

# SNP effects from ssGBLUP using BLUPF90 (postGSf90)

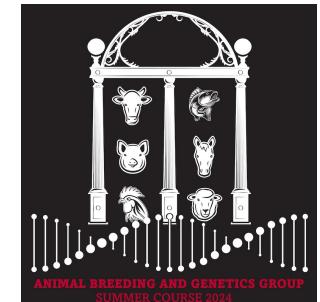
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BLUPF90 TEAM – 05/2024



UNIVERSITY OF  
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College of Agricultural &  
Environmental Sciences

*Animal Breeding and  
Genetics Group*



# Equivalence between GBLUP and SNP-BLUP

GBLUP

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{W} \\ \mathbf{W}'\mathbf{X} & \mathbf{W}'\mathbf{W} + \mathbf{G}^{-1}\lambda_1 \end{bmatrix} \begin{bmatrix} \hat{\beta} \\ \hat{u} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{W}'\mathbf{y} \end{bmatrix}$$



GEBV

SNP-BLUP (Ridge Regression)

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{Z} \\ \mathbf{Z}'\mathbf{X} & \mathbf{Z}'\mathbf{Z} + \mathbf{I}\lambda_2 \end{bmatrix} \begin{bmatrix} \hat{\beta} \\ \hat{a} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{Z}'\mathbf{y} \end{bmatrix}$$



SNP effects

$$\text{Var}(\mathbf{u}) = ?$$

$$\text{Var}(\mathbf{u}) = \mathbf{G}\sigma_u^2$$

$$\mathbf{u} = \mathbf{Z}\mathbf{a}$$

$$\text{Var}(\mathbf{u}) = ?$$

$$\text{Var}(\mathbf{u}) = \mathbf{G}\sigma_u^2$$

# Are GBLUP and SNP-BLUP equivalent?

- Assumption of GBLUP:  $\text{Var}(\mathbf{u}) = \mathbf{G}\sigma_u^2$
- In SNP-BLUP:  $\mathbf{u} = \mathbf{Z}\mathbf{a}$

$$\mathbf{u} = \mathbf{Z}\mathbf{a}$$

$$\text{Var}(\mathbf{u}) = \text{Var}(\mathbf{Z}\mathbf{a})$$

$$\text{Var}(\mathbf{u}) = \mathbf{Z} \text{Var}(\mathbf{a}) \mathbf{Z}'$$

$$\text{Var}(\mathbf{u}) = \mathbf{Z}\mathbf{Z}'\sigma_a^2$$

$$\sigma_a^2 = \frac{\sigma_u^2}{2 \sum_{i=1}^{SNP} p_i(1 - p_i)}$$

$$\text{Var}(\mathbf{u}) = \mathbf{Z}\mathbf{Z}' \frac{\sigma_u^2}{2 \sum_{i=1}^{SNP} p_i(1 - p_i)}$$

$$\text{Var}(\mathbf{u}) = \frac{\mathbf{Z}\mathbf{Z}'}{2 \sum_{i=1}^{SNP} p_i(1 - p_i)} \sigma_u^2$$

$$\mathbf{G} = \frac{\mathbf{Z}\mathbf{Z}'}{2 \sum_{i=1}^{SNP} p_i(1 - p_i)}$$

$$\text{Var}(\mathbf{u}) = \mathbf{G}\sigma_u^2$$



Genomic  
relationship matrix  
VanRaden (2008)

GBLUP assumption!!!

# GBLUP and SNP-BLUP are equivalent!

If we can get  $\mathbf{u}$  ( $\mathbf{u} = \mathbf{Z}\mathbf{a}$ ) from SNP-BLUP, we can get  $\mathbf{a}$  from GBLUP!



*Review*

## Single-Step Genomic Evaluations from Theory to Practice: Using SNP Chips and Sequence Data in BLUPF90

Daniela Lourenco <sup>1,\*</sup>, Andres Legarra <sup>2</sup>, Shogo Tsuruta <sup>1</sup>, Yutaka Masuda <sup>1</sup>, Ignacio Aguilar <sup>3</sup>, and Ignacy Misztal <sup>1</sup>

<https://www.mdpi.com/2073-4425/11/7/790>

Pages 11-12

# **ssGBLUP** and **ssSNP-BLUP** are also equivalent!

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{W} \\ \mathbf{W}'\mathbf{X} & \mathbf{W}'\mathbf{W} + \mathbf{H}^{-1} \frac{\sigma_e^2}{\sigma_u^2} \end{bmatrix} \begin{bmatrix} \hat{\boldsymbol{\beta}} \\ \hat{\mathbf{u}} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{W}'\mathbf{y} \end{bmatrix}$$

## **ssGBLUP**

Misztal et al. (2009)  
Legarra et al. (2009)  
Aguilar et al. (2010)  
Christensen & Lund (2010)

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{Z}\mathbf{M} & \mathbf{X}'_n\mathbf{Z}_n \\ \mathbf{M}'\mathbf{Z}'\mathbf{X} & \mathbf{M}'\mathbf{Z}'\mathbf{Z}\mathbf{M} + \mathbf{I} \frac{\sigma_e^2}{\sigma_\alpha^2} & \mathbf{M}'_n\mathbf{Z}'_n\mathbf{Z}_n \\ \mathbf{Z}'_n\mathbf{X}_n & \mathbf{Z}'_n\mathbf{Z}_n\mathbf{M}_n & \mathbf{Z}'_n\mathbf{Z}_n + \mathbf{A}^{nn} \frac{\sigma_e^2}{\sigma_g^2} \end{bmatrix} \begin{bmatrix} \hat{\boldsymbol{\beta}} \\ \hat{\boldsymbol{\alpha}} \\ \hat{\boldsymbol{\epsilon}} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{M}'\mathbf{Z}'\mathbf{y} \\ \mathbf{Z}'_n\mathbf{y}_n \end{bmatrix}$$

## **ssSNPBLUP or ssBR**

Fernando et al. (2014)  
Liu et al. (2014)  
Mantysaari & Strandén (2016)

 J. Dairy Sci. 101:10082–10088  
<https://doi.org/10.3168/jds.2018-14913>  
© 2018, The Authors. Published by FASS Inc. and Elsevier Inc. on behalf of the American Dairy Science Association®.  
This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

**Short communication: Genomic prediction using different single-step methods in the Finnish red dairy cattle population**

H. Gao,\*† M. Koivula,‡ J. Jensen,\* I. Strandén,‡ P. Madsen,\* T. Pitkänen,‡ G. P. Aamand,†  
and E. A. Mantysaari†

\*Center for Quantitative Genetics and Genomics, Department of Molecular Biology and Genetics, Aarhus University, DK-8830 Tjele, Denmark

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‡Natural Resources Institute Finland (Luke), FIN-31600 Jokioinen, Finland

We confirmed that regular ssGBLUP and ssBR with an extra polygenic effect led to the same predictions.

