0 mean in group 1
    0.067211652 0.002258386
> t.test(alt.y<sup>c</sup>arrier, var.equal=TRUE)
Two Sample t-test
data: alt.y by carrier
t = -5.0415, df = 298, p-value = 8.033e-07
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-0.8381897 -0.3675325
sample estimates:
mean in group 0 mean in group 1
                 0.4623361
     -0.1405250
```

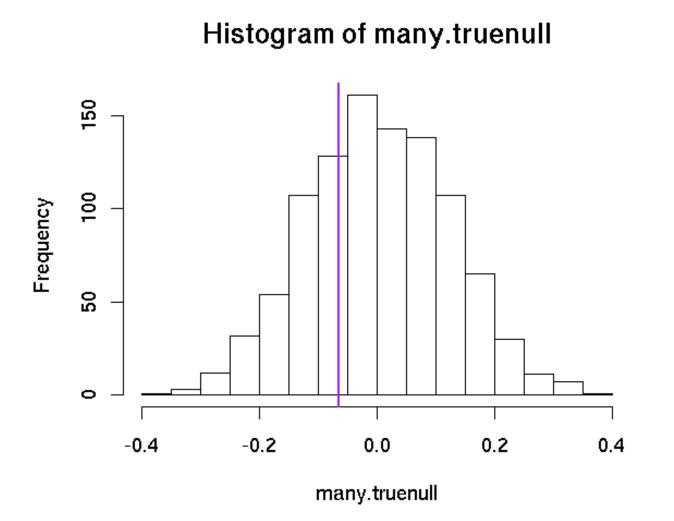
```
null.diff<-mean(null.y[carrier==1])-mean(null.y[carrier==0])
alt.diff<-mean(alt.y[carrier==1])-mean(alt.y[carrier==0])</pre>
```

```
one.test <- function(x,y) {
    xstar<-sample(x)
    mean(y[xstar==1])-mean(y[xstar==0])
    }</pre>
```

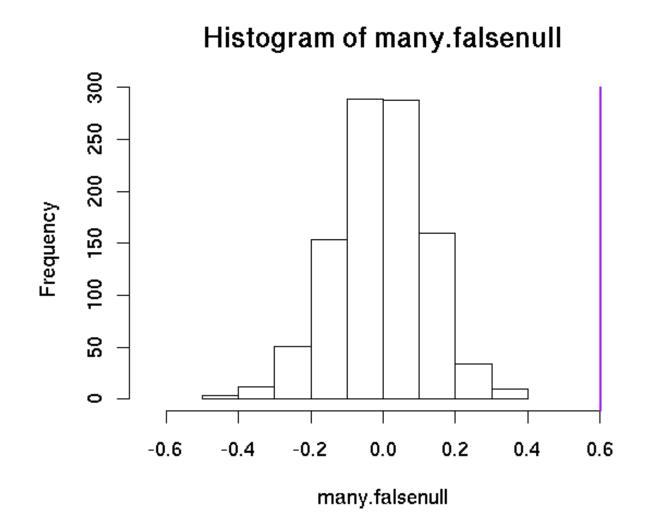
```
many.truenull <- replicate(1000, one.test(carrier, null.y))
many.falsenull <- replicate(1000, one.test(carrier, alt.y))</pre>
```

```
hist(many.truenull)
abline(v=null.diff, lwd=2, col="purple")
mean(abs(many.truenull) > abs(null.diff))
```

