

# A comparative study on PCR, PLS, Envelope and BayesPLS models

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# Overview

- Background
- Estimation methods under comparison
- Data Simulation
- Analysis, Results and Discussions

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- Cook et al. [2013] said that PLS is fundamentally an envelope in the population model

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- Using `simrel` [Sæbø et al., 2015] R-package, data with diverse nature are simulated.
- `simrel` allows to have control over latent structure (relevant component) of the data, fine analysis of strength and weakness of a models is possible

The common ground of all the methods is to best describe (fit) the multivariate linear model below,

$$y = X\beta + \epsilon \quad (1)$$

where,

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$y$	:	Response
$X$	:	Matrix of $p$ predictor variable
$\beta$	:	Regression Coefficients
$\epsilon$	:	Error $\epsilon \sim \text{NID}(0, \sigma^2)$

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Here, both  $y$  and  $X$  are considered to be centered.

# Statistical Model

All the models under this study consider a **subspace of predictor variables that is relevant for response**. They differ in the ways of finding the subspace and corresponding model estimates. The true estimates can also be written as,

$$\beta = \Sigma_{XX}^{-1} \sigma_{Xy} = \sum_{j=1}^p \frac{1}{\alpha_j} e_j e_j^t \sigma_{Xy} = \sum_{j=1}^p \gamma_j e_j$$

where,

---

$\gamma_j$	$\vdots \frac{e_j^t \sigma_{Xy}}{\lambda_j}$
$e_j$	: Eigenvector of $\Sigma_{xx}$
$\lambda_j$	: Eigenvalue of $\Sigma_{xx}$
$\sigma_{Xy}$	: Covariance between $y$ and $X$

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So, True regression estimates are the space spanned by the eigenvectors of population covariance matrix  $\Sigma_{xx}$ .

# Comparison of Methods

## PCR

- \* Regression of response on latent space of predictor
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## Envelope (MLE)

- \* Estimation using Maximum Likelihood
- \* Can not be used when predictor is larger than observations

## Bayes

- \* Estimation through MCMC approach with rotation of relevant space
- \* Heavy Computation when  $p$  is large

# Data Simulation

Models are analysed under diverse nature of data. Data are simulated using `simrel` package (R). In this study, I have included following four design;

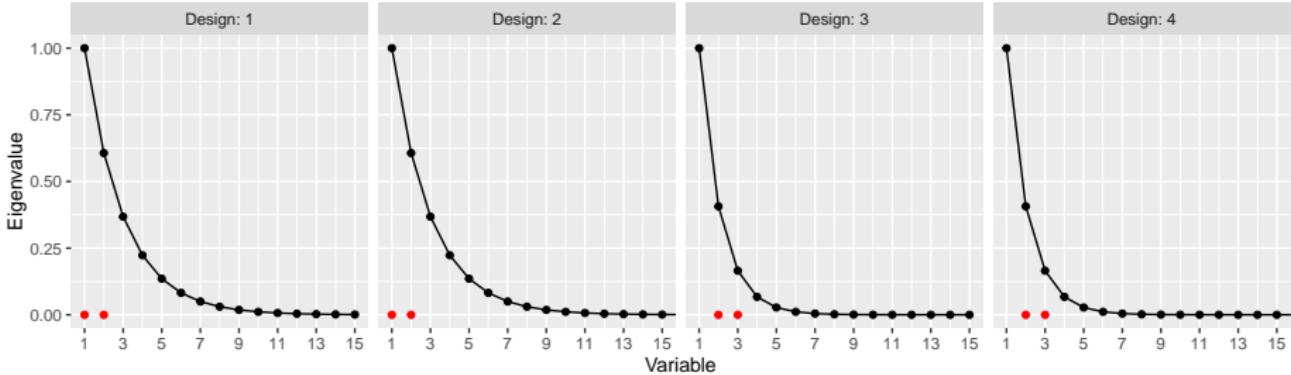
n	p	R2	relpos	gamma
50	15	0.5	1, 2	0.5
50	40	0.5	1, 2	0.5
50	15	0.9	2, 3	0.9
50	40	0.9	2, 3	0.9

---

n	:	Number of observations
p	:	Number of variables
R2	:	Variation explained by the model
relpos	:	Position of relevant components
gamma	:	Reduction factor of eigenvalue of X

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# Relevant Position and Eigenvalues



