Modeling Phenotype Products through Pre-Computed Summary Statistics

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IGES 30th Annual Meeting, General Session #3
October 15, 2021

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Introduction

A Question

What do we need to consider when we work with large biobank data?

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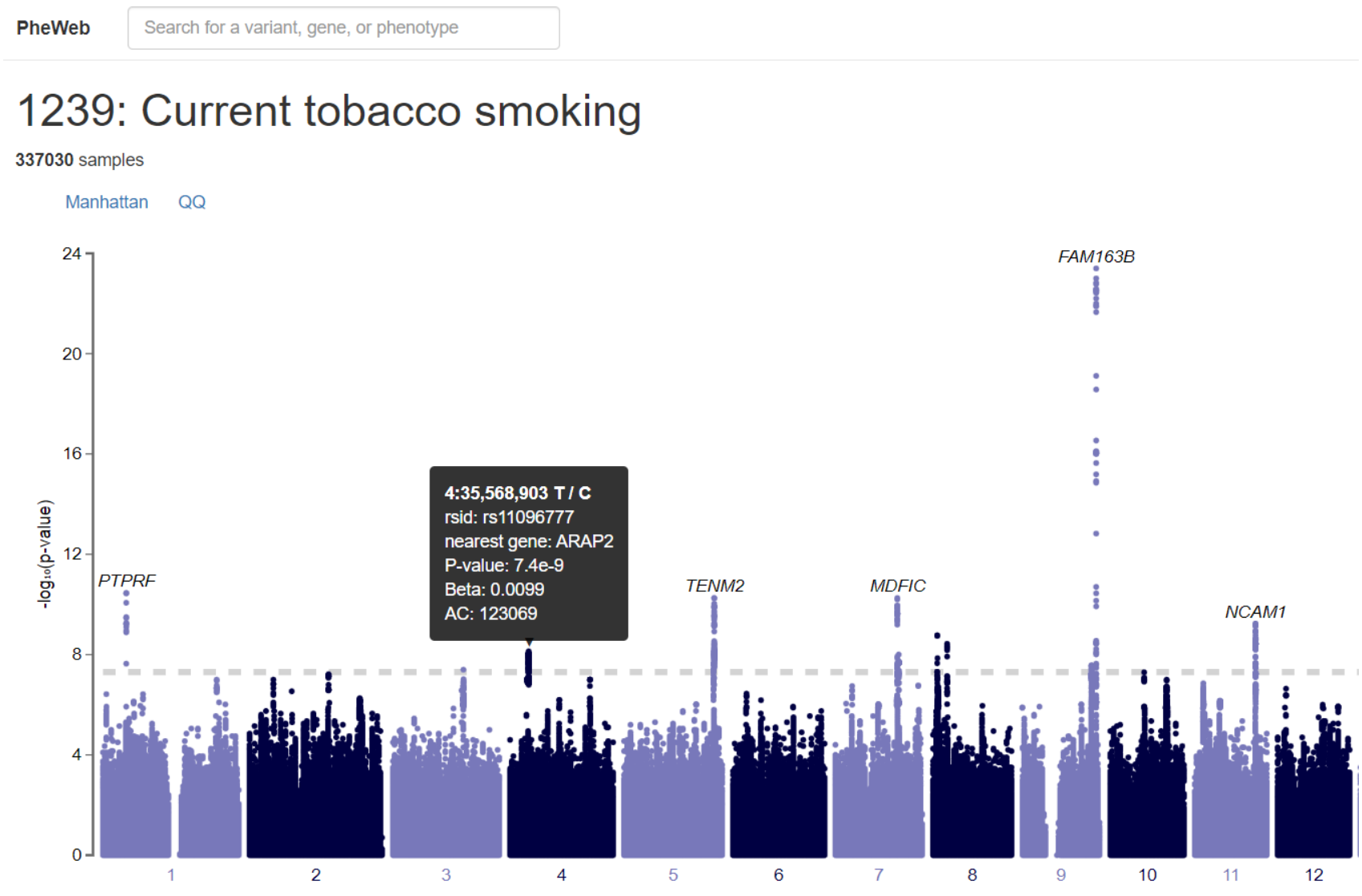
What do we need to consider when we work with large biobank data?

