Imputation of medium density genotypes from custom low density genotype panel in sheep

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Motivation

- •Aim to have multi-breed genomic evaluations in Irish sheep in 2017
- Accurate genomic predictions require a large reference population
- Average lamb value < cattle value

Need a low cost strategy for genomic selection



iaacmghzq abehkzi yufxbhjl

M__a_l is g_ing to g_ to _i_b__t_ ne_t y_ar in s__ing



iaacmghzq abehkzi yufxbhjl

John and Michael are good friends. Michael's uncle lives in Timbuktu

M___a_l is going to go to _i_b__t_ next year in spring



iaacmghzq abehkzi yufxbhjl

John and Michael are good friends. Michael's uncle lives in Timbuktu

Michael is going to go to Timbuktu next year in spring



iaacmghzq abehkzi yufxbhjl

John and Michael are good friends. Michael's uncle lives in Timbuktu

Mychael is going to go to Timbuktu next year in spring



Study objective

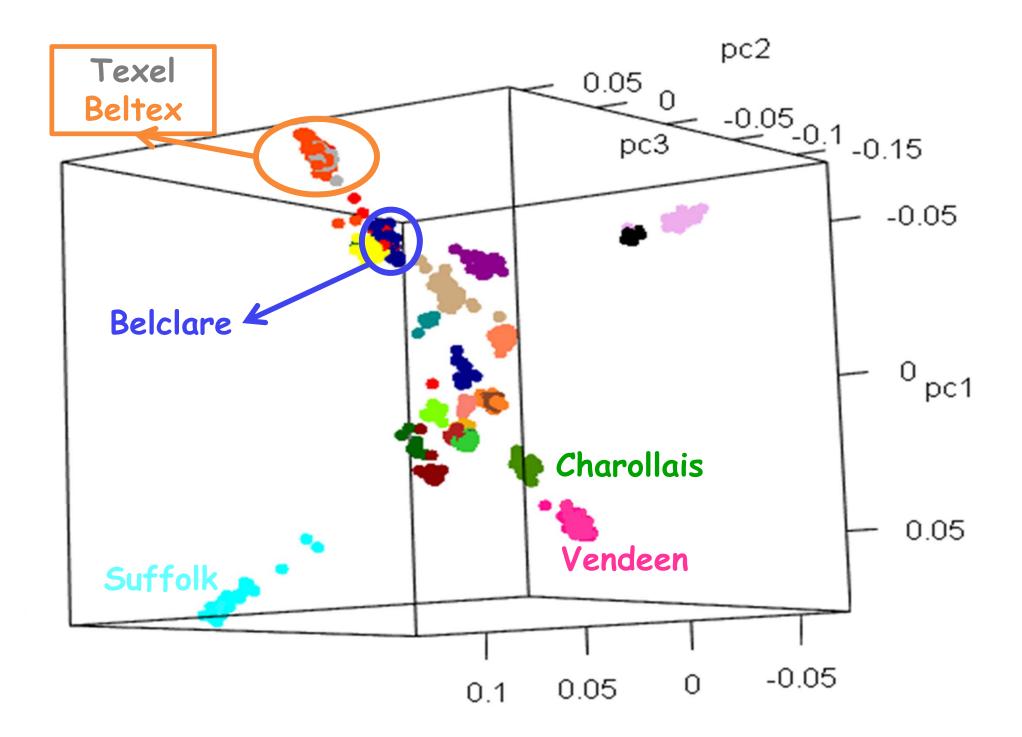
•Current thinking is at least a 50K panel (i.e., 50,000 markers) is necessary

•Generate several thousand "50K genotypes" at a low cost

•Many breeds

 Texel, Suffolk, Vendeen, Charollais, Belclare





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- •Many breeds
 - Texel, Suffolk, Vendeen, Charollais, Belclare (+Beltex)
- Scottish friends also shared Texel genotypes



Low density panel

- ·15,000 SNPs
- •Equally spaced SNPs across the genome for imputation
- •Greater density in putative QTL regions
- Several parentage panels
- (Putative) causal mutations
 - •Scrapie, spiderlamb, GDF, Booroola, Inverdale....



