

Testing against a high-dimensional alternative

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Statistical Learning

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Testing against a high-dimensional alternative

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Outline

- 1 **High-Dimensional Alternatives**
- 2 **A Locally Most Powerful Test**
 - A random regression coefficient model
 - Derivation of the test statistic
- 3 **Applications**
 - Theoretical: the linear model
- 4 **Summary**

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Testing in a high-dimensional model

Testing in linear regression

- Response vector y ($n \times 1$) and design matrix X ($n \times p$)
- Model: $y \sim \mathcal{N}(X\beta, \sigma^2 I)$
- Regression coefficients β ($p \times 1$) and variance σ^2 unknown
- How to test $H_0 : \beta = \mathbf{0}$ against $H_A : \beta \neq \mathbf{0}$?

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 - What if p is larger than n ?

What is a microarray?

- Microarray measures 'gene expression'
- Central dogma of molecular biology

DNA \rightarrow RNA \rightarrow protein

- DNA is the same in every cell at every time
But need for proteins is different
 \rightarrow amount of RNA is different
- Gene expression \approx concentration of RNA of a specific gene
- Microarray: simultaneous measurement of gene expression of around 20,000 genes in one tissue

A typical microarray experiment

- Make microarrays of tissue samples of n patients
- Sample size n usually between 10 and 300.
- Typical research question:
Which genes are associated with the response?

Typical procedure

- 1 Do 20,000 univariate tests
- 2 Correct for multiple testing
- 3 Report a list of associated genes

Alternative: focus on pathways

- Pathway = set of genes with a similar function in the cell
 - Apoptosis: programmed cell death
 - Cell Cycle: the process of cell division
 - Angiogenesis: generation of blood vessels
- Our research question:

which “pathways” are associated with y ?
- More general: take any predefined set of genes
 - Genes with similar annotation in Gene Ontology / KEGG
 - Genes with similar chromosomal location
 - All genes on the microarray
- $p = \# \text{genes}$: anything from 1 to 20,000 or more

Testing association of a pathway with a response

- Gene set = a set of covariates (x_1, \dots, x_p)
→ design matrix X ($n \times p$)
- How to test for association (x_1, \dots, x_p) and y ?
- (x_1, \dots, x_p) and y are associated \iff
part of the variance of y can be predicted using (x_1, \dots, x_p)
- In a linear model: $y \sim \mathcal{N}(X\beta, \sigma^2 I)$
- No association:

$$H_0 : \beta = \mathbf{0}$$

- Some association:

$$H_A : \beta \neq \mathbf{0}$$

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The alternative hypothesis

Alternative: “at least one $\beta_i \neq 0$ ”: very general if p large

Statistical problems

- Classical tests break down if $p > n$
- There are $\beta \neq \mathbf{0}$ which have $\mathbf{r} = X\beta = \mathbf{0}$
→ it is impossible to have power against all alternatives

Focus the power

- Focus the power using a distribution on β :
- Larger density of β
= alternative β is more interesting to detect

General Unprejudicedness Assumptions:

- 1 Alternative β as interesting as $-\beta$
- 2 Alternative β as interesting as permutation $\pi(\beta)$

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- Consequence: take β with $E(\beta) = \mathbf{0}$ and $E(\beta\beta') = \tau^2 I$
- The distribution of β may have any shape
→ We don't want to assume any shape

Marginal model

- Let $L(\beta; y)$ be the likelihood of β for given y
- Let $E_{\beta|\tau^2}(\cdot)$ denote the expectation over the chosen distribution of β for given τ^2
- The marginal density of y is then

$$\bar{L}(\tau^2; y) = E_{\beta|\tau^2}\{L(\beta; y)\}$$

which can be interpreted as the likelihood of τ^2 in a new marginal model of y

- In the new model, rejecting $H_0 : \tau^2 = 0$ implies rejecting the old $H_0 : \beta = \mathbf{0}$ as the two imply the same distribution of y

A model with random regression coefficients

Inference via the marginal model: integrate β out
 $\rightarrow p$ parameters in β become one parameter τ^2

New alternative

Test $H_0 : \tau^2 = 0$ against $H_A : \tau^2 > 0$

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Model interpretation 1: Empirical Bayes

Look at the distribution of β as a prior
to Inference on β goes through hyperparameter τ^2

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Test $H_0 : \tau^2 = 0$ against $H_A : \tau^2 > 0$

Model interpretation 2: Penalized likelihood

- The distribution of β gives a penalty to the likelihood

$$\log L(y, \beta) = \log L(y|\beta) + \log L(\beta)$$

- Do maximum likelihood on penalized likelihood
- Normal β : Ridge regression
- Double exponential β : LASSO

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Model interpretation 3: Random effects model

- β random \rightarrow linear predictor $\mathbf{r} = X\beta$ random
- Moments: $E(\mathbf{r}) = \mathbf{0}$; $E(\mathbf{r}\mathbf{r}') = XX'$
- Look at r_1, \dots, r_n as a random subject effect
- Similar gene expressions \rightarrow correlated random effect
- Similar gene expressions \rightarrow correlated response

The Neyman Pearson Lemma

- Why integrate out the β parameters?