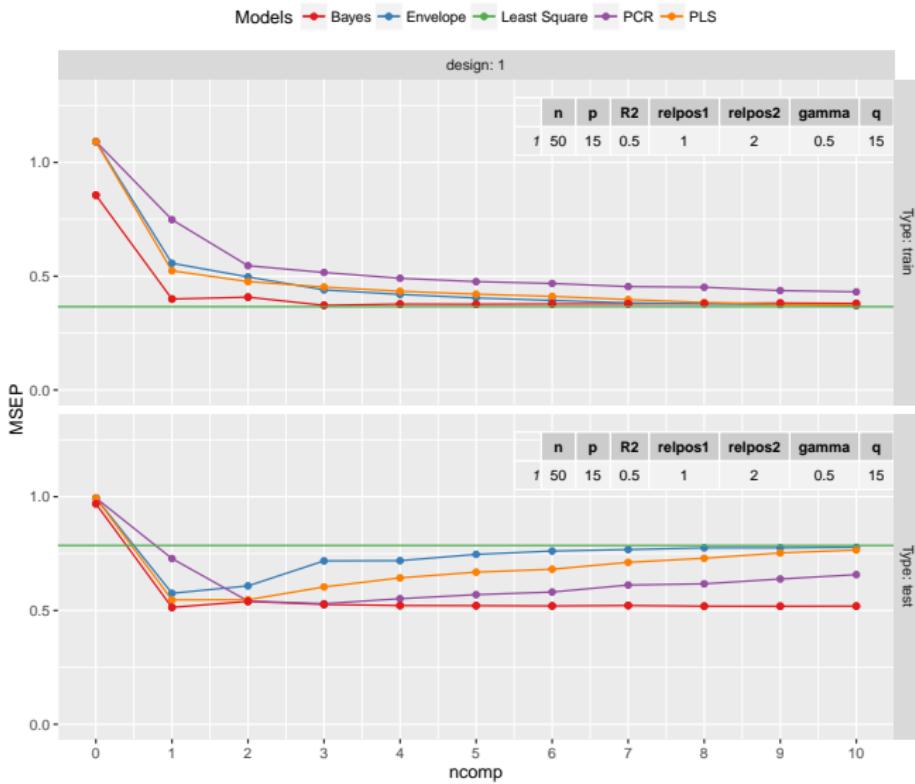
- When Relevant components are at the position of high eigenvalues, the situation is easier to model
- When Relevant components are at the position of low eigenvalues, for example 5, 10, then the most variation present in  $X$  are not relevant for  $Y$  and this will become a very difficult situation.

Models are compared on the basis of their prediction ability by measuring *test* and *training* **Mean Square Error of Prediction (MSEP)**. Mean prediction error is calculated as,

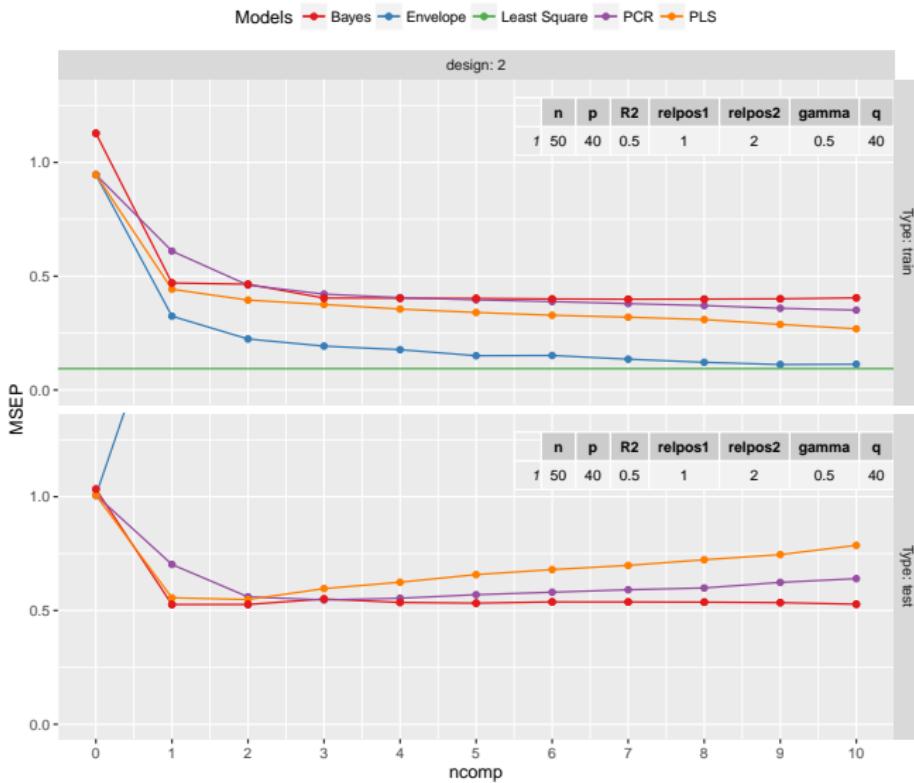
$$(\text{Prediction Error})_{\text{training}} = \frac{1}{n} \sum_{i=1}^n (\mathbf{y}_i - \hat{\mathbf{y}}_i)^2 = \frac{1}{n} \sum_{i=1}^n \left( \mathbf{y}_i - (\hat{\beta}_0 + \hat{\boldsymbol{\beta}} \mathbf{X}_i) \right)^2$$

