A practical evaluation of recent methods in high-dimensional inference

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Problem and motivation

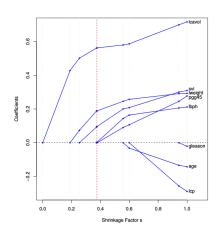
- $x \in \mathbb{R}^p, y \in \mathbb{R}$ have a joint distribution P where $y|x \sim N(x^T\beta, \sigma^2)$
- Observe $X = (x_1, ..., x_n)^T$, $Y = (y_1, ..., y_n)$ iid
- Problem: test H_i : $\beta_0 = i$ for i = 1, ..., p
- Motivation: x are SNPs (mutations), y is phenotype

Methods

	Control	p > n
Classical inference (Pearson 1930)	Marginal	No
Covariance test (Lockhart et al. 2014)	FWER?	Yes
Debiased lasso (Javanmard et al. 2014)	Marginal	Yes
Knockoffs (Barber et al. 2014)	FDR	?

The LASSO path

$$\hat{\beta}_{\lambda} = \operatorname{argmin}_{\beta} \frac{1}{2} ||X\beta - Y||^2 + \lambda ||\beta||_1$$



(Image credit: ??)



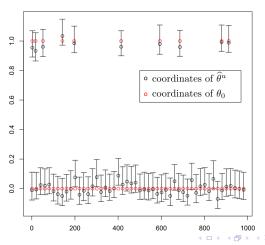
Covariance test

- (2014) Lockhart, Taylor, Tibshirani (x 2)
- Standard assumptions $Y \sim N(X\beta, \sigma^2 I) + \text{large } p$ asymptotics
- See also non-asymptotic exact test (Lee, Sun x 2, Taylor 2015)

Step	Predictor entered	Forward stepwise	Lasso
1	lcavol	0.000	0.000
2	lweight	0.000	0.052
3	svi	0.041	0.174
4	lbph	0.045	0.929
5	pgg45	0.226	0.353
6	age	0.191	0.650
7	lcp	0.065	0.051
8	gleason	0.883	0.978

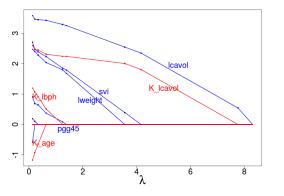
Debiased regularized M-estimators

- (2014) Javanmard and Montanari
- Standard assumptions + sparsity condition on β + large n and p asymptotics



Knockoff filter

- (2014) Barber and Candés
- Finite sample $Y \sim N(X\beta, \sigma^2 I)$, $n \leq p$, control FDR
- Extension to p > n, FWER control, etc. forthcoming...



lweight 22.5652 lcavol 20.5199 svi 4.4871 lbph 1.1865 age 0.0829 gleason 0.0387 lcp -0.2359 pgg45 -3.3742

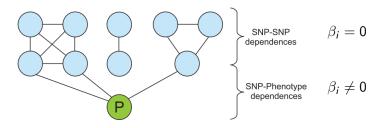
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But what's actually used in practice?

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Marginal screening	???	Yes

Regression vs Marginal Screening

Testing H_i : $\beta_i = 0$ is better than testing H_i : $Cov(X_i, Y) = 0$ when you are looking for X_i directly linked to Y



(Adapted from Mourad 2012)

Statistical Validation

- These procedures are derived under strong assumptions (lienarity, gasusianity, homoscedasticty)
- How well do they work in real data where these assumptions are violated?
- We could validate inference procedures in real data if only we knew the 'true' β , defined as

$$\beta = \mathbf{E}[\mathbf{x}\mathbf{x}^T]^{-1}\mathbf{E}[\mathbf{y}\mathbf{x}]$$

Statistical Validation

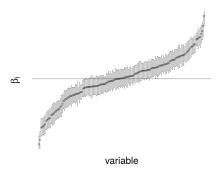
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• Possibility: take a dataset with large p and $humongous\ n$, so we can get an extremely precise estimate of β using OLS. Then test the high-dimensional inference procedures on subsamples of size $n_0 \le p < n$ of the data

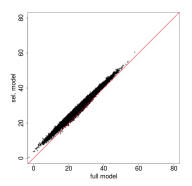
- Data with p = 163 survey questions from an online personality test, n = 49086 (after processing)
- Predict self-reported age of respondent, y, from their responses
- Is n large enough for us to confidently say which $\beta_i = 0$ (for use as ground truth?)

