

Effectiveness of genomic prediction of boar taint components in Pietrain sired breeding populations

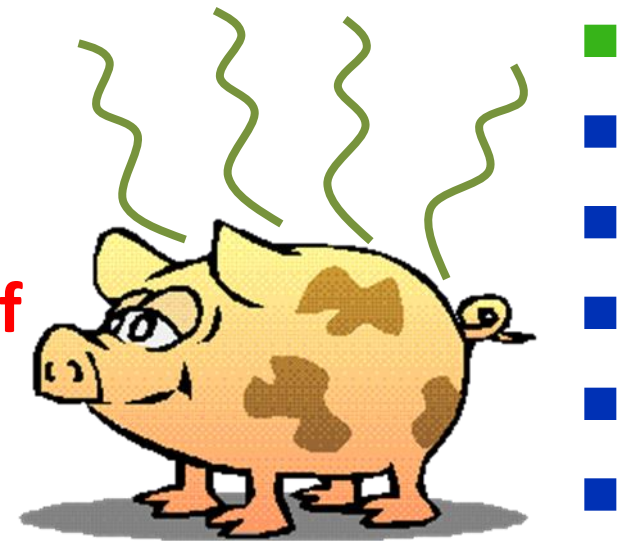
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Boar taint ?

- „... unpleasant flavour and odour of porcine meat“



- Androstenone + Skatole + others
(5 α -androst-16-ene-3-on) (3-methylindol) (indole, phenol, ...)

- Ban of surgical castration of piglets until 2018

→ Fattening of entire boars

- Pig breeding organizations
 - Selection of boars with low incidence of boar taint
 - Set up Pietrain breeding lines



- BUT: costly phenotyping and small Pietrain populations

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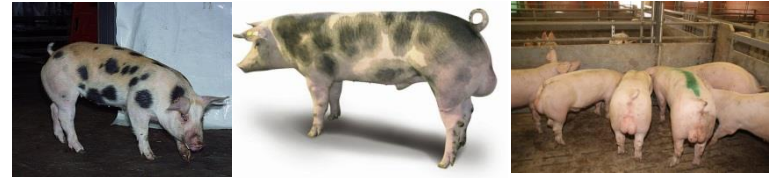
Objective: Genomic selection across breeding organizations

- using data of cross-bred boars

1. Combine the data sets of two boar taint related projects of Pietrain-sired progenies
2. Investigation and comparison of the genetic structure of the different local cross-bred Pietrain-populations
3. Evaluation of genomic prediction accuracy for boar taint components within Pietrain-sired populations

Population structure

Population	No. boars	Type of cross	Project	Year
E1	241	Pi x F1	ENZEMA	Nov. 2009 – Dec. 2010
E2	236	Pi x F1		
E3	120	Pi x F2		
G1	213	Pi x F1	GOGS	Oct. 2012 – May 2013
G2	262	Pi x DL		



- Project EN-Z-EMA (E): **597** crossbred boars
 - Androstenone (And): GC-MS (Gracia-Regueiro & Diaz 1989)
 - Skatole (Ska): RP-HPLC (Dehnhard et al. 1993)
 - **Lab.: IME, Fraunhofer Institut, Schmallingenberg**
- Project GOGS (G): **475** crossbred boars
 - And: rapid enzyme immunoassay (Claus et al. 1997)
 - Ska: UPLC - HPLC Hillenbrand (2000)
 - **Lab.: Tiergesundheitsdienst Bayern**

■ Genotypes

- DNA isolation from muscle samples
- Quality control: call rate > 0.95 , MAF > 0.01
- Project E:
 - Typing of DNA with porcineSNP60 Illumina BeadChip v2
 - **Lab: Life & Brain, Bonn**
- Project G:
 - Typing of DNA with porcineSNP60 Illumina BeadChip v1
 - **Lab: GeneControl, Poing**