Strategy •Imputation accuracy of low density to medium density

Reference population

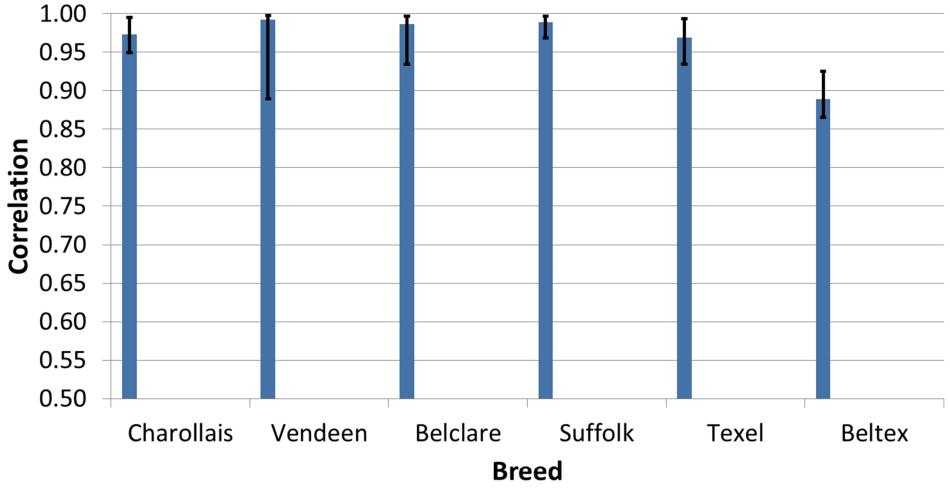
- Multi-breed Irish genotypes
- ·Within-breed Irish genotypes
- Multi-breed Irish+Scottish genotypes