- Data privacy and security
- Data access and availability
- Computational costs

Introduction



Introduction

Key Idea

How can we leverage pre-computed summary statistics (PCSS) from biobanks to estimate statistical models fit using individual participant data (IPD)?

Existing Methods:

- Multi-trait association tests (Ray & Boehnke, 2018; Dutta et al., 2019; Guo & Wu, 2019)
- Linear combinations of phenotypes (Gasdaska et al., 2019; Wolf et al., 2020)

Introduction

Goal

Approximate linear models for products of phenotypes of the form:

$$\prod_{k=1}^{m} \boldsymbol{y}_k = \boldsymbol{X}\beta + \epsilon$$

using PCSS with flexible choice of covariates.

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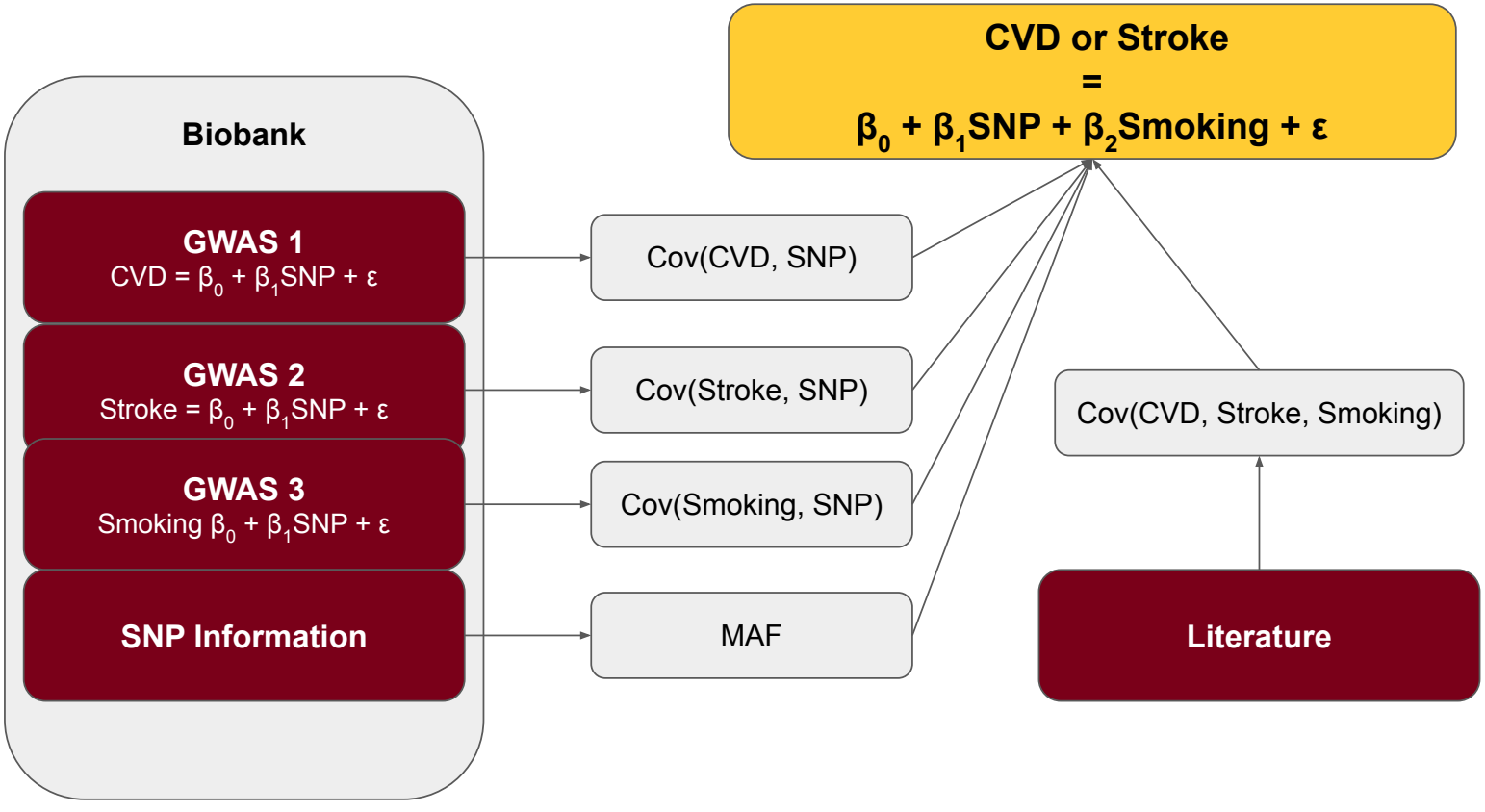
Why Products?