- Genotypes

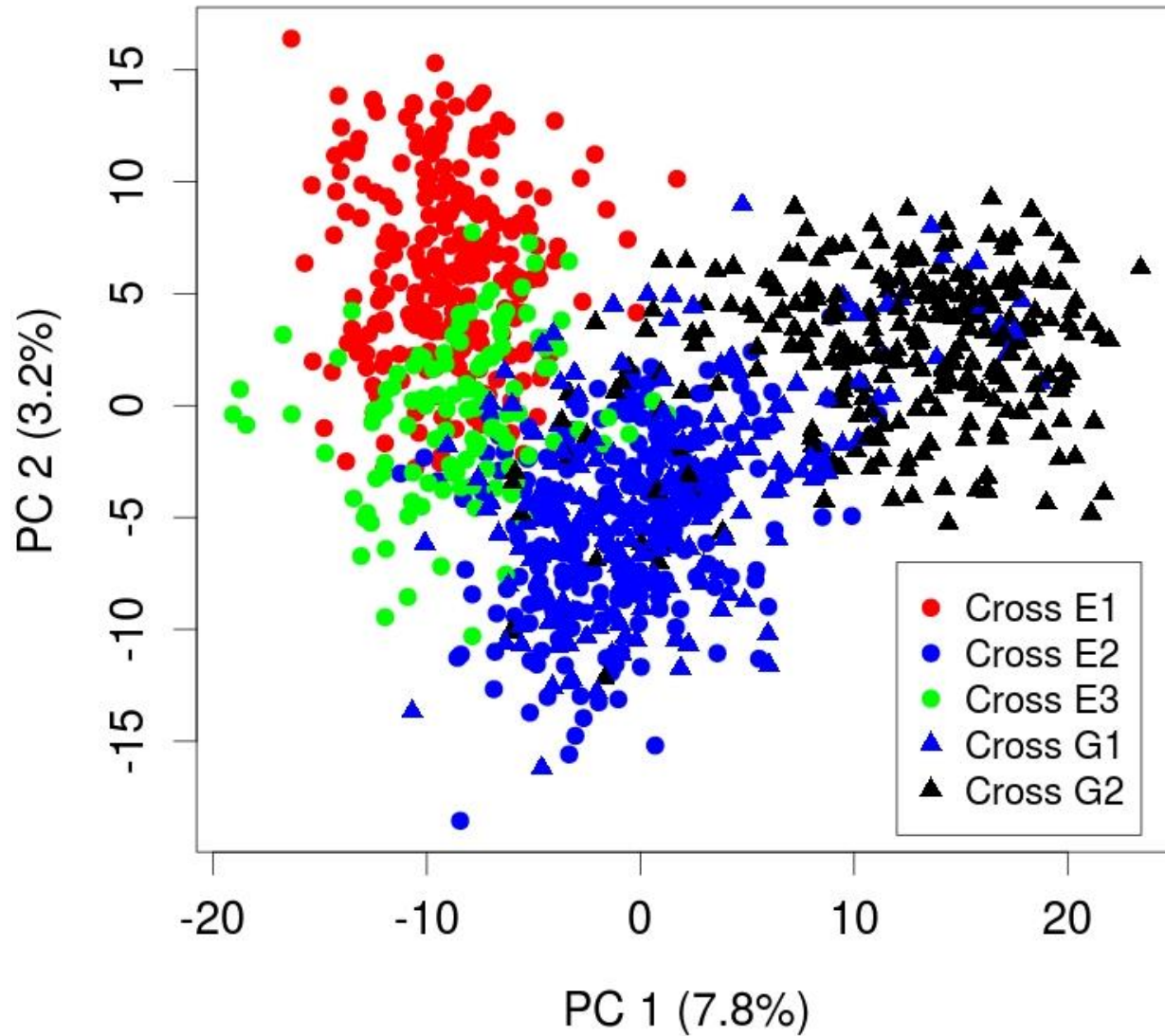
- DNA isolation from muscle samples
- Quality control: call rate > 0.95 , MAF > 0.01

– Dataset F

After quality control:

- 1077 boars
- 42'063 SNP markers

Population structure



- Estimation of the F_{ST} values across the regional populations (Weir 1996)

$$F_{ST} = \frac{\sum_i n_i (\tilde{p}_i - \bar{p})^2 / (r - 1) \bar{n}}{\bar{p}(1 - \bar{p})}$$

	● E1	● E2	▲ G1	▲ G2	● E3
● E1-F2	241	1.23E ⁻⁰²	1.26E ⁻⁰²	1.92E ⁻⁰²	1.44E ⁻⁰²
● E2-F2		236	0.52E ⁻⁰²	1.18E ⁻⁰²	1.38E ⁻⁰²
▲ G1-F2			213	0.97E ⁻⁰²	1.53E ⁻⁰²
▲ G2-F1				266	2.29E ⁻⁰²
● E3-F3					120

- Breeding value (BV) estimation: multivariate sire model
 - Wombat (Meyer 2006)
 - Estimation within the project groups
- Genomic BV estimation: ridge regression BLUP
 - R package: rrBLUP (Endelman et al. 2011)
 - $EBV = Zu + e$
 - $u \sim N(0, G \sigma_a^2)$, G: genomic relationship matrix (vanRaden 2008)

Accuracy of GS

- derived from 5-fold cross-validation
- Sampling number: 5000

1. r_{MP} : Correlation between observed and estimated breeding values

2. $r_{GS} = \frac{r_{MP}}{\sqrt{\overline{r_{EBV}}}}$ (Lande & Thompson 1990, Dekkers 2007)

- $\overline{r_{EBV}}$: mean accuracy of the conventional breeding value

3. $r_{PEV} = \sqrt{1 - (PEV/\sigma_g^2)}$ (Henderson 1975)

S1: CV across all groups of local populations

S2: CV within each local population

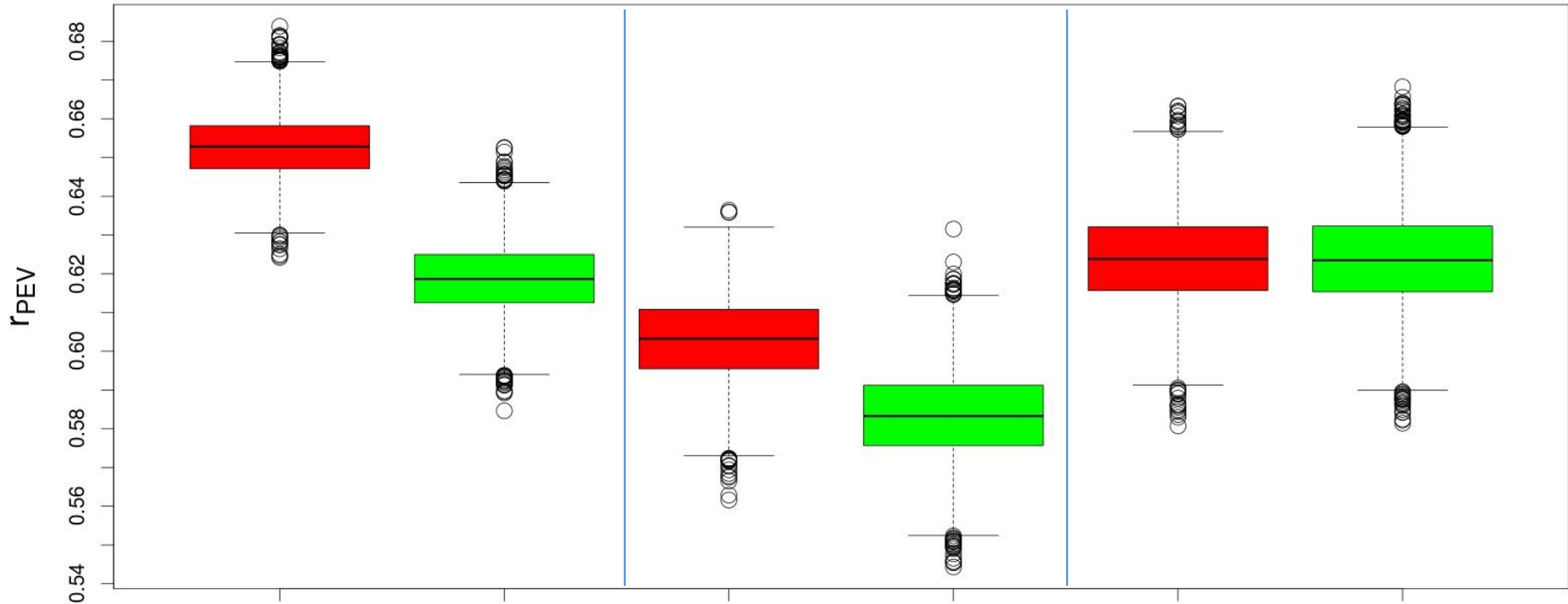
S3: Calibration (C) in one population and validation (V)
in another population