# SNP effects in ssGBLUP

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{W} \\ \mathbf{W}'\mathbf{X} & \mathbf{W}'\mathbf{W} + \mathbf{H}^{-1}\lambda_1 \end{bmatrix} \begin{bmatrix} \hat{\boldsymbol{\beta}} \\ \hat{\mathbf{u}} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{W}'\mathbf{y} \end{bmatrix}$$

$$\hat{\mathbf{a}} = \alpha b \frac{1}{2\sum p_i(1-p_i)} \mathbf{Z}' \mathbf{G}^{-1} \hat{\mathbf{u}}$$

↗ Matrix of SNP content  
↘ Genomic relationship matrix

$\alpha$  = blending parameter for  $\mathbf{G}$

$$b = 1 - \frac{\lambda}{2}$$

$$\lambda = \frac{1}{n^2} \left( \sum_i \sum_j \mathbf{A}_{22ij} - \sum_i \sum_j \mathbf{G}_{ij} \right)$$

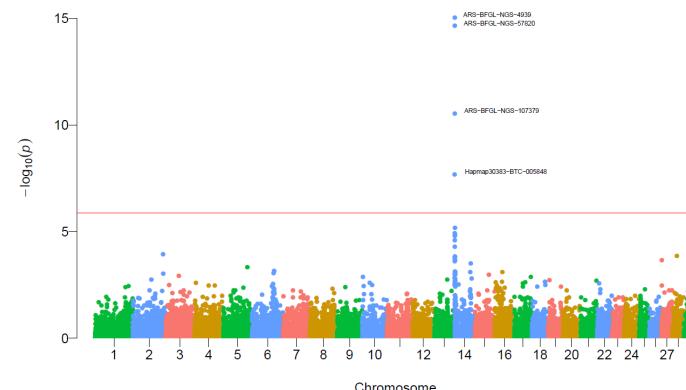
# What can we do with SNP effects?

- 1) Predictions for animals not included in the evaluation

Indirect Predictions

Indirect Genomic Predictions

- 2) Genome-Wide Association Studies (GWAS)



# 1) Indirect Predictions

- Interim evaluations
  - Between official runs
- Not all genotyped animals are in the evaluations
  - Animals with incomplete pedigree increase bias and lower  $R^2$
- Commercial products
  - e.g., GeneMax -> genomic testing for non-registered animals

# 1) Indirect Predictions

$$\begin{bmatrix} \mathbf{X}'\mathbf{X} & \mathbf{X}'\mathbf{W} \\ \mathbf{W}'\mathbf{X} & \mathbf{W}'\mathbf{W} + \mathbf{H}^{-1}\lambda_1 \end{bmatrix} \begin{bmatrix} \widehat{\boldsymbol{\beta}} \\ \widehat{\mathbf{u}} \end{bmatrix} = \begin{bmatrix} \mathbf{X}'\mathbf{y} \\ \mathbf{W}'\mathbf{y} \end{bmatrix} \quad \longrightarrow \quad \widehat{\mathbf{a}} = \alpha b \frac{1}{2\sum p_i(1-p_i)} \mathbf{Z}'\mathbf{G}^{-1} \widehat{\mathbf{u}}$$

Indirect Prediction:  $\mathbf{IP} = \mathbf{u}_m^* = \mathbf{Z}\widehat{\mathbf{a}}$

# 1) Indirect Predictions

Indirect Prediction:  $\mathbf{u}_m^* = \mathbf{Z}\hat{\mathbf{a}}$

→ Fine if comparing among animals with IP

- Not fine if comparing IP with GEBV ( $\hat{\mathbf{u}}$ ) from the main evaluation
  - Need to put IP in the pedigree scale

$$\mathbf{u}_m = \hat{\boldsymbol{\mu}} + \mathbf{u}_m^*$$

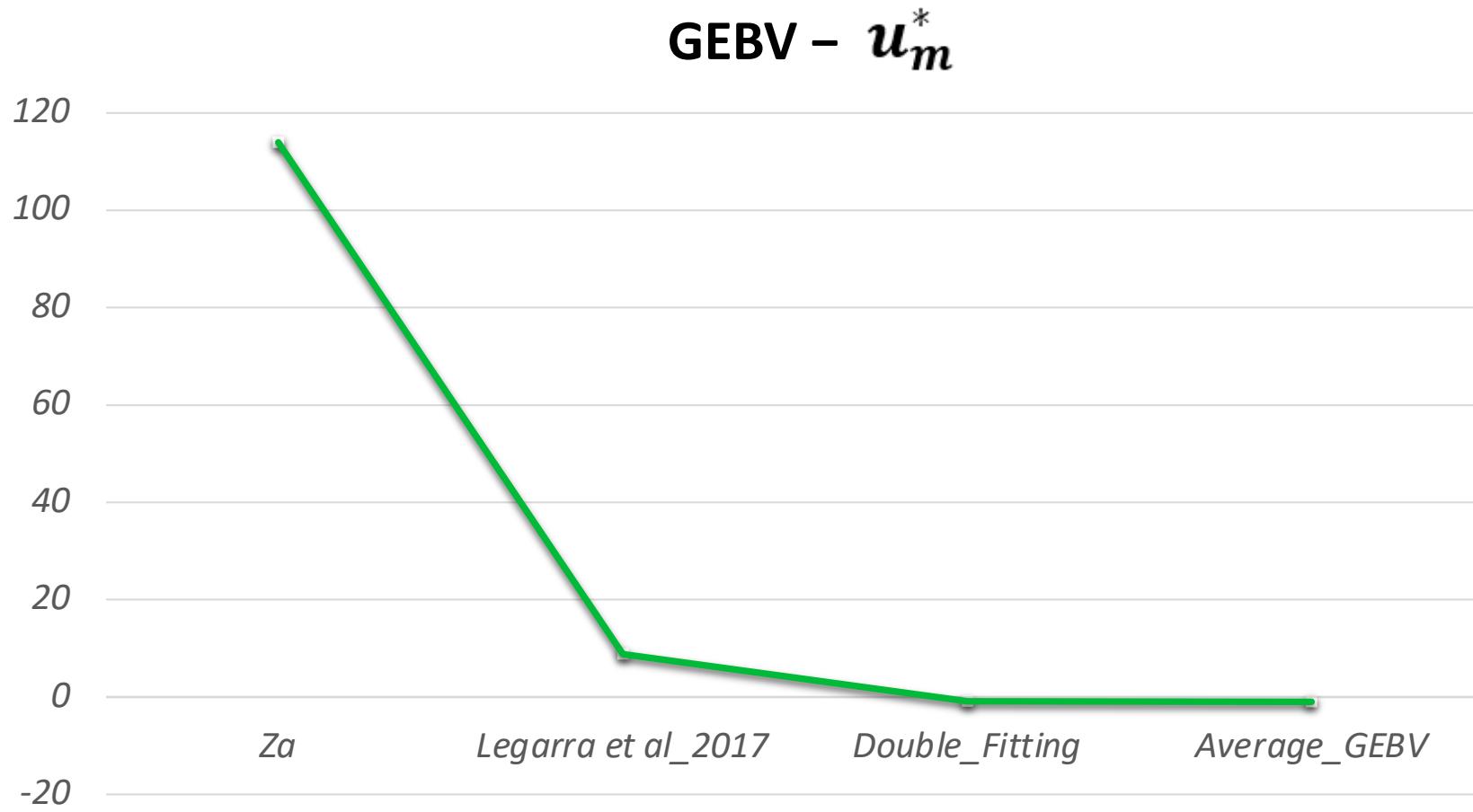
→

$$\hat{\boldsymbol{\mu}} = \alpha\lambda\mathbf{1}'\mathbf{G}^{-1}\hat{\mathbf{u}}$$

