

Improving high-dimensional prediction by empirical Bayes learning from co-data

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Our group: www.bigstatistics.nl

Setting

- **Prediction or Diagnosis**
- **Primary data**
 - ▶ Variables $i = 1, \dots, p$; Individuals $j = 1, \dots, n$; $p > n$
 - ▶ Focus on binary response Y_j (e.g. case vs control)
 - ▶ Measurements $\mathbf{X}_j = (X_{1j}, \dots, X_{pj})$
 - ▶ Goal: find f such that $Y_j \approx f(\mathbf{X}_j)$
 - ▶ Here, f : *logistic regression*
 - ▶ Some form of regularization required
- **Focus**
 - ▶ Differential regularization based on prior information:
co-data

Co-data

Definition Co-data: any information on the *variables* that does not use the response labels of the primary data

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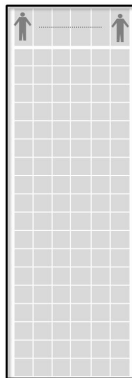
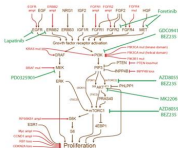


Databases



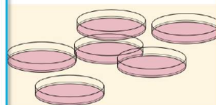
Related bio-molecules

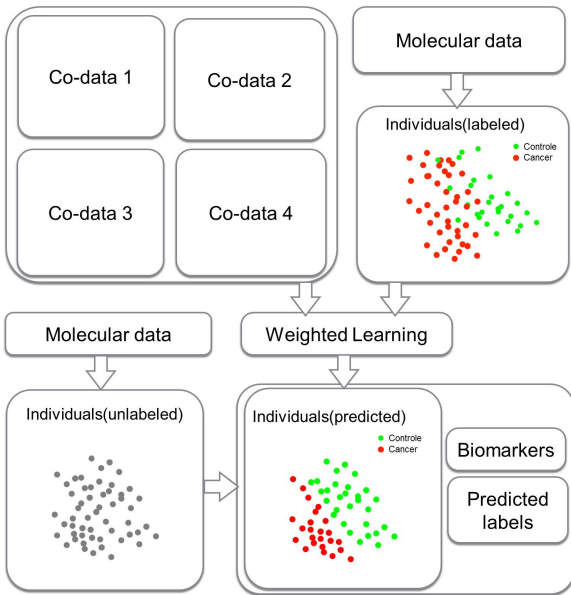
Pathways



Primary Data

Cell lines





Use of co-data

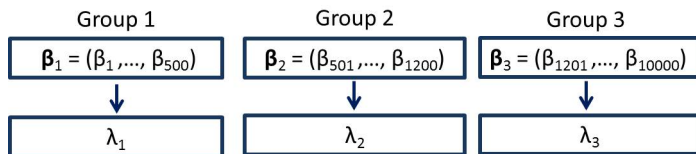
Groups: Co-data determine G prior groups of variables

Idea: Use different penalty weights $\lambda_1, \dots, \lambda_G$ across G co-data-based groups.

Use of co-data

Groups: Co-data determine G prior groups of variables

Idea: Use different penalty weights $\lambda_1, \dots, \lambda_G$ across G co-data-based groups. $G = 3$:



E.g. Ridge: $\operatorname{argmax}_{\beta} \{ \mathcal{L}(\mathbf{Y}; \beta) - \sum_{g=1}^G \lambda_g \|\beta_g\|_2 \}$

→ **CV** not attractive

Empirical Bayes (EB)

Empirical Bayes: estimate hyper-parameters from data

Relation penalty parameters \leftrightarrow hyper-parameters (prior)

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E.g. logistic ridge: $\beta_i \sim N(0, \sigma_g^2), i \in \text{group}_g; \lambda_g = 1/(2\sigma_g^2)$:

$$\operatorname{argmax}_{\beta} \{ \mathcal{L}(\mathbf{Y}; \beta) - \sum_{g=1}^G \lambda_g \|\beta_g\|_2 \} = \hat{\beta}_{\lambda} = \hat{\beta}_{\sigma}^{\text{MAP}} = \operatorname{mode}(\pi_{\sigma}(\beta | \mathbf{Y}))$$