- a) C- and V-population are genetically close related
- b) C- and V-population are genetically less related

Heritability and accuracy of estimated breeding values

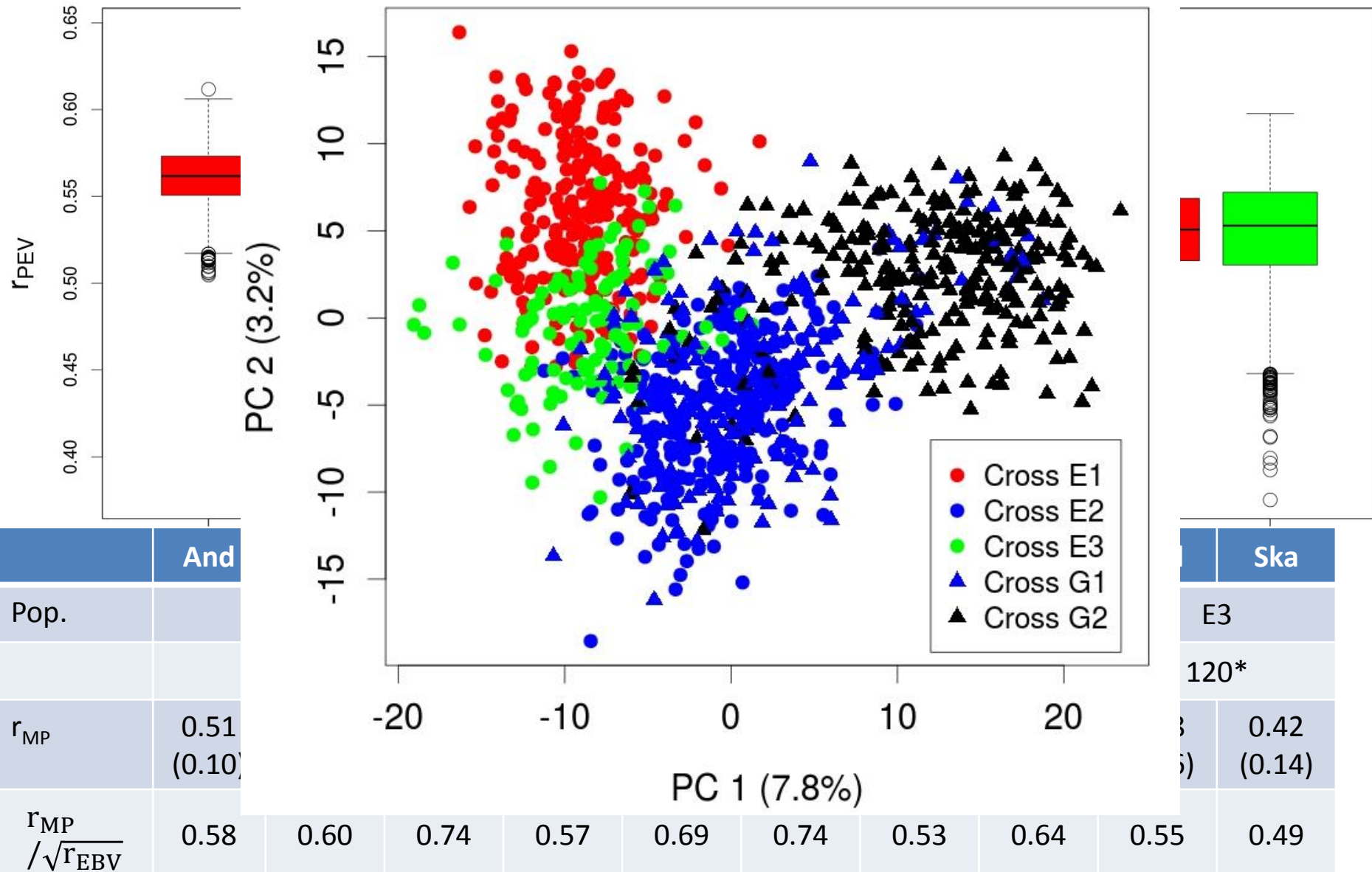
Data	N	Trait	h^2	r_{EBV}
All ENZMA	597	AND	0.58	0.76
		SKA	0.52	0.72
● E1	241	AND	0.63	0.78
		SKA	0.61	0.77
● E2	236	AND	0.67	0.81
		SKA	0.61	0.77
● E3	120	AND	*	*
		SKA	*	*
All GOGS	481	AND	0.58	0.77
		SKA	0.42	0.66
▲ G1	214	AND	0.70	0.78
		SKA	0.57	0.71
▲ G2	267	AND	0.40	0.66
		SKA	0.30	0.58