Coefficient estimates \pm 3 sd

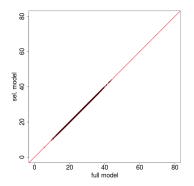


Consider declaring all variables whose intervals cross 0 to be null. Then $p_1=105$ (out of 163)

- If n were large enough, then for the selected model S we should have $\hat{y} = \sum_{i=1}^{p} X \hat{\beta}_i$ close to $\hat{y}_S = \sum_{i \in S} X_i \hat{\beta}_i$
- But...



- Here n is not large enough for p = 163
- If we reduce the dimensionality to 15 by subsampling columns, it looks more convincing that we selected the correct 10 variables



Dillemma

- It is by no means *impossible* to get large enough data to estimate high-dimensional β , with say, p > 100
- But if were *easy* to get such large *n* data... we wouldn't need these new inference techniques in the first place!

Why not use simulations?

- Simulations can be used to test robustness of the procedure
- In simulations, we can add all the nonlinearities, nongaussianity, etc. that we want

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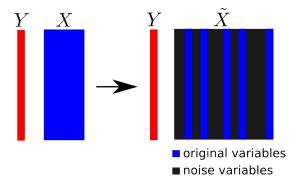
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- Advantage: We can vary simulation parameters and get a lot of insight about the procedure being tested
- **Disadvantage:** Are these simulations relevant? How can we tell the simulated models are realistic?

Idea

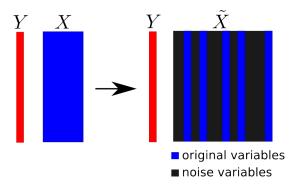
I give you real data mixed in with noise variables



• Can you identify the original columns from the noise columns?

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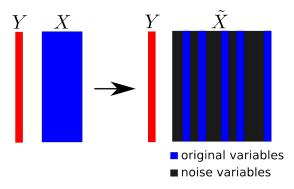
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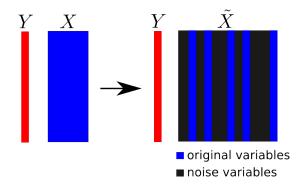
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I give you real data *mixed in* with noise variables



- Can you identify the original columns from the noise columns?
- I can test your procedure this way, because I know the ground truth!
- Caveat: this test is unrealistically 'easy' (due to lack of correlations)



- Synthetic negative controls (SNCs) are artificial columns which are correlated to X, yet still have zero (population) regression coefficients
- Suppose I give you real data + SNCs, then you apply high-dimensional inference. If you reject any SNCs, we know these are errors!
- This gives us some measure of performance on "real" data (maybe?)

- Given random vector $x \in \mathbb{R}^p$, let e be noise in \mathbb{R}^p independent of x.
- Let Γ be a fixed $p \times q$ matrix. Define synthetic negative controls $z \in \mathbb{R}^q$ by by

$$z = x'\Gamma + e$$

and let $\tilde{x} = (x, z)$, so that

$$\tilde{x}_1 = x_1, \dots, \tilde{x}_p = x_p$$

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Then

$$\forall i \in \{1, \dots, p\} : \beta_i = \tilde{\beta}_i$$
$$\forall i \in \{p+1, \dots, p+q\} : \tilde{\beta}_i = 0$$



- Recall that $\hat{\beta}_i$ is the *univariate regression* coefficient of Y on $X_{i|-i}$, where $X_{i|-i}$ is the *residual of* X_i after X_i is regressed on the other columns..
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$$P_X^{\perp} \tilde{X} = P_X^{\perp} X \Gamma_i + P_X^{\perp} E_i = 0 + E_i$$

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• Since $E_i \perp y$, we have $Cor(P_X^{\perp} \tilde{X}_{p+i}, y) = 0$, hence $\tilde{\beta}_{p+i} = 0$



• Population version: $\beta_i = 0$ if the projection of X_i on the null space of the other covariates is uncorrelated with Y

- Since $E_i \perp y$, we have $\operatorname{Cor}(P_X^{\perp} \tilde{X}_{p+i}, y) = 0$, hence $\tilde{\beta}_{p+i} = 0$
- And since $\beta_j = 0$ for all the added variables $j = p + 1, \dots p + q$, it follows that $\tilde{\beta}_i$ is unchanged for $i = 1, \dots, p$.