Previous work

- **EB**: Morris, Carlin & Louis, Efron, George, Casella, Van Houwelingen, etc.
- Blog: David Robinson: varianceexplained.org
- Review: EB for high-dimensional prediction*
 - ▶ High-dimensional vs low-dimensional
 - ▶ Theory on EB estimator ($p \uparrow$) for simple linear case
 - ▶ Various EB methodologies
 - ▶ Spike-and-slab

*VdW, Münch, arXiv, to appear: *Scand J Stat*

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 - ▶ Spike-and-slab
- **Groups:** group-lasso (Meier et al.) + many versions thereof

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Formal EB: Maximum marginal Likelihood

$\beta = (\beta_1, \dots, \beta_p)$. Prior(s): $\pi_{\alpha}(\beta)$, $\alpha = (\alpha_1, \dots, \alpha_K)$

Marginal likelihood maximization:

$$\hat{\alpha} = \operatorname{argmax}_{\alpha} \operatorname{ML}(\alpha), \text{ with } \operatorname{ML}(\alpha) = \int_{\beta} \mathcal{L}(\mathbf{Y}; \beta) \pi_{\alpha}(\beta) d\beta,$$

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Optimization hard, because of the high-dimensional integral

- Laplace approximation (Shun & McCullagh, *JRSSB*, 1995)
- EM on Gibbs samples (Casella, *Biostatistics*, 2001) or on Variational Bayes approximation (Part II: Elastic Net).
- Moment estimation

EB using moments: group-regularized ridge

Estimate σ_g^2 ($\lambda_g \propto \sigma_g^{-2}$), for ridge: $\beta_i \sim N(0, \sigma_g^2)$, $i \in \text{group } g$

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Intuitive Idea:

1. Run an initial ridge regression with one λ
2. For $g = 1, 2$, consider mean squares of coefficients:

$$MS_g = \frac{1}{p_g} \sum_{i \in \text{group } g} \hat{\beta}_i^2$$

3. If MS_g is large then σ_g^2 should be large (hence λ_g small)

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More difficult, because $E(MS_g)$ depends also on variables *not in group g* (biased estimation)

EB using moment estimation[†]

Two-group example: estimate σ_1^2, σ_2^2 ($\lambda_g \propto \sigma_g^{-2}$), for ridge:

$$\beta_i \sim N(0, \sigma_1^2), i \in \text{group 1}, \beta_i \sim N(0, \sigma_2^2), i \in \text{group 2}$$

Idea: equate empirical moment(s) to theoretical ones

[†]Details: Van de Wiel et al., *Stat Med*, 2016

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$$\frac{1}{p_1} \sum_{i \in \text{group 1}} \hat{\beta}_i^2 \approx \frac{1}{p_1} \sum_{i \in \text{group 1}} E_{\beta} \left[E[\hat{\beta}_i^2(\mathbf{Y}) | \beta] \right] := f_1(\sigma_1^2, \sigma_2^2)$$
$$\frac{1}{p_2} \sum_{i \in \text{group 2}} \hat{\beta}_i^2 \approx \frac{1}{p_2} \sum_{i \in \text{group 2}} E_{\beta} \left[E[\hat{\beta}_i^2(\mathbf{Y}) | \beta] \right] := f_2(\sigma_1^2, \sigma_2^2),$$

Result: System of equations $\mathbf{b}_{\text{data}} = \mathbf{A}\mathbf{x}, \lambda_g^{-1} \propto \hat{\sigma}_g^2 = x_g$.

[†]Details: Van de Wiel et al., *Stat Med*, 2016

Shrink the shrinkage parameters[‡]

Co-data may consist of many groups (e.g. pathways)

→ $\hat{\sigma}^2 = A^{-1} \mathbf{b}_{\text{data}}$ instable → over-fitting.

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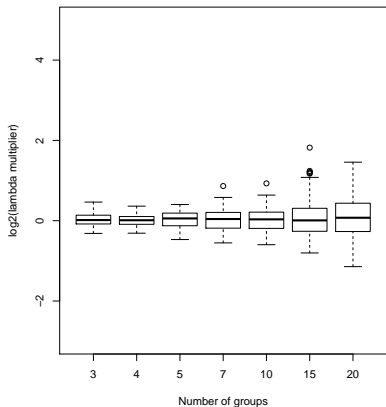
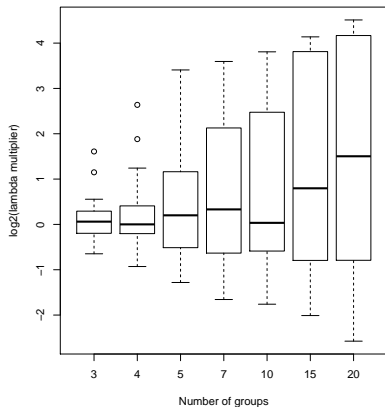
Solution: shrink A to stable target matrix, e.g. $T = \text{diag}(A)$:

$$\tilde{A}_q = qA + (1 - q)T$$

[‡]Details: Novianti et al., *Bioinformatics*, 2017

Effect of shrinkage

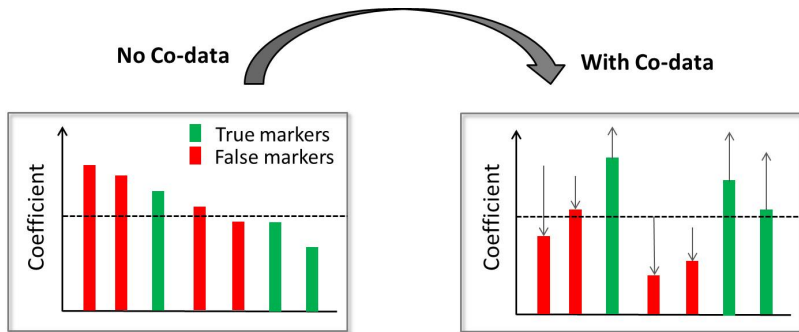
Real data, *random* groups of variables; Penalties: $\lambda_g = \lambda'_g \lambda$
 λ'_g : lambda multiplier; $\log_2(\lambda'_g)$ should $\approx \log_2(1) = 0$



Left: No Shrinkage; Right: Shrinkage

Suppose we want variable selection...

Why can co-data help?



Suppose we want variable selection...

Nicest solution: A coherent framework for EB estimation in a group-regularized elastic net setting[§]

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Ad-hoc solution:

1. Estimate group penalties from ridge regression, possibly for multiple groupings
2. Select k variables by introducing non-grouped L_1 penalty
3. Refit the model using the selected variables and their respective L_2 penalties

Software¶

R-package GRRidge, Github + Bioconductor

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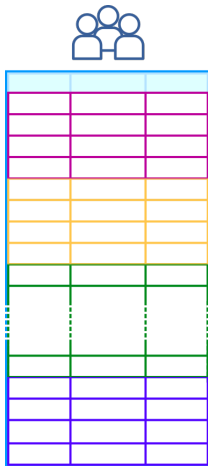
R-package GRRidge, Github + Bioconductor

- Allows iteration; CVlik as stopping criterion
- Allows *multiple* sources of co-data, as groups
- Allows *overlapping* groups, e.g. pathways
- Auxiliary functions for co-data processing
- Built-in CV for comparison with ridge & lasso

¶To be discussed during course

Part II: Group-regularized elastic net^{||}

Group of feature j : g_j .



$$g_j = 1$$

$$g_j = 2$$

\vdots

$$g_j = G$$

Group-regularized elastic net

Model

$$Y_i | \beta \sim \text{Bern}(\text{expit}(\mathbf{X}_i^T \beta)),$$

$$\beta_j \stackrel{\text{ind}}{\sim} \exp \left[-\frac{1}{2} \left(\alpha \lambda \cdot \sqrt{\lambda'_{g(j)}} |\beta_j| + (1 - \alpha) \lambda \cdot \lambda'_{g(j)} \beta_j^2 \right) \right]$$

- Shrinks estimates towards zero
- ‘Global’ α and λ determine overall shrinkage
- Elastic net with penalty weights $w_{g(j)} = (\lambda'_{g(j)})^{1/2}$:
 $\alpha \lambda |w_{g(j)} \cdot \beta_j| + (1 - \alpha) \lambda (w_{g(j)} \cdot \beta_j)^2$

Penalty parameter estimation

Cross-validation

- Prohibitively slow and unstable with even few groups

Hybrid CV and Empirical Bayes

- Fix α and estimate λ by CV for global shrinkage
- Empirical Bayes estimation of λ' by MML

Maximum marginal likelihood (MML)

$$\hat{\lambda}' = \operatorname{argmax}_{\lambda'} \int_{\beta} \mathcal{L}(\mathbf{Y}; \beta) \pi_{\lambda'}(\beta) d\beta$$