Scenario 1: 5-fold cross validation across the data sets

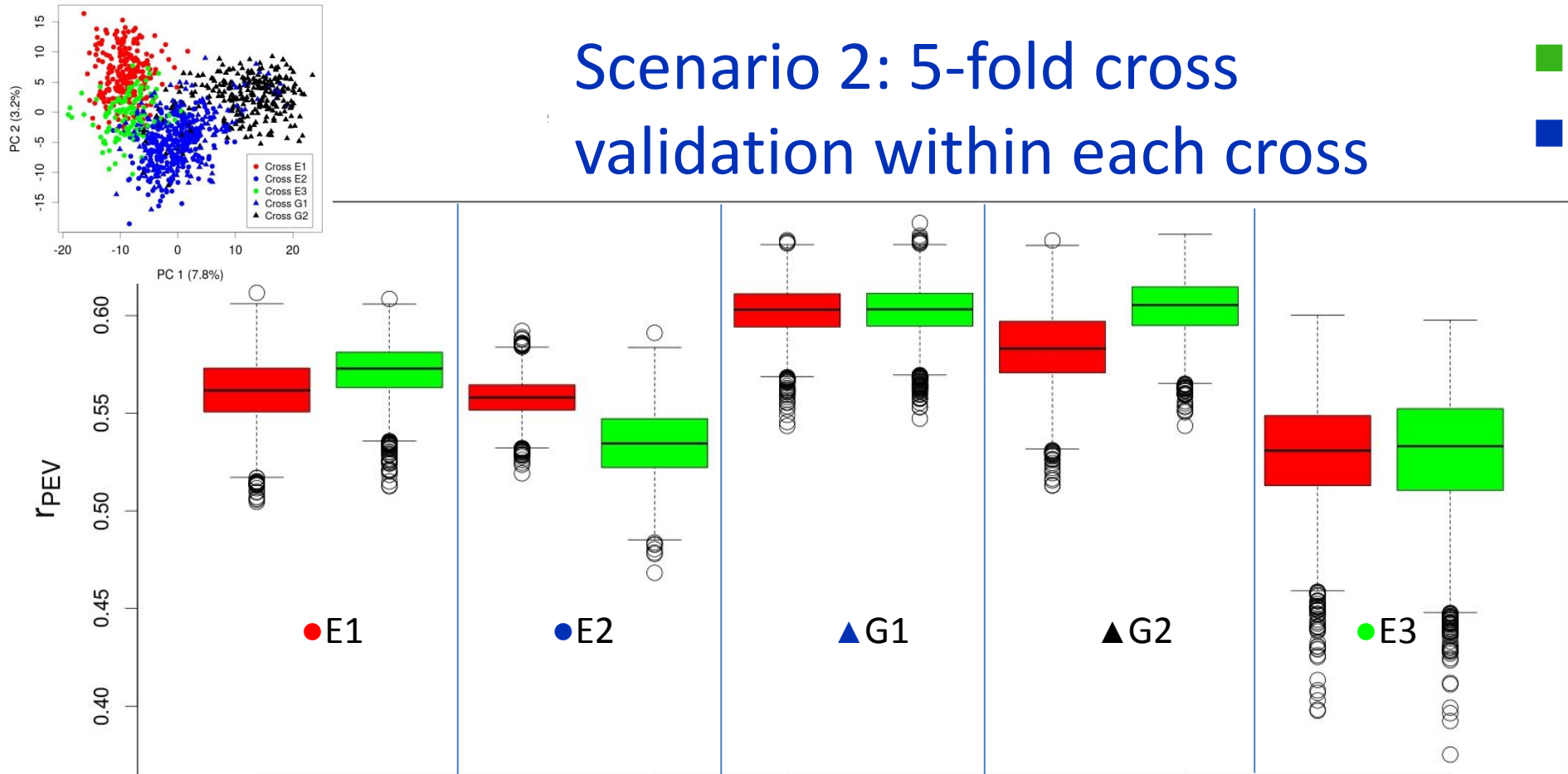


	And	Ska	And	Ska	And	Ska
Pop	All		ENZMA		GOGS	
Val./Cal.	862/215		478/120		380/95	
r_{MP}	0.56 (± 0.05)	0.49 (± 0.05)	0.56 (± 0.06)	