$\alpha$  = blending parameter for  $\mathbf{G}$

$$\lambda = \frac{1}{n^2} \left( \sum_i \sum_j \mathbf{A}_{22ij} - \sum_i \sum_j \mathbf{G}_{ij} \right)$$

# 1) Indirect Predictions



# How to compute Indirect predictions

1) Pedigree + phenotypes + genotypes

2) **renumf90**

3) preGSf90 to save clean files

4) **blupf90+** (save the clean files)

- Good practice to save time: OPTION saveGInverse + OPTION saveA22Inverse

5) **postGSf90** (with clean files)

- BLUPF90 family software to compute SNP effects (+more)
- Same parameter file as blupf90+
- Good practice to save time: OPTION readGInverse + OPTION readA22Inverse

# Output from postGSf90

snp\_sol

<http://nce.ads.uga.edu/wiki/doku.php?id=readme.pregsf90>

contains solutions of SNP and weights

- 1: trait
- 2: effect
- 3: SNP
- 4: Chromosome
- 5: Position
- 6: SNP solution
- 7: weight

snp\_pred

- 1<sup>st</sup> line: model, tuning, blending information
- 2<sup>nd</sup> line: Trait/effect info
- AF in 10 columns
- mu\_hat, var\_mu\_hat
- SNP effects

# How to compute Indirect Predictions

## 6) predf90

- Have to provide a SNP file for the new genotyped animals to receive IP
  - same SNP as in the clean file

```
predf90 --snpfile newgen.txt --use_mu_hat
```

- The last statement adds the base, so that we have:  $u_m = \hat{\mu} + u_m^*$

# Output from predf90

## SNP\_predictions

Animal ID	SNP call rate	Indirect Predictions
-----------	---------------	----------------------

UGA50014	1.00	0.17414457
UGA50016	1.00	0.72332874E-01
UGA50042	1.00	1.0016705
UGA50058	1.00	0.17190497
UGA50060	1.00	0.98674759E-01
UGA50065	1.00	-0.60623702E-01
UGA50073	1.00	-0.17860851
UGA50077	1.00	-0.21597147
UGA50079	1.00	-0.69586390
UGA50084	1.00	1.0600574
UGA50085	1.00	-0.28602412
UGA50088	1.00	-0.12758011

**predf90** can also compute accuracy of indirect predictions

```
OPTION snp_p_value      #in blupf90+
OPTION snp_var          #in postGSf90
--acc                  #in predf90
```

Garcia et al. *Genetics Selection Evolution* (2022) 54:66  
<https://doi.org/10.1186/s12711-022-00752-4>



RESEARCH ARTICLE

Open Access



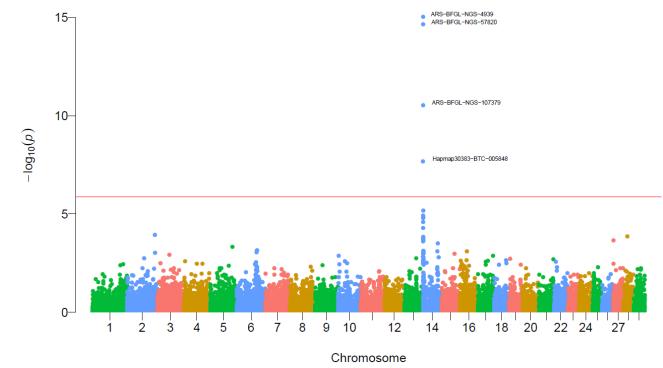
Theoretical accuracy for indirect predictions  
based on SNP effects from single-step GBLUP

Andre Garcia<sup>1\*</sup>, Ignacio Aguilar<sup>2</sup>, Andres Legarra<sup>3</sup>, Shogo Tsuruta<sup>1</sup>, Ignacy Misztal<sup>1</sup> and Daniela Lourenco<sup>1</sup>

## 2) Genome-wide Association Studies

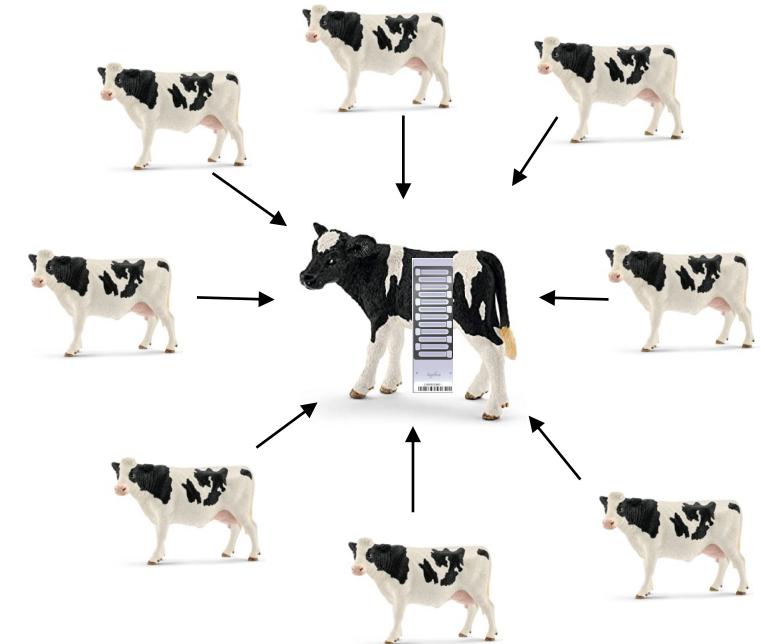
# Current standard for GWAS

- Single marker regression with  $\mathbf{G}$  to compensate for relationships
  - $\mathbf{y} = \mathbf{X}\mathbf{b} + \mathbf{z}_i\mathbf{a}_i + \mathbf{u} + \mathbf{e}$ 
    - $\mathbf{z}$ : gene content {0,1,2}
    - $\mathbf{a}$ : SNP effect
- Estimate SNP effects
- Get p-values as  $pval_i = 2 \left( 1 - \Phi \left( \left| \frac{\hat{a}_i}{sd(\hat{a}_i)} \right| \right) \right)$
- Apply Bonferroni to correct for multiple testing
- **Assumption: Genotyped individuals have phenotypes**