Latent variables

Extra latent variables (Polson et al., 2013; Li & Nin, 2010)

- $\omega|\beta \sim \prod_{i=1}^n \mathcal{PG}(1, |\mathbf{X}_i^\top \beta|)$, independent of Y_i
- $\beta|\tau \sim \prod_{j=1}^p \mathcal{N}\left(0, \frac{\tau_j - 1}{\lambda'_{g(j)}(1-\alpha)\lambda\tau_j}\right)$ and
 $\tau \sim \prod_{j=1}^p \mathcal{TG}\left(\frac{1}{2}, \frac{8(1-\alpha)}{\alpha^2\lambda}, (1, \infty)\right)$

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Computational reasons

- ω renders logistic part ‘easy’: it disappears in the calculations
- τ makes posterior calculations of β easier

EM algorithm

Recap Casella (2001):

$$\boldsymbol{\lambda}'^{(k+1)} = \operatorname{argmax}_{\boldsymbol{\lambda}'} \mathbb{E}_{\boldsymbol{\omega}, \boldsymbol{\beta}, \boldsymbol{\tau} | \mathbf{Y}} \left[\log \mathcal{L}_{\boldsymbol{\lambda}'}(\mathbf{Y}, \boldsymbol{\omega}, \boldsymbol{\beta}, \boldsymbol{\tau}; \boldsymbol{\lambda}'^{(k)}) \right].$$

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Exact expectation is difficult, options:

- Monte Carlo approximation: slow
- Laplace approximation: not accurate in high dimensional space
- Variational Bayes: fast and accurate (for the posterior mean)

Empirical-variational Bayes

Variational Bayes

Approximate posterior factorizes:

$$p(\omega, \beta, \tau | \mathbf{Y}) \approx q(\omega)q(\beta)q(\tau) =: Q$$

\downarrow

$$\mathbb{E}_{p(\omega, \beta, \tau | \mathbf{Y})} [\log \mathcal{L}_{\lambda'}(\cdot)] \approx \mathbb{E}_Q [\log \mathcal{L}_{\lambda'}(\cdot)] =: f(\lambda')$$

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EM algorithm

- E-step is an iterative VB algorithm itself to find Q .
- M-step, $\operatorname{argmax}_{\lambda'} f(\lambda')$, is now convex and easily solved.

Automatic feature selection

Feature selection

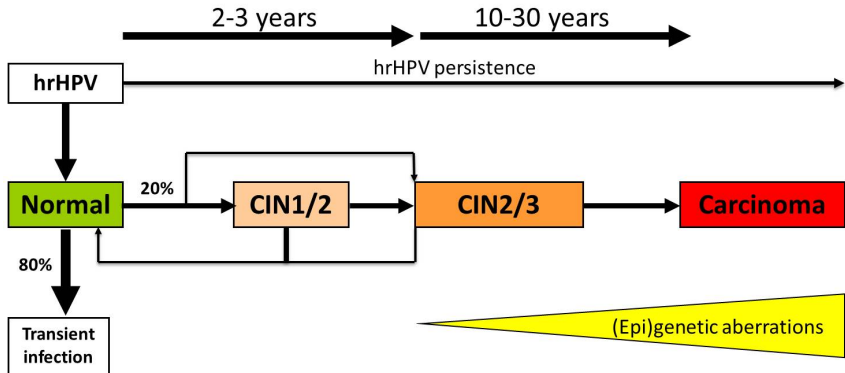
1. Plug estimated penalty parameters into frequentist elastic net:

$$\hat{\beta} := \operatorname{argmax}_{\beta} \log \mathcal{L}(\mathbf{Y}; \beta) + \frac{\alpha \lambda}{2} \sum_{j=1}^p \sqrt{\lambda'_{g(j)}} |\beta_j| + \frac{(1 - \alpha) \lambda}{2} \sum_{j=1}^p \lambda'_{g(j)} \beta_j^2$$

2. Adjust λ until desired number of features selected

- The L_1 -norm penalty term ensures automatic feature selection
- Estimated penalty multipliers may enhance predictive performance