0.51 (± 0.07)	0.54 (± 0.07)	0.55 (± 0.07)
$r_{MP}/\sqrt{r_{EBV}}$	0.64	0.58	0.64	0.60	0.62	0.68

Scenario 2: 5-fold cross validation within each cross

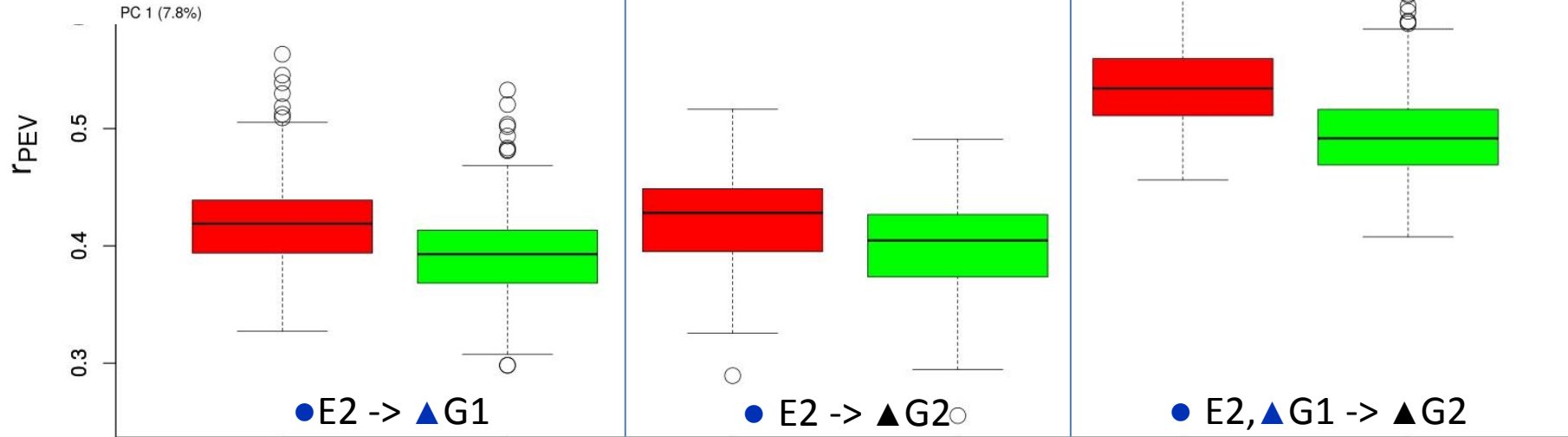
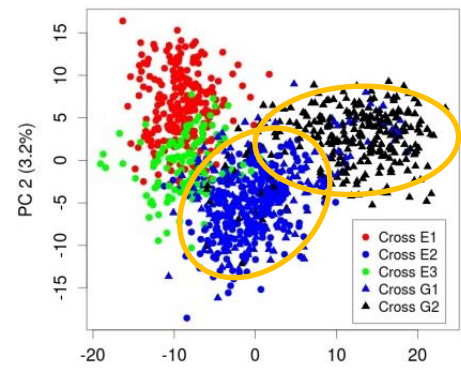