- Ratios of phenotypes
- Logical combinations of phenotypes

$$y_{i1} \wedge y_{i2} = y_{i1} y_{i2},$$
$$y_{i1} \vee y_{i2} = 1 - (1 - y_{i1})(1 - y_{i2})$$

Introduction



CVD = Cardiovascular disease; Cov = Covariance

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Assumed PCSS

$$\underbrace{\begin{bmatrix} \sigma_{\boldsymbol{x}_1,\boldsymbol{x}_1} & \sigma_{\boldsymbol{x}_1,\boldsymbol{x}_2} & \cdots & \sigma_{\boldsymbol{x}_1,\boldsymbol{x}_p} \\ & \ddots & \ddots & \vdots \\ & & \ddots & \vdots \\ & & & \sigma_{\boldsymbol{x}_p,\boldsymbol{x}_p} \end{bmatrix}}_{p\times p} \qquad \underbrace{\begin{bmatrix} \sigma_{\boldsymbol{x}_1,\boldsymbol{y}_1} & \sigma_{\boldsymbol{x}_1,\boldsymbol{y}_2} & \cdots & \sigma_{\boldsymbol{x}_1,\boldsymbol{y}_m} \\ \sigma_{\boldsymbol{x}_2,\boldsymbol{y}_1} & \ddots & & \vdots \\ \vdots & & \ddots & \vdots \\ \sigma_{\boldsymbol{x}_p,\boldsymbol{y}_m} & \cdots & \cdots & \sigma_{\boldsymbol{x}_p,\boldsymbol{y}_m} \end{bmatrix}}_{p\times m}$$

$$\underbrace{\begin{bmatrix} \bar{x}_1 & \bar{x}_2 & \cdots & \bar{x}_p \end{bmatrix}}_{1\times p} \qquad \underbrace{\begin{bmatrix} \sigma_{\boldsymbol{y}_1,\boldsymbol{y}_1} & \sigma_{\boldsymbol{y}_1,\boldsymbol{y}_2} & \cdots & \sigma_{\boldsymbol{y}_1,\boldsymbol{y}_m} \\ & \ddots & \ddots & \vdots \\ & & \ddots & \vdots \\ & & & \sigma_{\boldsymbol{y}_m,\boldsymbol{y}_m} \end{bmatrix}}_{m\times m}$$

$$\underbrace{\begin{bmatrix} \bar{y}_1 & \bar{y}_2 & \cdots & \bar{y}_m \end{bmatrix}}_{1\times m}$$

Regression with PCSS

Theorem

For the regression model $\boldsymbol{y} = \boldsymbol{X}\beta + \boldsymbol{\epsilon}$, *with* $\epsilon_i \overset{iid}{\sim} N(0, \sigma^2)$, *the ordinary least squares estimate for* β *is*

$$\hat{\beta} = (\boldsymbol{X}'\boldsymbol{X})^{-1}\boldsymbol{X}'\boldsymbol{y}$$