Example: Cervical cancer



Goal: Detect CIN3 lesions, to be removed surgically

Example: Diagnostics for cervical cancer

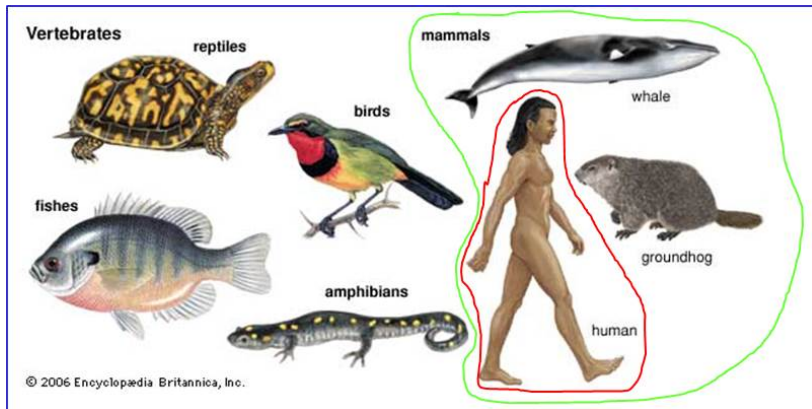
Goal: Select markers for classifying Normal vs CIN3
→ final goal is a cheap PCR assay

Data:

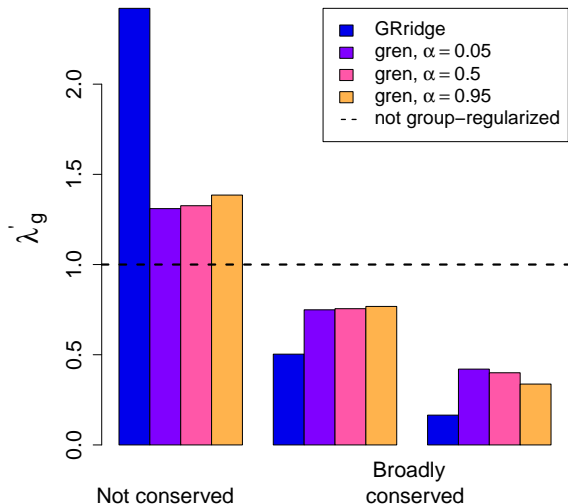
- microRNA sequencing data on *self-samples*
- $n = 56$: 32 Normal, 24 CIN3
- $p = 772$ (after filtering lowly abundant ones).
- Sqrt-transformed
- Standardized

Co-data: Conservation status

1. Non-conserved, **human** only (552)
2. Conserved across **mammals** (72)
3. Broadly conserved, across most **vertebrates** (148)



Co-data results

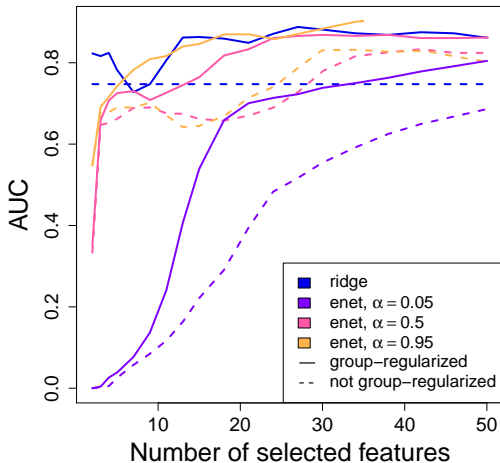


Clinician:

“That’s all nice, but does the predictive accuracy improve?”

Performance under variable selection

AUC assessed by LOOCV



Extensions, other co-data applications

Generalized ridge: covariance structures (in progress)

Random Forest: Allows flexible co-data.**

Networks: Bayesian SEM: VB + EB + prior network††

Hybrid Bayes-Empirical Bayes: $\lambda_g = \lambda \lambda'_g$, $\lambda \sim$ hyper-prior, λ'_g fixed. Example in the Review.

**Te Beest, et al., *BMC Bioinf*, 2017

††Leday, Kpogbezan, et al., *Ann Appl Stat*, 2016; *Biom J*, 2017

Thanks

Magnus Münch (Leiden Univ / VUmc)



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Cervical cancer data: Saskia Wilting (Erasmus MC), Barbara Snoek (VUmc)

Co-data: Putri Novianti (VUmc)

Stats: Wessel van Wieringen, Carel Peeters (VUmc); Aad van der Vaart (Leiden Univ)

QUESTIONS?^{††}

COURSE: Please install GRridge, gren and dependencies prior to the course.

See https://magnusmunch.github.io/co-data_learning/