Scenario 2: 5-fold cross validation within each cross



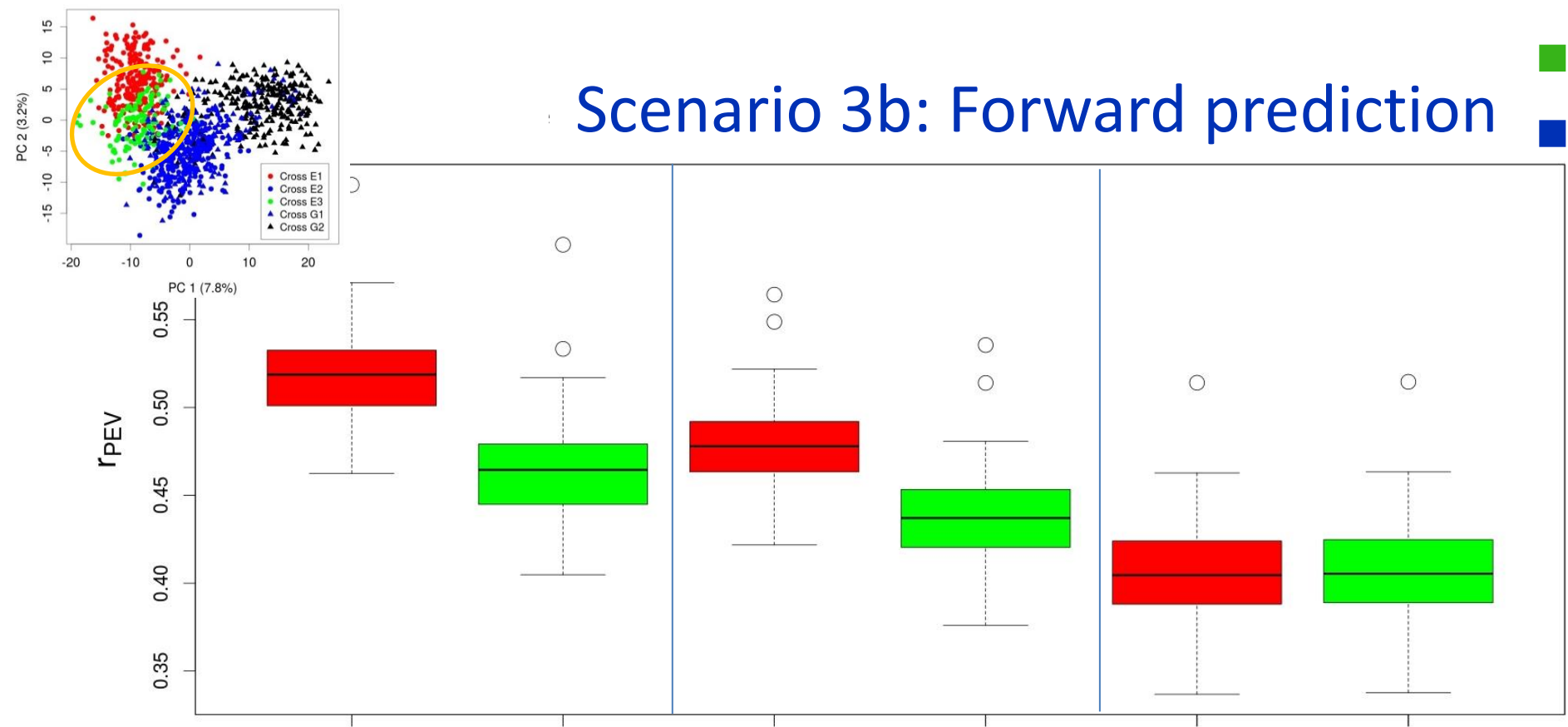
	And	Ska	And	Ska	And	Ska	And	Ska	And	Ska
Pop.	● E1		● E2		▲ G1		▲ G2		● E3	
	242		236		213		262		120*	
r_{MP}	0.51 (0.10)	0.53 (0.11)	0.67 (0.08)	0.50 (0.10)	0.61 (0.09)	0.60 (0.09)	0.47 (0.10)	0.52 (0.09)	0.48 (0.16)	0.42 (0.14)
$\frac{r_{MP}}{\sqrt{r_{EBV}}}$	0.58	0.60	0.74	0.57	0.69	0.74	0.53	0.64	0.55	0.49