This can be computed via PCSS using the facts that:

$$\boldsymbol{X}'\boldsymbol{X} = (n-1)S(\boldsymbol{X}) + n\bar{\boldsymbol{x}}\bar{\boldsymbol{x}}' \tag{1}$$

$$\boldsymbol{X}'\boldsymbol{y} = (n-1)(s_{y,x_1}, \ldots, s_{y,x_p})' + n\bar{y}\bar{\boldsymbol{x}} \tag{2}$$

(Wolf et al., 2020)

Regression with PCSS

Theorem

The estimated variance of $\hat{\beta}$ is[*]

$$\widehat{\text{Var}}(\hat{\beta}) = \hat{\sigma}^2(\boldsymbol{X}'\boldsymbol{X})^{-1}$$

This can be calculated via PCSS using previous equalities and the fact that:

$$\hat{\sigma}^2 = [(n-1)s_y^2 + n\bar{y}^2 - \hat{\beta}'\boldsymbol{X}'\boldsymbol{y}]/(n-p) \tag{3}$$

(Wolf et al., 2020)

Modeling Phenotype Products

To approximate the covariance between $\boldsymbol{x}_j$ and the product $\boldsymbol{w} = \boldsymbol{y}_1 \boldsymbol{y}_2$ we estimate the conditional mean of $\boldsymbol{w}$ given $\boldsymbol{x}_j$ as

$$g(w|x) = g(y_1|x)g(y_2|x) + h(y_1, y_2|x), \tag{4}$$

which gives the covariance estimate

$$s_{x_j,w} \approx \sum_{x \in \mathcal{S}_j} f_j(x)(x - \bar{x}_j)g(w|x) \tag{5}$$

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Simulation Studies

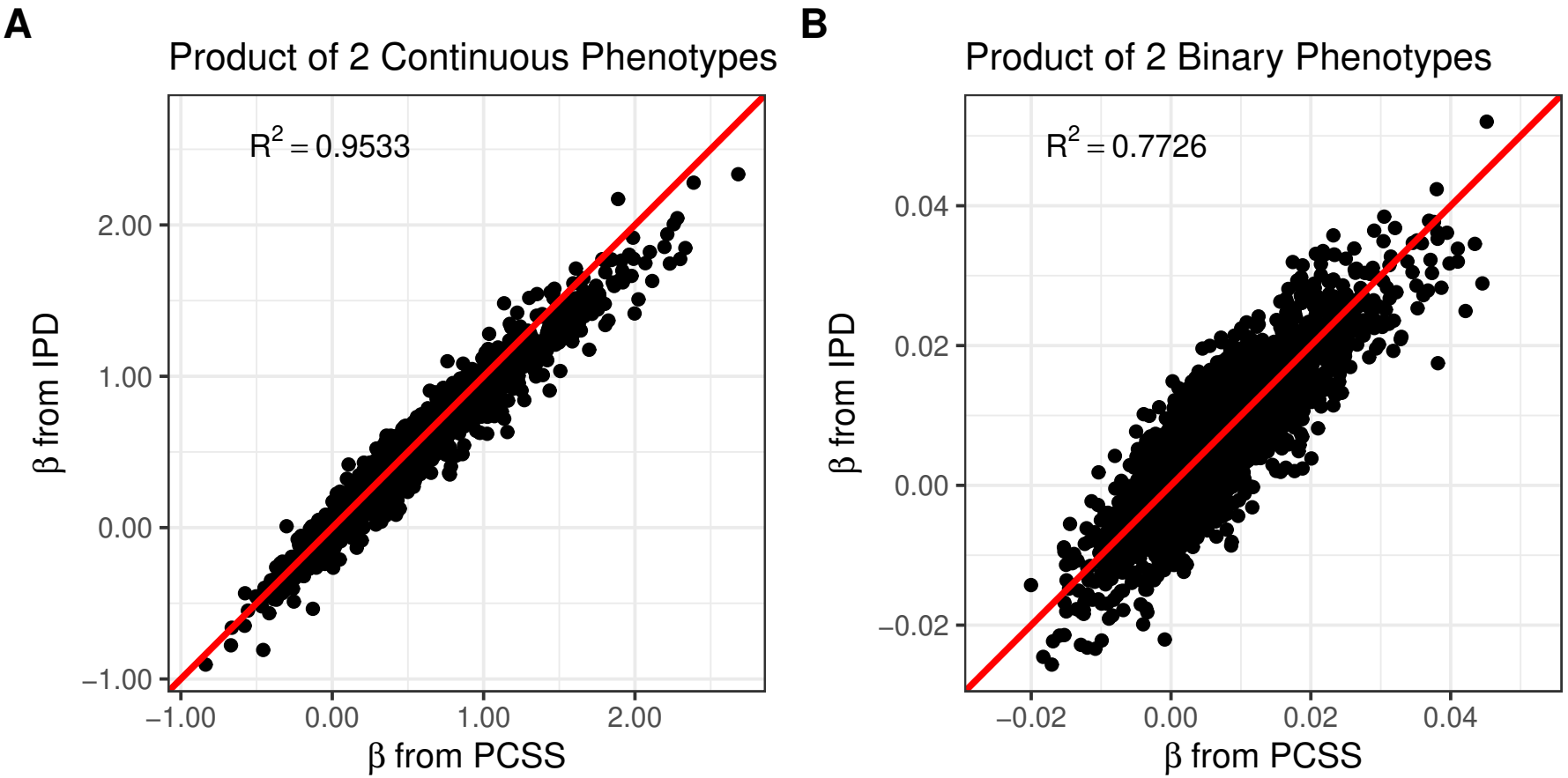
We simulated data through the model:

$$u(y_{ik}) = \beta_{k0} + \sum_{j=1}^{3} x_{ij}\beta_{kj} + \epsilon_{ik}$$

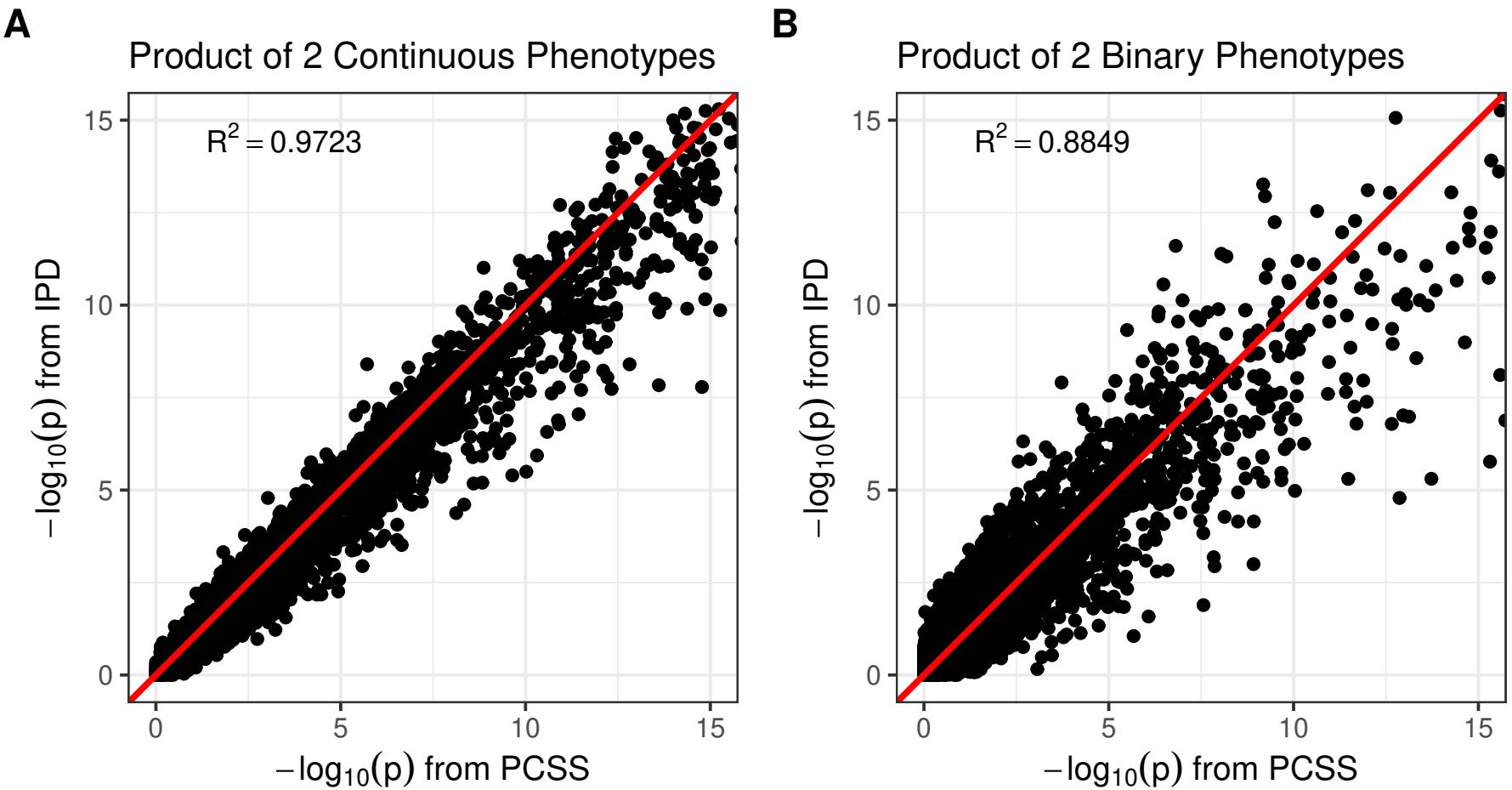
where

- $u(y_{ik}) = y_{ik}$ or $\text{logit}(\Pr(Y_{ik} = 1))$
- $\boldsymbol{x}_1 =$ SNP's minor allele counts
- $\boldsymbol{x}_2 =$ continuous covariate
- $\boldsymbol{x}_3 =$ binary covariate

Simulation Study Estimating β



Simulation Study Estimating p-values



Real Data Analysis

Fatty acids and conversion ratios

- Fatty acids are biomarkers of various cardiometabolic and cognitive health outcomes
- Conversion ratios illustrate how fatty acids are converted from one fatty acid to the next

Real Data Analysis

Framingham Heart Study (Mailman et al., 2007)

- 12 fatty acid conversion ratios
- 362,330 SNPs
- 4,347,960 models: FA Ratio $\sim$ SNP $+$ age $+$ sex

Real Data Analysis

Framingham Heart Study (Mailman et al., 2007)

- 12 fatty acid conversion ratios
- 362,330 SNPs
- 4,347,960 models: FA Ratio $\sim$ SNP $+$ age $+$ sex
- Disagreement rate of $10/(4.3 \times 10^6)$
- Of the 10 disagreements:
  - 4 where PCSS failed to reject when IPD rejected H_0,
  - 6 where PCSS rejected when IPD failed to reject

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Discussion

Takeaway

We can approximate linear models for products and logical combinations of phenotypes with a **flexible choice of covariates** using only readily available pre-computed summary statistics.

Discussion

Limitations and Future Work

- Assessing the compounding of errors when modeling the product of ≥ 4 phenotypes
- Measuring sensitivity to missing data and other assumption violations
- Assumes access to certain PCSS
- Accounting for related individuals through kinship matrices

Acknowledgements

This work was made possible by the generous support of the NIH through grant 2R15HG006915-03.

I would also like to thank the wonderful collaborators who have contributed to this research:

- Dr. Nathan Tintle
- Jason Westra
- Martha Barnard

Thank you!

Slides:	`https://bit.ly/IGESProduct`
R Package:	`pcsstools`
Twitter:	`@_jackmwolf`
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