Using SNCs to evaluate procedures

- Take low-dimensional real data mixed with SNCs (synthetic negative controls), apply inference procedure
- Proxy for Type I error: Rejected SNCs
- Proxy for Power: Rejected original variables

A step-by-step tutorial (in R)

1. Take the prostate data

```
> data(prostate)
> x <- prostate[, 1:8]
> y <- prostate[, 9]
> colnames(x)
[1] "lcavol" "lweight" "age" "lbph" "svi"
     "lcp" "gleason" "pgg45"
> dim(x)
[1] 97 8
```

A step-by-step tutorial

2. Construct 20 synthetic negative controls

```
> GAMMA <- matrix(rnorm(8 * 20), 8, 20)
> E <- matrix(rnorm(97 * 20), 97, 20)
> sncs <- as.matrix(x) %*% GAMMA + 2 * E
> sncs <- data.frame(sncs)
> colnames(sncs)
  [1] "X1" "X2" "X3" "X4" "X5" "X6" ...
[19] "X19" "X20"
```

- 3. Create combined design matrix
- $> x2 \leftarrow cbind(x, sncs)$

A step-by-step tutorial

4. Try marginal screening

A step-by-step tutorial

```
5. Try covariance test
> library(covTest)
> covTest(lars(as.matrix(x2), y), as.matrix(x2), y)
$results
Predictor_Number Drop_in_covariance P-value
                                         0.0000
                 1
                               69.0292
                 5
                                1.5390 0.2219
                                6.8094 0.0020
                11
                                0.8559 0.4294
(Numbers 1, 5, 2 are original, 11 is a SNC)
```

A step-by-step tutorial

```
6. Try debiased lasso (code at
http://web.stanford.edu/ montanar/sslasso/)
> res <- SSLasso(as.matrix(x2), y)
[1] "10% done"
...
[1] "90% done"
> rej <- (res$up < 0) | (res$low > 0)
> names(x2)[rej]
[1] "lcavol" "lweight" "svi"
```

A step-by-step tutorial

Disclaimer!

- I am not proposing SNCs as a methodology for inference
- There is a danger of inferring that Type I error has been controlled from lack of rejection of SNCs. There are no formal guarantees of this!
- One should interpret results from experiments with SNCs in the same way one interprets simulation results with purely synthetic data

Experiments, part 1

Data	n	p_1	Linear?	Gaussian?	Constant σ^2 ?
Personality	49k	163	No	No	No
fMRI	1750	44	No	OK	No
HIV	842	207	No	Yes?	OK?
Galaxy	323	4	No	OK	No

- We add $n/2 p_1$ synthetic negative controls
- X is scaled, Γ is a gaussian matrix, $Var(E) = Var(X\Gamma)$
- \bullet Multiple trials averaging over the randomness of generating SNCs

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- Example (GWAS): There is a new disease (say, liver cancer) where we have little prior info. I have data for a 'similar' disease, (say, stomach cancer) for which I have partial information about the relevant causes. In particular, I know the most important the genes involved stomach cancer
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- Testing your procedure directly on the stomach cancer data is uninformative: if you reject a gene which is not a priori known to cause stomach cancer... it could still be a new discovery!
- Rather than test your inference procedure on the full stomach cancer data, test it on the subset of the known genes plus SNCs
- Tricky part: choosing the size of the noise for the SNCs. To be safe, try several noise levels.

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- Possible to run a Kaggle-style competition for inference rather than prediction
- Recognizing that different procedures can have differing strengths creates room for a diversity of approaches

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 is right framework for the real-world problem we are trying to solve?
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 which are a better fit to the problem.
- Ultimately, we need the practitioner to measure the value of the results we obtain from the inference procedure.

Closing thoughts

"Both the client and the statistician... must base their thinking on a recognition that their assumptions will always require review and reappraisal..."

John Tukey

Acknowledgements

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References

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