```
hist(many.falsenull)
abline(v=alt.diff, lwd=2, col="purple")
mean(abs(many.falsenull) > abs(alt.diff))
```



616 shuffled differences exceeed true difference: p = 617/1001



No shuffled difference exceeds true difference: p = 1/1001.

With 1000 permutations the smallest possible p-value is 0.001, and the uncertainty near p = 0.05 is about $\pm 1\%$

If we have multiple testing we may need *much* more precision.

Using 100,000 permutations reduces the uncertainty near p = 0.05 to $\pm 0.1\%$ and allows *p*-values as small as 0.00001.

A useful strategy is to start with 1000 permutations and continue to larger numbers only if p is small enough to be interesting, eg p < 0.1.

Parallel computing of permutations is easy: just run R on multiple computers.

R error messages are sometimes opaque, because the error occurs in a low-level function.

traceback() reports the entire *call stack*, which is useful for seeing where the error really happened.

Suppose our outcome variable were actually a data frame with one column rather than a vector:

```
> one.test(carrier,ywrong)
Error in '[.data.frame'(y, xstar == 1) : undefined columns selected
```

We didn't know we were calling '[.data.frame', so we don't understand the message

Debugging

```
> traceback()
5: stop("undefined columns selected")
4: '[.data.frame'(y, xstar == 1)
3: y[xstar == 1]
2: mean(y[xstar == 1])
1: one.test(carrier, ywrong)
```

so the problem happens in our line of code mean(y[xstar==1]) and is a problem with computing y[xstar==1].

We might want to have a look at y and xstar

Debugging

The post-mortem debugger lets you look inside the code where the error occurred.

```
> options(error=recover)
> one.test(carrier,ywrong)
Error in '[.data.frame'(y, xstar == 1) : undefined columns selected
```

Enter a frame number, or 0 to exit

```
1: one.test(carrier, ywrong)
2: mean(y[xstar == 1])
3: y[xstar == 1]
4: '[.data.frame'(y, xstar == 1)
```

```
Selection: 1
Called from: eval(expr, envir, enclos)
```

```
Browse[1]> str(xstar)
  num [1:300] 1 1 1 1 1 1 1 0 1 1 ...
Browse[1]> str(y)
'data.frame': 300 obs. of 1 variable:
  $ null.y: num 1.265 0.590 -0.722 0.676 -0.431 ...
```

Turn the post-mortem debugger off with

```
options(error=NULL)
```

Little point in permutation test for the mean: same result as t-test

Permutation test is useful when we do not know how to compute the distribution of a test statistic.

Suppose we test additive effects of 8 SNPs, one at a time, and we want to know if the most significant association is real.

For any one SNP the *z*-statistic from a logistic regression model has a Normal distribution.

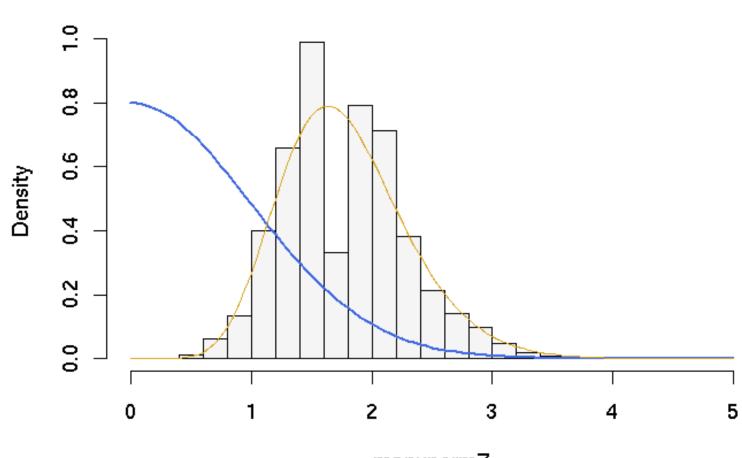
We need to know the distribution of the most extreme of eight zstatistics. This is not a standard distribution, but a permutation test is still straightforward. dat <- data.frame(y=rep(0:1,each=100), SNP1=rbinom(200,2,.1), SNP2=rbinom(200,2,.2),SNP3=rbinom(200,2,.2), SNP4=rbinom(200,2,.4),SNP5=rbinom(200,2,.1), SNP6=rbinom(200,2,.2),SNP7=rbinom(200,2,.2), SNP8=rbinom(200,2,.4))

>	head(dat)								
	у	SNP1	SNP2	SNP3	SNP4	SNP5	SNP6	SNP7	SNP8
1	0	0	0	0	0	0	1	0	0
2	0	0	1	0	1	0	1	0	2
3	0	0	1	0	1	1	0	0	0
4	0	0	0	1	1	0	0	0	0
5	0	0	1	0	1	1	0	0	0
6	0	0	0	0	1	0	1	0	1