# GWAS in livestock populations

- Most animals are non-genotyped
- Animals may not have phenotypes
- Some traits are sex-limited
  - milk, fat, protein
- Single marker regression
  - Only genotyped animals with phenotypes
  - Deregressed EBV
- Need a method that fits the livestock data
  - ssGWAS



# Single-step GWAS (historical)

SNP  
effects

GEBVs

VanRaden 2008  
Stranden and Garrick 2009  
Wang et al. 2012

$$\hat{\mathbf{a}} = \alpha b \frac{1}{2\sum p_i(1-p_i)} \mathbf{Z}' \mathbf{G}^{-1} \hat{\mathbf{u}}$$

a) Quadratic SNP variance (Falconer & Mackay, 1996)

$$d_i = \hat{a}_i^2 2p_i(1-p_i)$$

b) NonlinearA SNP variance (VanRaden, 2008)

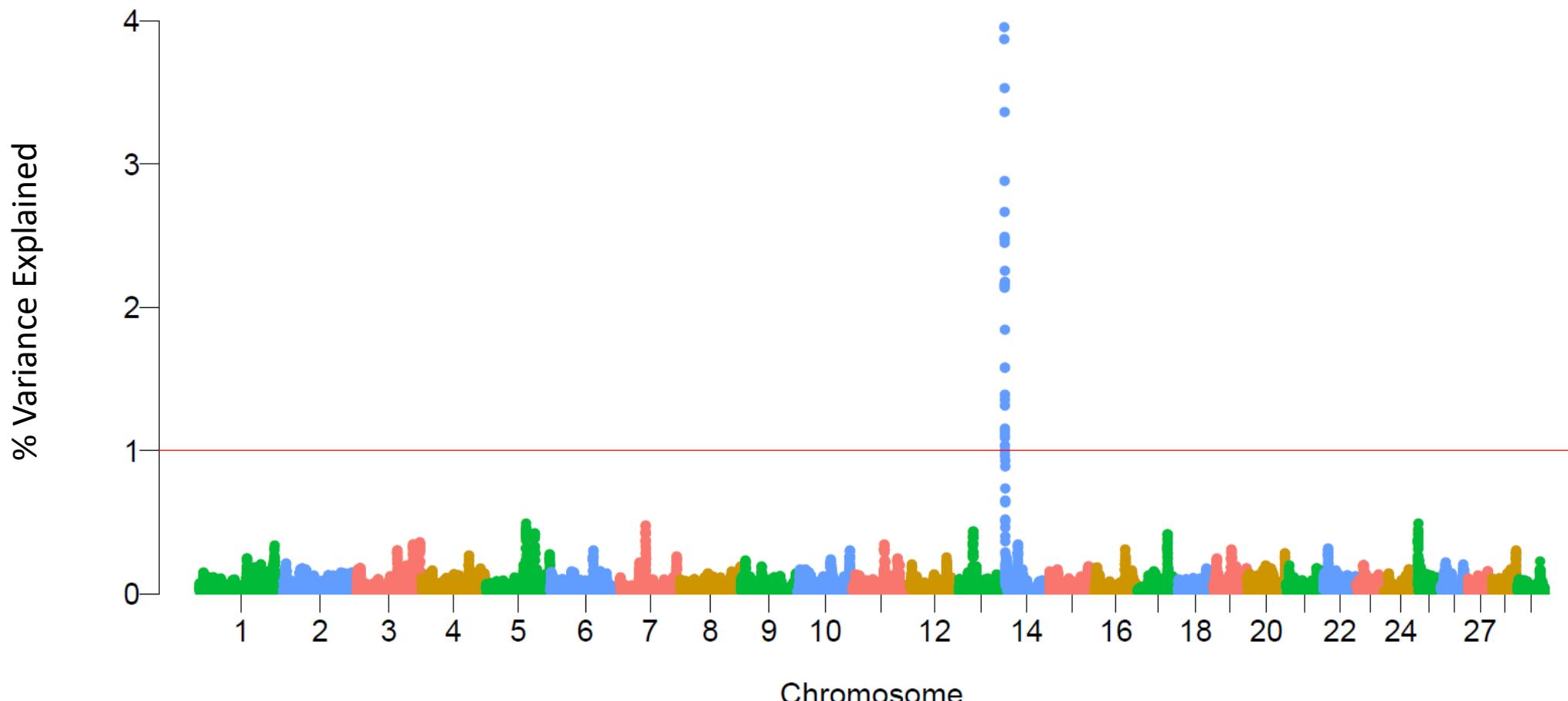
$$d_i = 1.125 \frac{|\hat{a}_i|}{sd(\hat{a})} - 2$$

# Single-step GWAS

Fat – US Holsteins

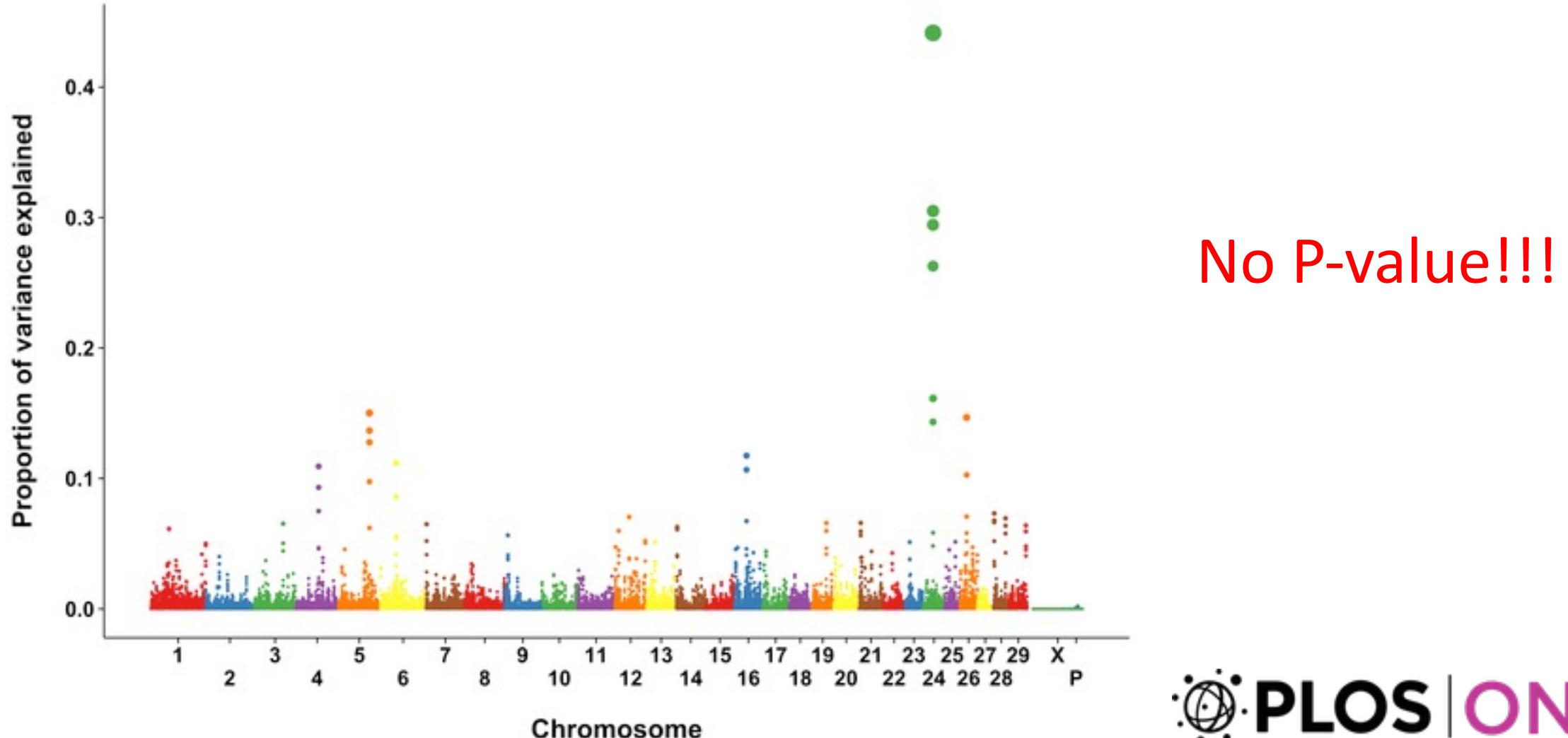
No P-value!!!