The Neyman-Pearson lemma:

The likelihood ratio test of $H_0 : \beta = \mathbf{0}$ against $H_A : \beta = \beta_A$ has optimal power to detect H_A among all tests of at most the same size

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Corollary to Neyman-Pearson:

The likelihood ratio test of $H_0 : \tau^2 = 0$ against $H_A : \tau^2 = \tau_A^2$ has optimal expected power to detect $H_A : \beta = \beta_A$ among all tests of at most the same size α

- Expectation is taken over the distribution of β_A under H_A

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- Expectation is taken over the distribution of β_A under H_A
- Drawback:** distribution of β must be fully specified

Score testing

- Score test always one-sided: $H_0 : \theta = 0$ against $H_A : \theta > 0$
- Score test = limit for $\theta_1 \downarrow 0$ of the LR test of H_0 against $H_A : \theta = \theta_1$

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- Score test also “locally most powerful test”

Corollary 2 to Neyman-Pearson:

The score test of $H_0 : \theta = 0$ has optimal slope of the power function in $\theta = 0$ among all tests of at most the same size

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- Practical advantage: no need to estimate θ

The locally most powerful test

- Back to our model with random parameters
- Instead of a Likelihood Ratio test we do a score test

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Corollary 3 to Neyman-Pearson:

The score test of $H_0 : \tau^2 = 0$ against $H_A : \tau^2 > 0$ has optimal expected slope of the power function in $\beta = \mathbf{0}$ among all tests of at most the same size

- Expectation is w.r.t. uniformly choosing a random direction in p -space
- Practical advantage: no need to estimate τ^2

The high dimensional score test statistic

- Test statistic is $S = S(0)$ with $S(\tau^2) = \frac{d}{d\tau^2} \log E_{\beta|\tau^2} L(\beta; y)$
- Is equal to:

$$S = \frac{1}{2} s' s - \frac{1}{2} \text{trace}(\mathbf{I})$$

- $s = \frac{\partial}{\partial \beta} \log L(0; y)$, the score of β
 $\mathbf{I} = -\frac{\partial^2}{\partial \beta \partial \beta'} \log L(0; y)$ is the observed Fisher inform. of β
- Nice properties:
 - *Easy: no evaluation of p -dimensional integrals*
 - *Test statistic only depends on the first two moments of β*

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- Nice properties:
 - *Easy: no evaluation of p -dimensional integrals*
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- Difficult part: What is the distribution of S ?
 - Solved for linear (exact), GLM, Cox PH (asymptotic)
 - Alternative: permutations

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Back to the linear model

- What is the locally most powerful test in the linear model?
- Get rid of the nuisance parameter $\sigma^2 \rightarrow$ profile likelihood
- For β with $E\beta = \mathbf{0}$ and $E(\beta\beta') = \tau^2 I$:

$$S = \frac{y'XX'y}{y'y}$$

- For β with $E\beta = \mathbf{0}$ and $E(\beta\beta') = \tau^2 \Sigma$:
Score test statistic:

$$S_{\Sigma} = \frac{y'X\Sigma X'y}{y'y}$$

The F-test revisited

- The F-test is only defined when $p < n$
- Equivalent test statistic $F = \frac{y'X(X'X)^{-1}X'y}{y'y}$ (beta distributed)

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The F-test as a score test

F-test is equivalent to a score test with $\Sigma = (X'X)^{-1}$

- $S = F$ in an orthogonal design (i.e. $X'X \propto I$)
- F-test optimizes power over a prior for β
with $E(\beta\beta') = \tau^2(X'X)^{-1}$ (g-prior) for τ^2 small

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Discussion: gene expression data

Analyzing micorarray data in terms of pathways

- Enables use of prior biological knowledge
- Alleviates the multiple testing problem
- Gives better reproducible results across platforms
- Global Test opens door to real inference:
→ testing hypotheses on biological mechanisms based on theory or past research

Discussion: high dimensional testing

A locally most powerful test in high dimensions

- Useable whatever the dimensionality
- Good power against interesting alternatives
- Useful applications in and outside microarray data analysis
- Applicable in survival analysis and generalized linear models

Read more?



Goeman, Van de Geer, De Kort, Van Houwelingen (2004).

A global test for groups of genes.

Bioinformatics, **20** (1) 93–99.



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JRSSB, **68** (3) 477–493.



Goeman and Oosting.

R package: globaltest

www.bioconductor.org