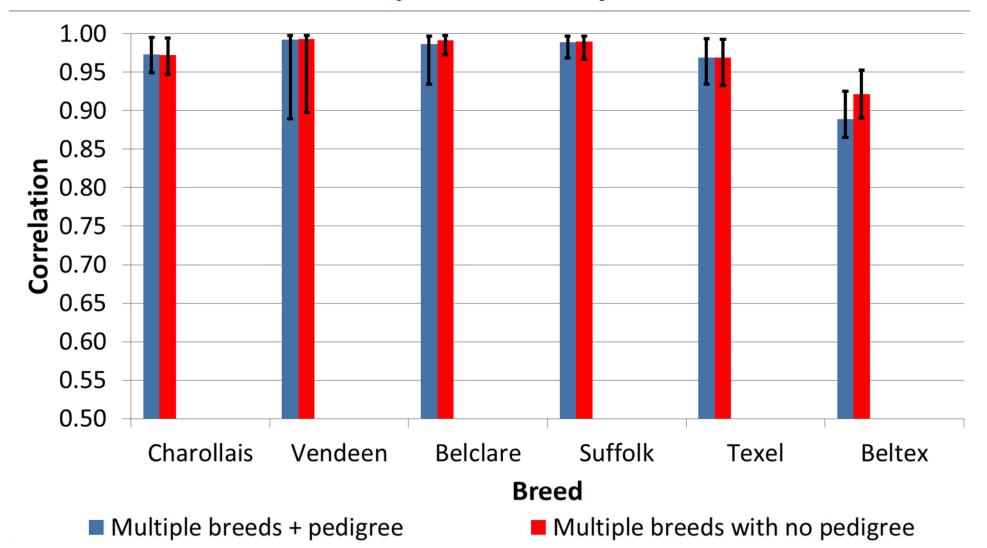
Data

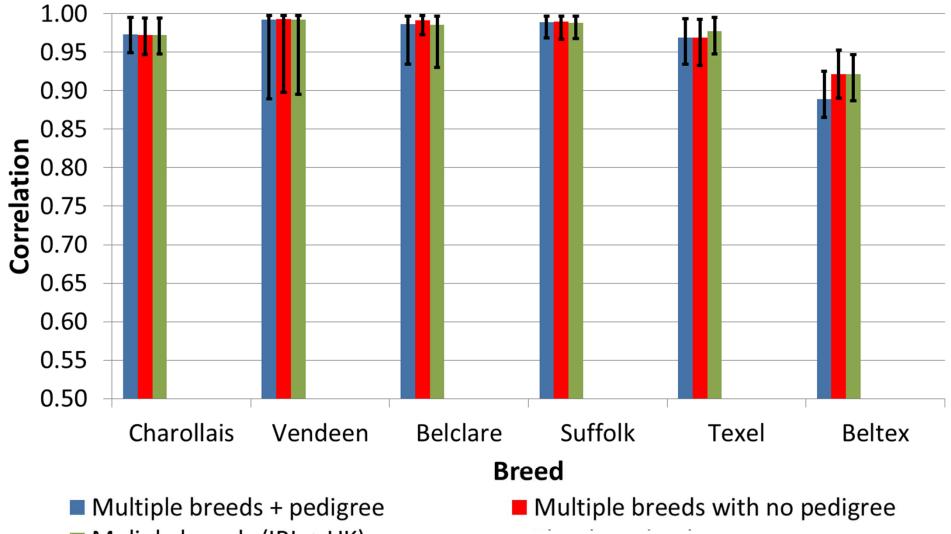
Breed	Reference	Validation
Belclare	559	75
Charollais	564	75
Suffolk	684	75
Vendeen	551	75
Texel	393 + 1796	75
Beltex		64