**Manhattan plot of Variances**



# Single-step GWAS

Figure 2. Proportion of SNP variance explained by 5-SNP moving windows for rectal temperature from a **single-step GBLUP** analysis



# Can we have p-values in ssGWAS?

Guadán Duarte et al. BMC Bioinformatics 2014, 15:246  
http://www.biomedcentral.com/1471-2105/15/246

METHODOLOGY ARTICLE



Open Access

Rapid screening for phenotype-genotype associations by linear transformations of genomic evaluations

Jose L Guadán Duarte<sup>1</sup>, Rodolfo JC Cantet<sup>1</sup>, Ronald O Bates<sup>2</sup>, Catherine W Ernst<sup>2</sup>, Nancy E Raney<sup>2</sup>  
and Juan P Steibel<sup>2,3\*</sup>

**Genome-Wide Association Analyses Based on Broadly Different Specifications for Prior Distributions, Genomic Windows, and Estimation Methods**

Chunyu Chen,<sup>1</sup> Juan P. Steibel, and Robert J. Tempelman  
Department of Animal Science, Michigan State University, East Lansing, Michigan 48824  
ORCID ID: 0000-0002-7833-6730 (R.J.T.)

Aguilar et al. *Genet Sel Evol* (2019) 51:28  
https://doi.org/10.1186/s12711-019-0469-3

SHORT COMMUNICATION

ANIMAL GENETICS Immunogenetics, Molecular Genetics and Functional Genomics



doi: 10.1111/age.12378

Meta-analysis of genome-wide association from genomic prediction models

Y. L. Bernal Rubio<sup>\*†</sup>, J. L. Guadán Duarte<sup>\*</sup>, R. O. Bates<sup>\*</sup>, C. W. Ernst<sup>\*</sup>, D. Nonneman<sup>‡</sup>,  
G. A. Rohrer<sup>‡</sup>, A. King<sup>‡</sup>, S. D. Shackelford<sup>‡</sup>, T. L. Wheeler<sup>‡</sup>, R. J. C. Cantet<sup>†§</sup> and J. P. Steibel<sup>\*¶</sup>



J. Dairy Sci. 101:3140–3154  
https://doi.org/10.3168/jds.2017-13364  
© American Dairy Science Association<sup>®</sup>, 2018.

Genome-wide association analyses based on a multiple-trait approach for modeling feed efficiency

Y. Lu,<sup>\*</sup> M. J. Vandehaar,<sup>\*</sup> D. M. Spurlock,<sup>†</sup> K. A. Weigel,<sup>‡</sup> L. E. Armentano,<sup>‡</sup> E. E. Connor,<sup>§</sup> M. Coffey,<sup>#</sup>  
R. F. Veerkamp,<sup>||</sup> Y. de Haas,<sup>||</sup> C. R. Staples,<sup>||</sup> Z. Wang,<sup>\*\*</sup> M. D. Hanigan,<sup>††</sup> and R. J. Tempelman<sup>\*¶</sup>



Open Access



Frequentist p-values for large-scale-single step genome-wide association, with an application to birth weight in American Angus cattle

Ignacio Aguilar<sup>1</sup>, Andres Legarra<sup>2\*</sup>, Fernando Cardoso<sup>3,4</sup>, Yutaka Masuda<sup>5</sup>, Daniela Lourenco<sup>5</sup>  
and Ignacy Misztal<sup>5</sup>

# P-values in ssGWAS

1) Factorize and Invert LHS of ssGBLUP with YAMS (Masuda et al., 2014)

2) Solve the MME for  $\begin{bmatrix} \hat{\beta} \\ \hat{u} \end{bmatrix}$  using the sparse Cholesky factor

**blupf90+**



Ignacio  
Aguilar

3) Extract coefficients for genotyped animals ( $\mathbf{C}^{u_2 u_2}$ ) from LHS<sup>-1</sup>

4) Obtain individual prediction error variance of SNP effects:

$$Var(\hat{a}_i) = ab \frac{1}{2\sum p_i(1-p_i)} \mathbf{z}'_i \mathbf{G}^{-1} (\mathbf{G}\sigma_u^2 - \mathbf{C}^{u_2 u_2}) \mathbf{G}^{-1} \mathbf{z}_i \frac{1}{2\sum p_i(1-p_i)} ab$$

(Gualdrón-Duarte et al., 2014)

**postGSf90**



Andres  
Legarra

5) Backsolve GEBV to SNP effects ( $\hat{a}$ ):  $\hat{a} = ab \frac{1}{2\sum p_i q_i} \mathbf{Z}' \mathbf{G}^{-1} \hat{u}$

6)  $p\text{-value}_i = 2 \left( 1 - \Phi \left( \left| \frac{\hat{a}_i}{sd(\hat{a}_i)} \right| \right) \right)$

$\Phi$  is the cumulative standard normal function



Yutaka  
Masuda

# How to run ssGWAS with p-values in BLUPF90

- After renumf90 and preGSf90 to save clean files:

- blupf90+ to estimate GEBV
  - OPTION SNP\_file `snp.dat_clean`
  - OPTION map\_file `mrkmap.txt_clean`
  - OPTION saveGInverse
  - OPTION saveA22Inverse
  - OPTION.snp\_p\_value
- postGSf90 to backsolve GEBV to SNP effect
  - OPTION SNP\_file `snp.dat_clean`
  - OPTION map\_file `mrkmap.txt_clean`
  - OPTION readGInverse
  - OPTION readA22Inverse
  - OPTION.snp\_p\_value
  - OPTION.windows\_variance X #if need variance explained by X SNP