```
oneZ<-function(outcome, snp){
    model <- glm(outcome~snp, family=binomial())
    coef(summary(model))["snp","z value"]
}
maxZ<-function(outcome, snps){
    allZs <- sapply(snps,
        function(snp) oneZ(outcome, snp))
    max(abs(allZs))
}</pre>
```

```
true.maxZ<-maxZ(dat$y, dat[,-1])</pre>
```

```
manypermZ<-replicate(10000, maxZ(sample(dat$y), dat[,-1]))</pre>
```



Histogram of manypermZ

manypermZ

The histogram shows the permutation distribution for the maximum Z-statistic.

The blue curve is the theoretical distribution for one Z-statistic

The yellow curve is the theoretical distribution for the maximum of eight independent Z-statistics.

Clearly the multiple testing is important: a Z of 2.5 gives p = 0.012 for a single test but p = 0.075 for the permutation test.

The theoretical distribution for the maximum has the right range but the permutation distribution is quite discrete. The discreteness is more serious with small sample size and rare SNPs.

[The theoretical distribution is not easy to compute except when the tests are independent.]

Permutation tests on other people's code might reveal a lack of robustness.

For example, a permutation might result in all controls being homozygous for one of the SNPs and this might give an error message

We can work around this with tryCatch()

Now oneZ will return a vector of NA if there is an error in the model fitting.

Permutation tests cannot solve all problems: they are valid only when the null hypothesis is 'no association'

Suppose we are studying a set of SNPs that each have some effect on outcome and we want to test for interactions (epistasis).

Permuting the genotype data would break the links between genotype and outcome and created shuffled data with no main effects of SNPs.

Even if there are no interactions the shuffled data will look different from the real data.

Caution: weak null hypothesis

A polymorphism could increase the variability of an outcome but not change the mean.

In this case the strong null hypothesis is false, but the hypothesis of equal means is still true.

- If we want to detect this difference the permutation test is unsuitable because it has low power
- If we do not want to detect this difference the permutation test is invalid, because it does not have the correct Type I error rate.

When groups are the same size the Type I error rate is typically close to the nominal level, otherwise it can be too high or too low.

To illustrate this we need many replications of a permutation test. We will do 1000 permutation tests for a mean, each with 1000 permutations.

```
meandiff<-function(x,trt){
mean(x[trt==1])-mean(x[trt==2])
}
meanpermtest<-function(x,trt,n=1000){
observed<-meandiff(x,trt)
perms<-replicate(n, meandiff(x, sample(trt)))
mean(abs(observed)>abs(perms))
}
```

```
trt1<-rep(c(1,2),c(10,90))</pre>
```

```
perm.p<-replicate(1000, {
    x1<-rnorm(100, 0, s=trt1)
    meanpermtest(x1,trt1)})</pre>
```

```
table(cut(perm.p,c(0,.05,.1,.5,.9,.95,1)))/1000
```

(0,0.05] (0.05,0.1] (0.1,0.5] (0.5,0.9] (0.9,0.95] (0.95,1]The p-values are too small, relative to a uniform distribution. If we reverse the standard errors we get (0,0.05] (0.05,0.1] (0.1,0.5] (0.5,0.9] (0.9,0.95] (0.95,1]If the two groups each have 50 observations we get (0,0.05] (0.05,0.1] (0.1,0.5] (0.5,0.9] (0.9,0.95] (0.95,1]

which is much better.