Scenario 3a: Forward prediction



	And	Ska	And	Ska	And	Ska
Calib.	● E2	● E2	● E2	● E2	● E2, ▲ G1	● E2, ▲ G1
Valid.	▲ G1	▲ G1	▲ G2	▲ G2	▲ G2	▲ G2
r_{MP}	0.28	0.06	0.10	0.11	0.34	0.26
$r_{MP}/\sqrt{r_{EBV}}$	0.32	0.07	0.11	0.13	0.39	0.31

Scenario 3b: Forward prediction

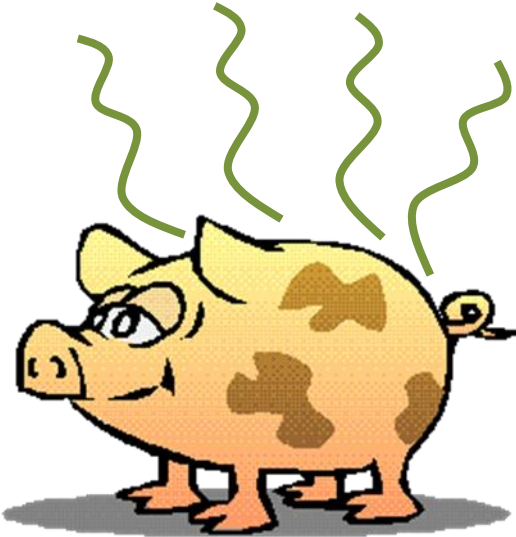


	And	Ska	And	Ska	And	Ska
Calib.	E1,E2,G1,G2	E1,E2,G1,G2	E2,G1,G2	E2,G1,G2	G1, G2	G1, G2
Valid.	● E3	● E3	● E3	● E3	● E3	● E3
r_{MP}	0.26	0.13	0.20	0.10	0.20	-
$r_{MP}/\sqrt{r_{EBV}}$	0.30	0.16	0.23	0.12	0.23	-

- Genomic selection against boar taint using information from commercial crossbreds is promising ($= r_{EBV}$)
- Genetic differences between regional populations
 - Calibrations **cannot** be transferred between subpopulations without considerable **loss of accuracy**

■ Perspectives:

- Combining pure- and cross-bred information
 - Investigation of the relationship between fertility traits and boar taint
- Enlarge the data set



Thank you for your attention!

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