# Output from postGSf90

chrsnp\_pval

contains data to create plot by GNUPLOT

- 1: trait
- 2: effect
- 3: -log10(p-value)
- 4: SNP
- 5: Chromosome
- 6: Position in bp

chrsnp

contains data to create plot by GNUPLOT

- 1: trait
- 2: effect
- 3: values of SNP effects to use in Manhattan plots →  $[\text{abs}(\text{SNP}_i)/\text{var}(\text{SNP})]$
- 4: SNP
- 5: Chromosome
- 6: Position

Pft1e2.gnuplot

Pft1e2.R

Sft1e2.gnuplot

Sft1e2.R

# Output from postGSf90

chr snp var

contains data to create plot by GNUPLOT

- 1: trait
- 2: effect
- 3: variance explained by  $n$  adjacents SNP
- 4: SNP
- 5: Chromosome
- 6: Position

Vft1e2.gnuplot

Vft1e2.R

# Output from postGSf90

snp\_sol

contains solutions of SNP and weights

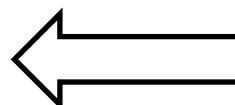
- 1: trait
- 2: effect
- 3: SNP
- 4: Chromosome
- 5: Position
- 6: SNP solution
- 7: weight

if `OPTION windows_variance` is used

- 8: variance explained by  $n$  adjacents SNP.

if `OPTION.snp_p_value` is used

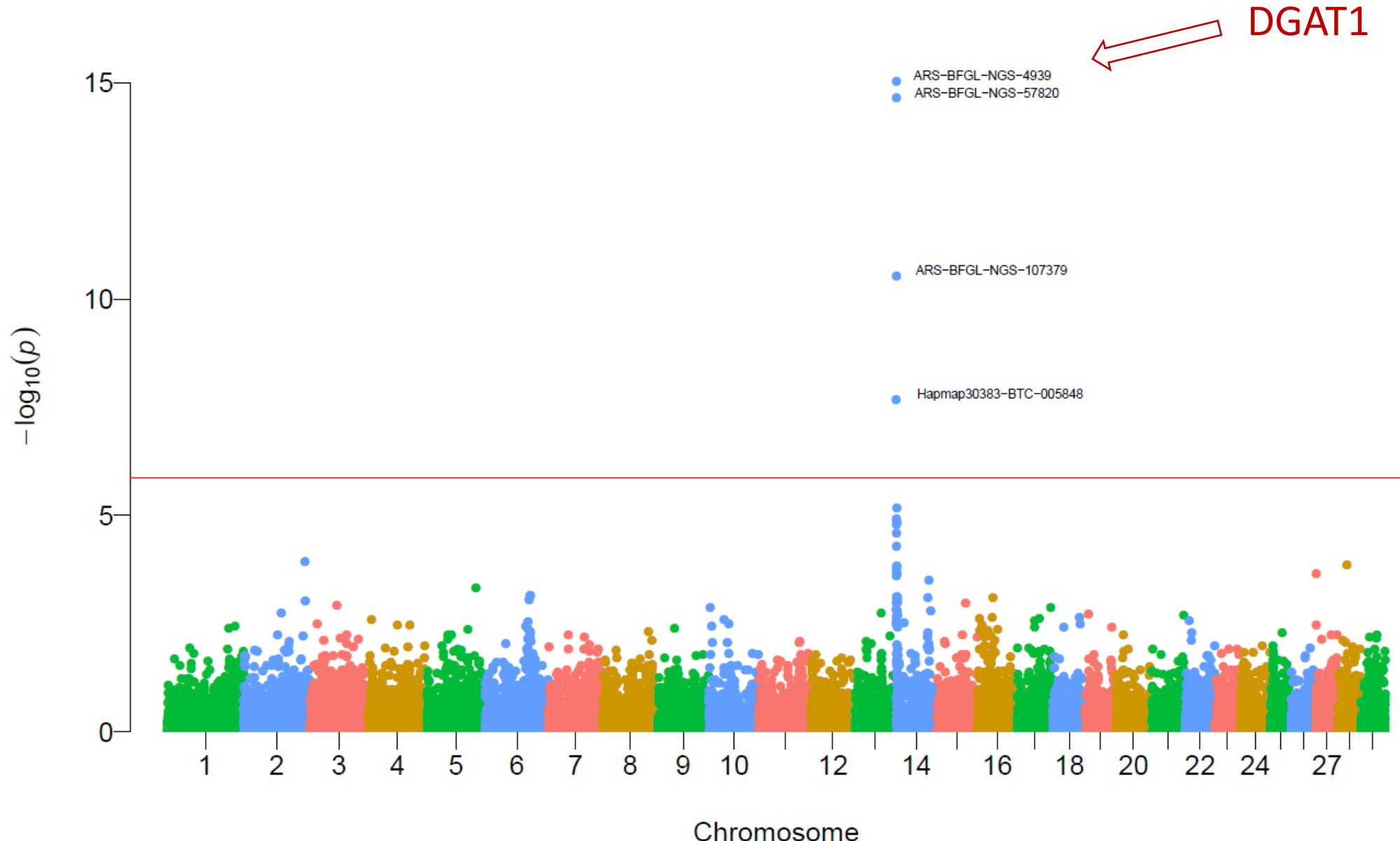
- 9: variance of the SNP solution (used to compute the p-value)



