Genomic BLUP

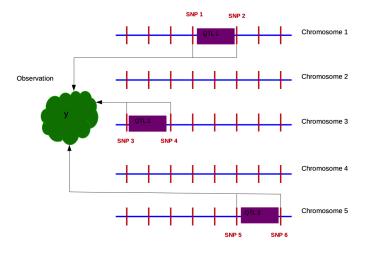
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So Far

- Estimate effect of few SNP loci linked to QTL
- ▶ Use parameter estimates to predict genomic breeding values
- ▶ **New**: Many SNP, find the important ones

Situation



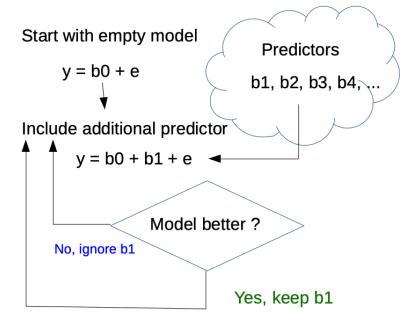
Goal: Find SNP 1 - SNP 6 out of the many SNPs

Approaches in Fixed Linear Model Framework

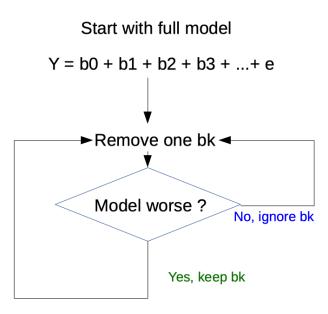
Two Approaches

- 1. Forward selection: Start with empty model, include predictors that improve model
- 2. Backward elimination: Start with full model, remove predictors as long as model does not get worse

Forward Selection



Backward Elimination



Model Selection With Genomic Data

- Only backward elimination really works in practical problems
- ► Large number of predictors $(1.5 * 10^5)$
- How to determine sequence of predictors to eliminate
- Fitting the full model is problematic

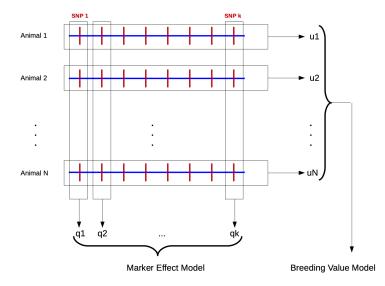
Mixed Linear Effect Model

- One solution: replace fixed linear effect model by mixed linear effect model (mle)
- ► MLE: additional random effect besides error term
- Random effects are specified by expected value and variance
- In livestock breeding MLE have a good reputation from BLUP animal model

MLE In Genomics

- Two different parametrizations
- 1. Marker Effect Model (MEM)
- 2. Breeding Value Model (BVM)

Overview



Marker Effect Model

In MEM random effects of markers are directly included in the model. For an idealized data set we can write

$$y = 1_n \mu + Wq + e$$

where

y vector of length n with observations

 μ general mean denoting fixed effects

 1_n vector of length n of all ones

q vector of length m of random SNP effects

W design matrix relating SNP-genotypes to observations

e vector of length *n* of random error terms

Breeding Value Model

e

$$y = Xb + Zg + e$$

vector of length *n* of random error terms

where

y vector of length n with observations
b vector of length r with fixed effects
X incidence matrix linking elements in b to observations
g vector of length t with random genomic breeding values
Z incidence matrix linking elements in g to observations