Genome wide diversity: heterozygosity vs. IBD

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Introduction

- Genetic diversity in livestock
- Loss of diversity
- Average diversity (pedigree) diversity on specific regions on the genome (non-neutral trait, i.e. growth and fertility)
- Dense marker maps (SNPs)
 Predict diversity on untyped regions (i.e. unknown QTL)





Genetic diversity evaluation



 Molecular / population genetics
 Heterozygosity % individuals that have 2 different alleles

Quantitative genetics

- Average relatedness
 - % DNA that is identical by descent (IBD)



Aim

Estimation of genetic diversity on untyped regions over the genome

Original Compare two methods: IBD and heterozygosity

→ Which method best predicts genetic diversity between markers?



Simulation

1 chromosome of 1 Morgan
1,800 SNP markers
Effective population size = 100
100 + 3 generations
3 generations: 2,100 genotyped animals



Genetic diversity over the genome





True genetic diversity



Heterozygosity of the untyped marker





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IBD-based genetic diversity







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Population level



IBD

Diversity-IBD = 1 – average P(IBD)
 (Details in Meuwissen (1997))

Heterozygosity

• He = # heterozygotes / # animals







Genetic diversity over marker intervals



True diversity

→ Large variation between markers



Genetic diversity over marker intervals



True diversity

→ Difference in level







Correlations: IBD vs. heterozygosity

Marker intervals

	True	4	10	20	40
IBD	0.27	0.42	0.57	0.72	0.82
Не	0.16	0.23	0.38	0.57	0.70



Conclusions

IBD-method better predictor for genetic diversity on untyped regions of the genome

More insight into genetic diversity on specific regions of the genome

Can be taken into account when selecting animals for conservation



Questions?

IBD-method better predictor for genetic diversity on untyped regions of the genome

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