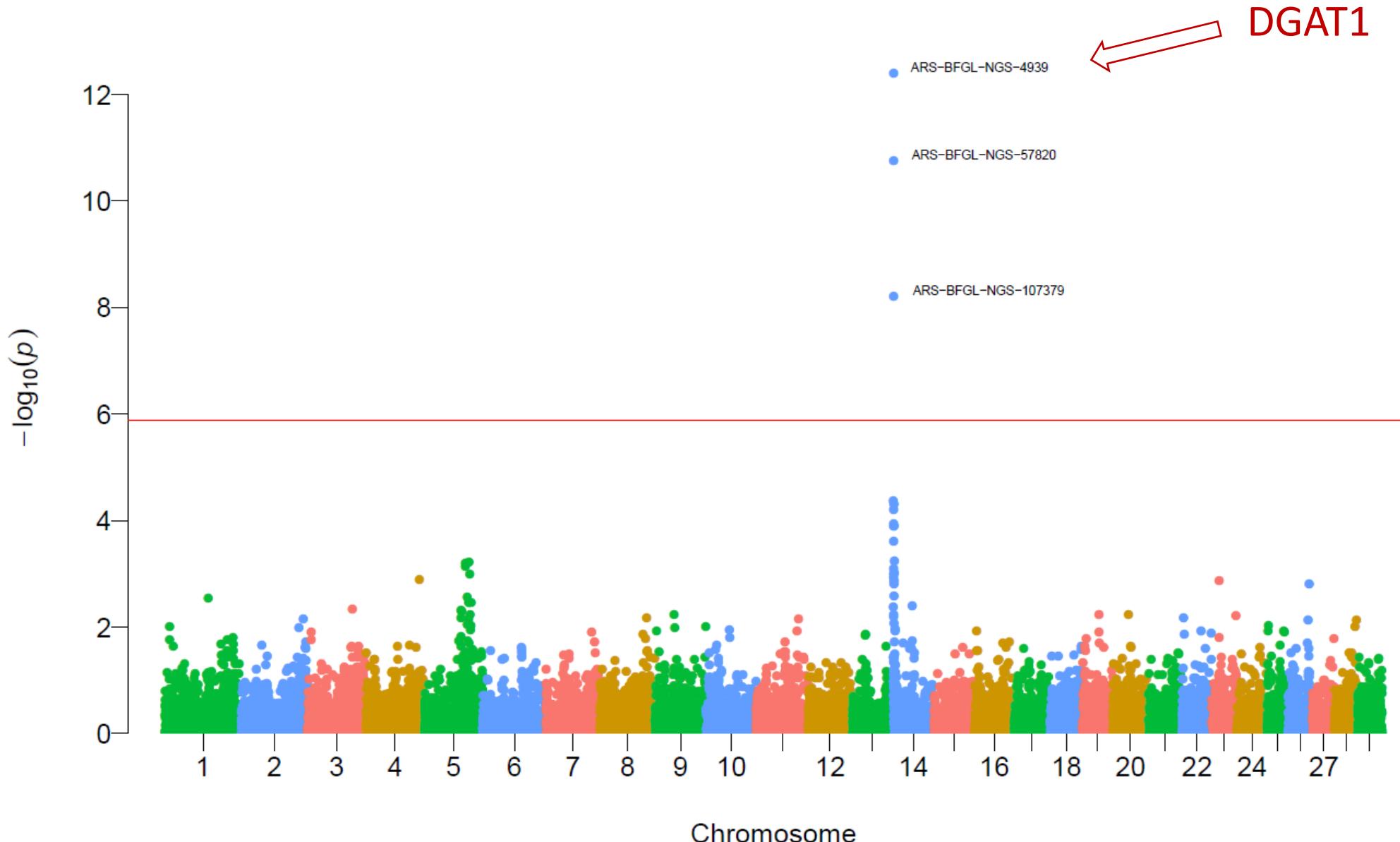
# P-values in ssGWAS for US Holsteins

- US HOL 2009 data: milk, fat, protein
- Single-trait models
  - 10k genotyped bulls
  - 752k records for 100k daughters
  - 303k animals in ped