$$\begin{aligned} (\text{Prediction Error})_{\text{test}} &= \frac{1}{n} \sum_{i=1}^{\text{ntest}} \left( \mathbf{y}_{i(\text{test})} - \hat{\mathbf{y}}_{i(\text{test})} \right)^2 \\ &= \frac{1}{n} \sum_{i=1}^{\text{ntest}} \left( \mathbf{y}_{i(\text{test})} - (\hat{\beta}_0 + \hat{\boldsymbol{\beta}} \mathbf{X}_{i(\text{test})}) \right)^2 \end{aligned}$$

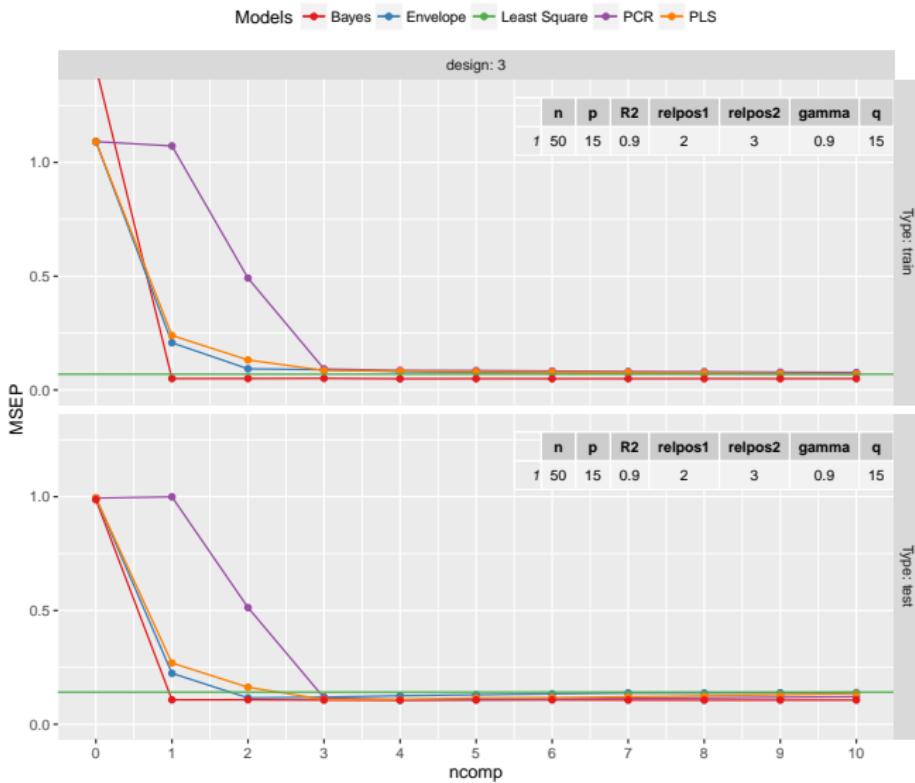
# Analysis Results



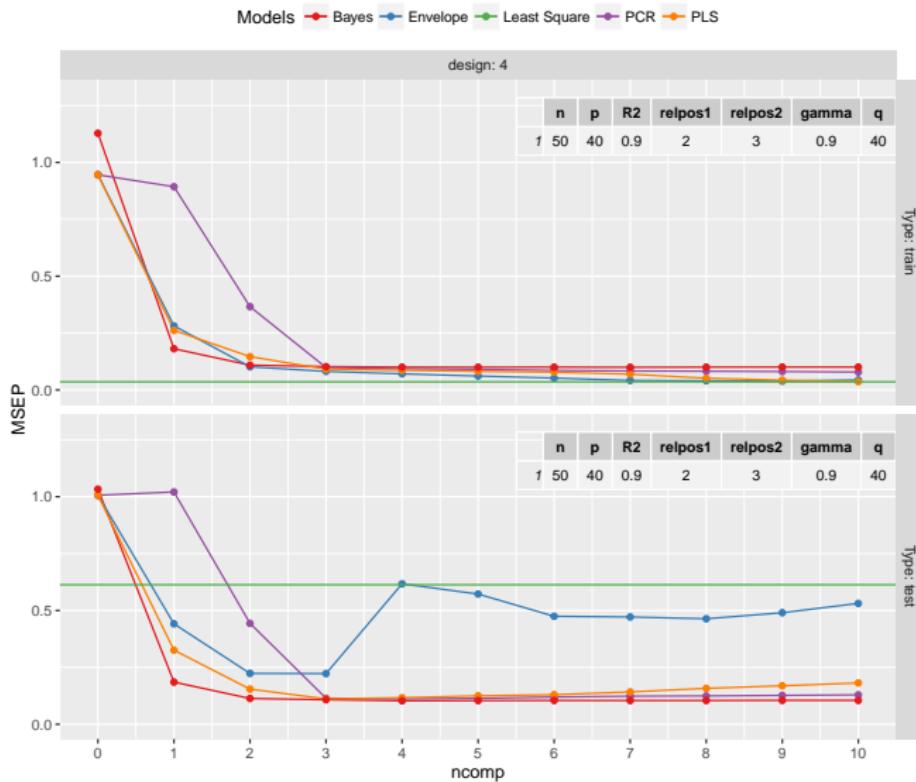
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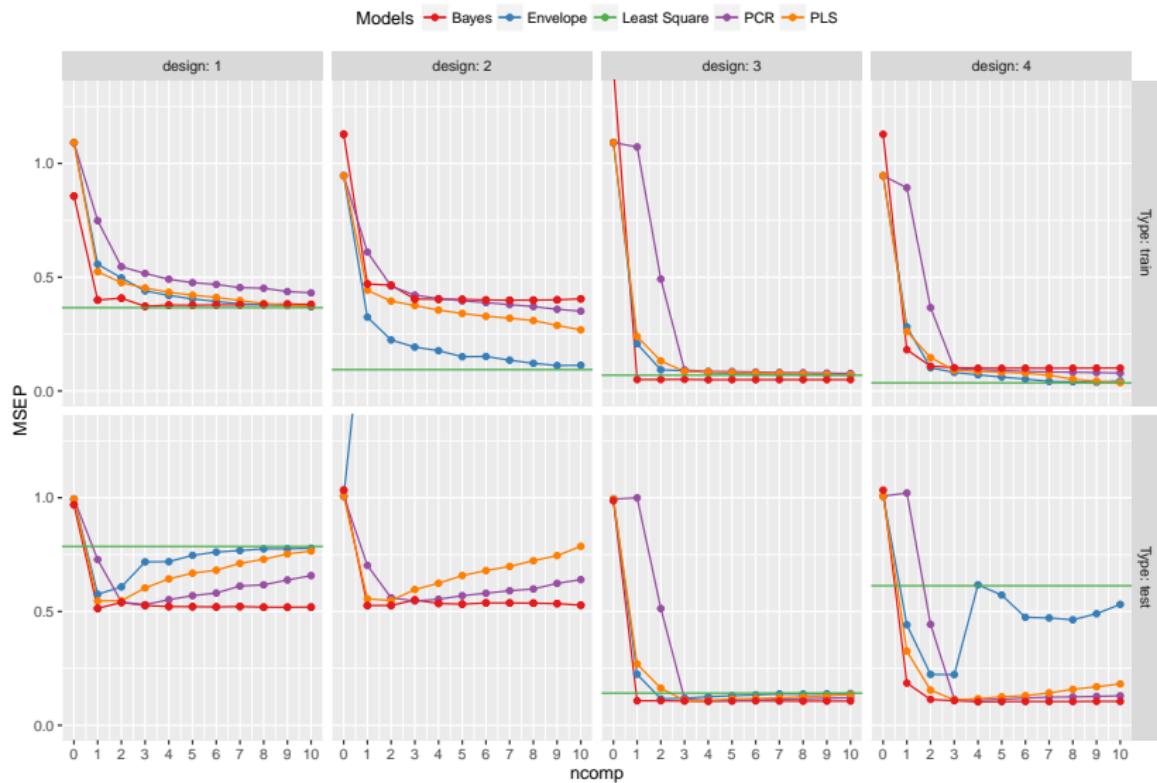


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- The computation regarding BayesPLS is intensive which will not be fisible in case of wide dataset (very common in genomic data)
- All the models are performing better than the least square solution

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TAKK HVALA  
Eυχαριστώ 감사합니다  
GRAZZII DANKE  
Rахмет kiiatos  
ARIGATO  
Suwun  
ধন্যবাদ  
MERCI  
ありがとう DANKE  
Благодарам  
grazie  
спасибо  
TAKK ASANTE  
多謝 SALAMAT  
gracias

teşekkür ederim SUWUN  
TAKK hvala Salamat  
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