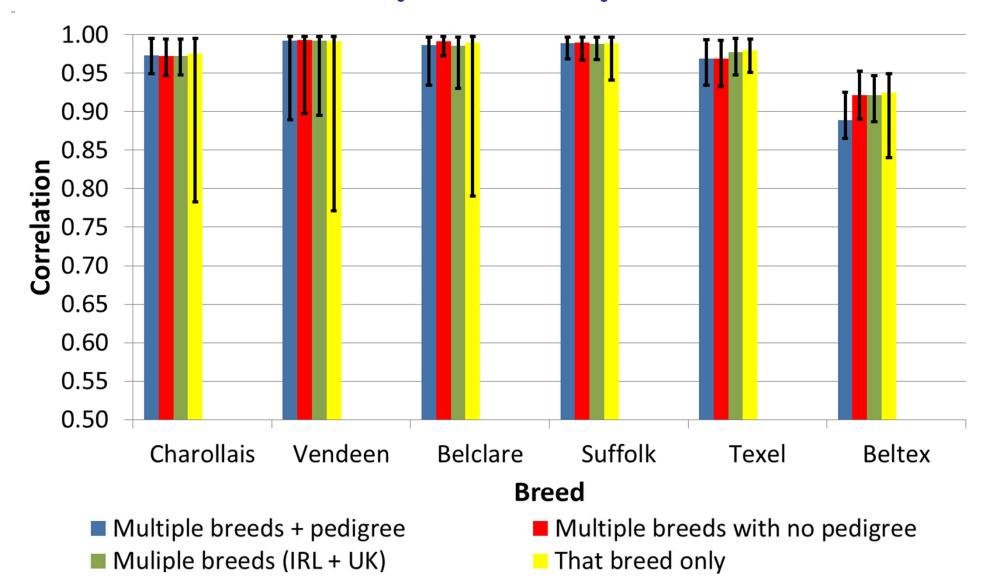


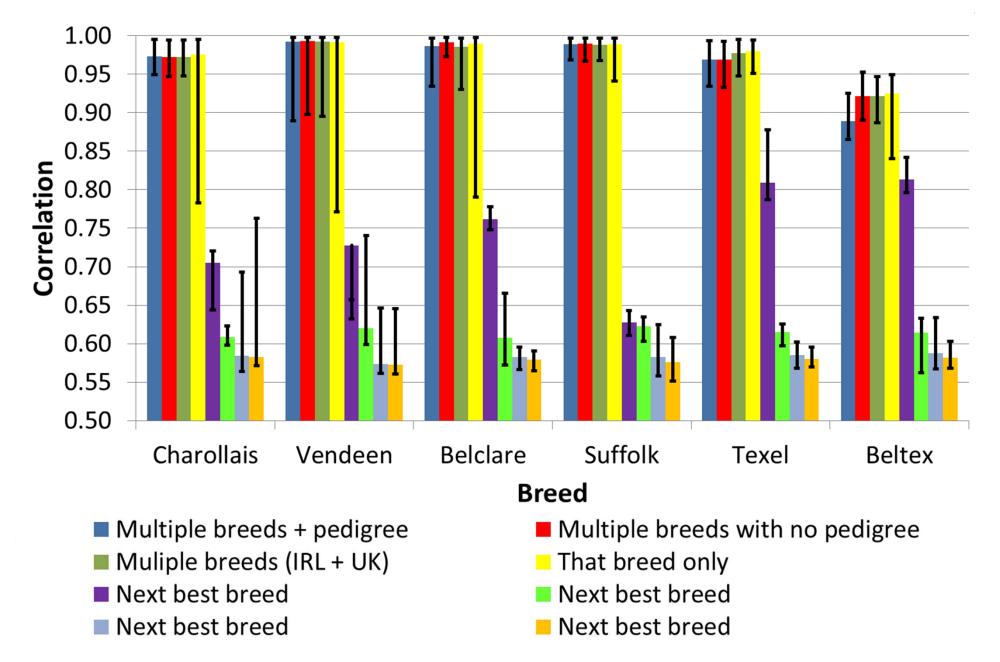
Multiple breeds + pedigree



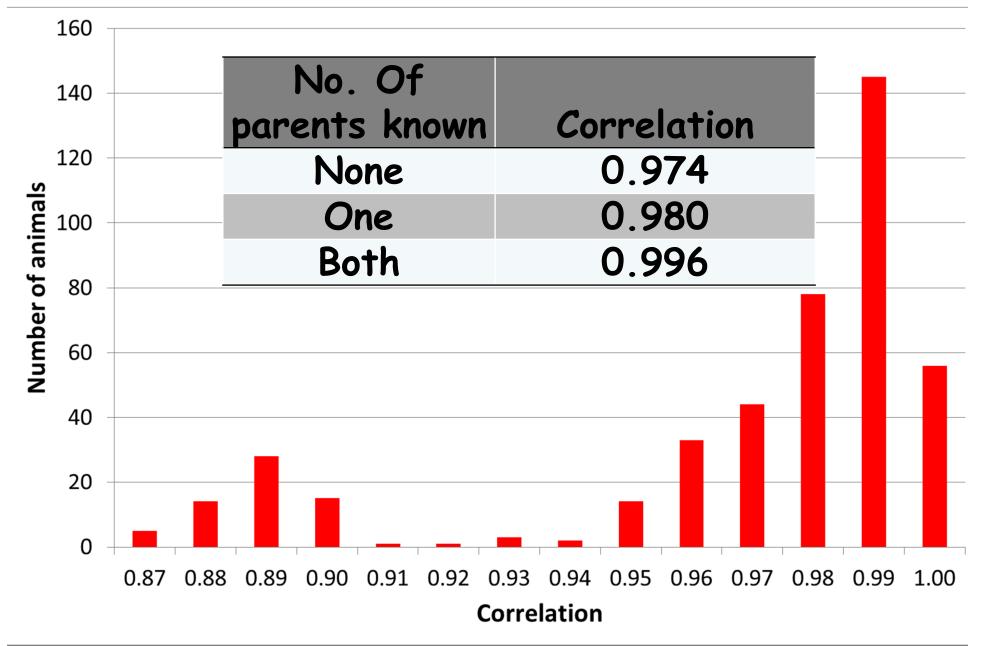


Muliple breeds (IRL + UK)





Accuracy per animal



Conclusions

- Information content on low density panel is excellent
 - •Almost all SNPs segregating
- High accuracy of imputation from low to high density
 - No impact of imputing all breeds in one step
 Probably better off to avoid the "rogues"
- Greater accuracy when parents (haplotypes) in the reference population



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