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Estimation of heritability for dairy traits, combining pedigree with dense snp information on some animals

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Introduction

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- Genetics of feed utilisation complex: milk, fat and protein production, feed intake, BCS and liveweight
- Interesting for agroproductie 21^{ste} century:
 Energy balance: robustness, health & fertility
 - Feed inefficiency: greenhouse gas emissions
 - Feed efficiency: economic efficiency
- "Something new to be learned in this area?"
 → better insight using using genomics

Questions Data GWAS Dutch dataset: 639 Holstein first lactation heifers, fed ad libitum TMR but also utilise SNP information to improve estimation of genetic parameters? Recording of milk yield, milk composition, Replace pedigree by genomic relationships? liveweight and feed intake • Utilise information on individual SNP effects? → average first 15 weeks of lactations Difficulty: animals with scarce and expensive Ca 580 animals DNA → genotyped Illumina 50k SNP panel historical phenotypes but without DNA available Combine genotyped and none genotyped relationships? ■ Quality control checks → 517 animals left Animal Breeding & Genomics Centre Animal Breeding & Genomics Centre LIVESTOCK RESEARCH



Statistical models				
 Relationship matrix: A based on pedigree (639 pheno 3363 in pedigree) 	typed animals plus			
 G from SNPs (517 genotyped+pt 	neno. animals)			
$G = \frac{\mathbf{Z}\mathbf{Z}'}{2\sum p_i(1-p_i)} \qquad (Var$	nRaden 2008)			
 H⁻¹ combine A⁻¹ and G⁻¹ (639 animals) (Aguilar et al. 2010) 				
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Statistical models	
WG: weighted genom	ic relationship matrix
Step 1: estimate SNP	effects BAYESC
$y = 1_n \mu + fixed _effect$	$bs + \sum_{j=1}^{m} (X_j(q_j v_j)) + e$
Step 2: weighted G	
$WG = \frac{ZDZ'}{2\sum p_i(1-p_i)}$	D diagonal matrix weights per SNP: squared allele substitution effects, rescaled to be 1.0 across all loci.
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	h^2				SI	2		
RM	A	A	G	H	Α	A	G	H
# phenotypes	517	639	517	639	517	639	517	639
Milk	0.48				0.13			
Fat %	0.89				0.10			
DMI	0.83				0.11			
LW	0.50				0.13			
fferences A&	G pai	oer Ver	bvla e	et al W	/CGA	LP		

RM P. Prob.	G none	WG p>0.10	WG p>0.05	WG p>0.01	WG p>0.001	WG All
# SNP		3	7	116	1237	43011
-SINI				110	4237	45011
h ²	0.77					
SE	0.08					

Conclusion

- Using SNP relationships improves estimation of variance components even when 517 phenotypes were used instead of 639 phenotypes Combining SNP and pedigree information gave best estimate
- Differences between estimates with G and A
 Scale A and G to same base population?
 IBD v. IBS (Janss)

 - → Also an issue when creating H matrix

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