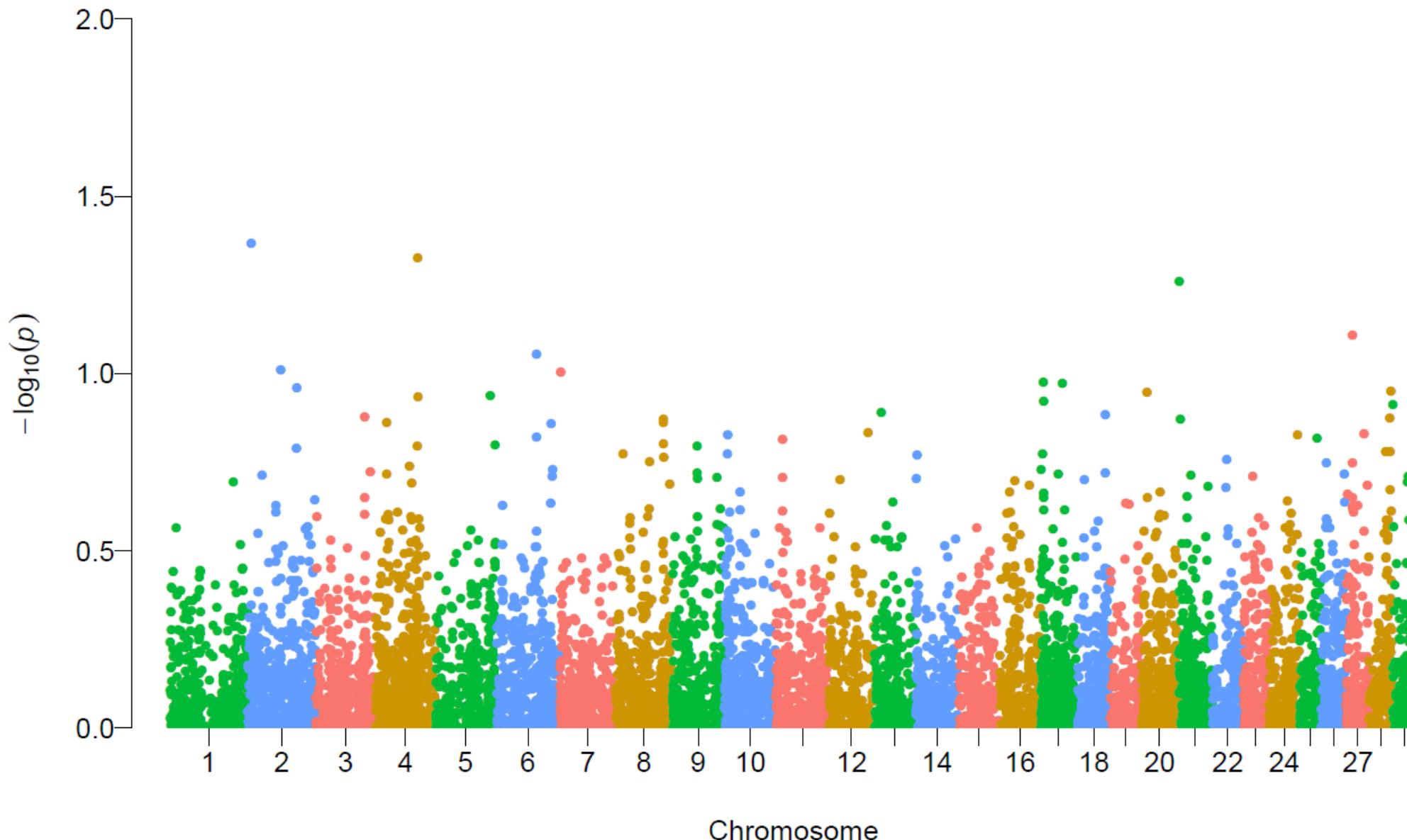
# P-values in ssGWAS - Milk



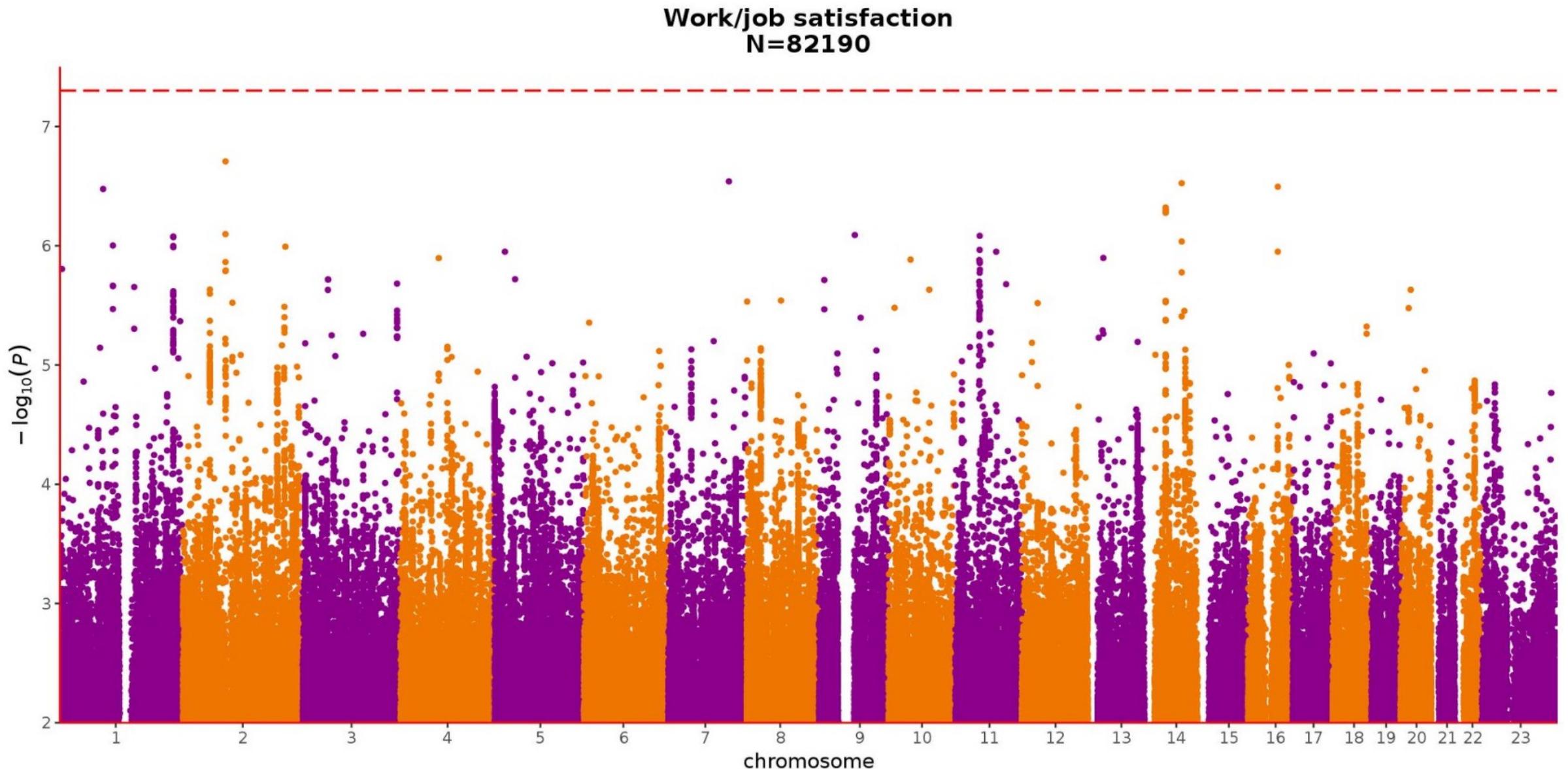
# P-values in ssGWAS - Fat



# P-values in ssGWAS - Protein



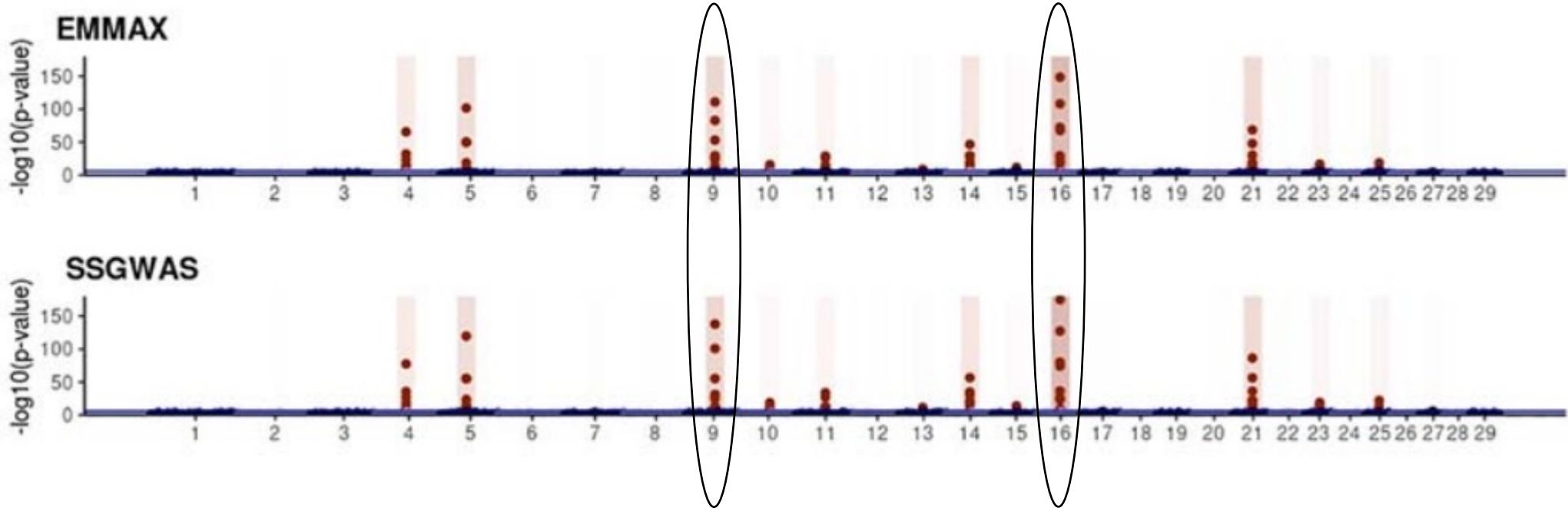
# Non-significant hits



# ssGWAS vs. EMMAX

- Simulated population (1 QTN per CHR)

14k genotyped sires  
Deregressed EBV  
(10 daughters)



14k genotyped sires  
500k Pedigree  
250k phenotypes

Association	EMMAX (Khang et al., 2010)	ssGWAS (Aguilar et al., 2019)
True Positive	55.2 <sup>a</sup> (3.7)	61.6 <sup>a</sup> (8.7)
False Positive	0.0	0.0

# ssGWAS

- ssGWAS works!!!
- Heavy computations
- Soft limit is the same as REML
  - 10k genotyped animals
  - 1M animals in pedigree
  - 1M phenotypes
  - Less than 1 day to finish

# postGSf90 additional options

<http://nce.ads.uga.edu/wiki/doku.php?id=readme.pregsf90>