Lab: Applications for Ordinary Least Squares and Mixed Models

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OLS and MM example 1: Balanced maize data

Learning objectives:

1. Brief overview of ordinary least squares (OLS) and mixed models (MM)
2. Estimate genetic values and h^2 using OLS and MM.
3. Learn to deal with unbalanced data

Maize Dataset

- 62 recombinant inbred line (RILs) from a cross between B73 and MO17.
- Randomized complete block design
- Two replications at four locations
- Traits: days to pollen, days to silking, anthesis/silking interval (ASI) and plant height.
  - We'll use height as the response variable.
- See Isik, Holland and Maltecca (2017)

Loading the data.

```
maize <- read.csv("~/Downloads/MaizeRILs.csv")

head(maize)
```

```
##   location rep block plot   RIL pollen silking ASI height
## 1      ARC   1     4   28 RIL-1     73      77   4  182.0
## 2      ARC   2     6   47 RIL-1     74      79   5  169.2
## 3      CLY   1     5   36 RIL-1     71      74   3  213.0
## 4      CLY   2     4  223 RIL-1     73      77   4  203.0
## 5     PPAC   1     8   64 RIL-1     97     101   4  155.6
## 6     PPAC   2     5   40 RIL-1     95     100   5  177.6
```

Obtaining genetic values with OLS

- For this dataset we can fit the following model:

$$y_{ijk} = \mu + L_i + Rep(L)_{ij} + G_k + GL_{ik} + e_{ijk}$$

- y_{ijk} is the phenotype (height)
- L_i is the fixed effect of location i
- $Rep(L)_{ij}$ is the fixed effect of replicate j nested within location i
- G_k is the fixed effect of RIL k, GL_{ik} is the interaction of RIL k and location i and e_{ijk} is the residual.

Here's everything except the error term is considered as a fixed effect

Obtaining genetic values with OLS

- Fit the linear model with lm in R

```
#rep is coded as 1 and 2. So make sure R knows its a factor
maize$rep <- as.factor(maize$rep)
mod1 <- lm(height ~ location*RIL + rep:location, data = maize)

anova(mod1)
```

```
## Analysis of Variance Table
##
## Response: height
##                Df Sum Sq Mean Sq  F value    Pr(>F)
## location        3  84931 28310.4 436.3090 < 2.2e-16 ***
## RIL            61 154938  2540.0  39.1448 < 2.2e-16 ***
## location:RIL  183  20999   114.8   1.7685 1.643e-05 ***
## location:rep    4   3594   898.6  13.8482 3.408e-10 ***
## Residuals     244  15832    64.9
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Obtaining genetic values with OLS

- Use the output of lm to estimate the marginal means
- For RIL-11 we can calculate the marginal means as:

$$RIL11 = \mu + \bar{L} + G_{RIL11} + \bar{GL}_{RIL11} + \bar{Rep}(L)$$

Obtaining genetic values with OLS

```r
#intercept
MU <- as.numeric(coef(mod1)["(Intercept)"] )
#locations
LOC.eff <- sum(as.numeric(coef(mod1)[c("locationCLY",
          "locationPPAC","locationTPAC")] ))/4
#RIL
RIL1.eff <- as.numeric(coef(mod1)["RILRIL-11"] )
#RIL x Location
RIL1.LOC.eff <- sum(as.numeric(coef(mod1)
               [c("locationCLY:RILRIL-11",
                  "locationPPAC:RILRIL-11",
                  "locationTPAC:RILRIL-11")] ))/4
#Rep within location
Rep.eff <- sum(as.numeric(coef(mod1)[c("locationARC:rep2",
               "locationCLY:rep2", "locationPPAC:rep2",
               "locationTPAC:rep2")] ))/8

RIL_11 <- MU + LOC.eff + RIL1.eff + RIL1.LOC.eff + Rep.eff

print(RIL_11)
```

```
## [1] 182.875
```

Estimating heritability from ANOVA/OLS

$$h^2 = \frac{\sigma^2_{RIL}}{\sigma^2_{RIL} + \frac{\sigma^2_{RIL \times LOC}}{n_l} + \frac{\sigma^2_e}{n_r n_l}}$$

- EMS from ANOVA
  - Location: $\sigma^2_e + n_b\sigma^2_{GL} + n_g\sigma^2_B(L) + n_b n_g \sigma^2_L$
  - B(L): $\sigma^2_e + n_g\sigma^2_B(L)$
  - RIL: $\sigma^2_e + n_b\sigma^2_{GL} + n_g\sigma^2_B(L)$
  - RIL x Loc: $\sigma^2_e + n_b\sigma^2_{GL}$

Estimating heritability from ANOVA/OLS

- Since the design is balanced we can estimate H^2 using ANOVA

$$h^2 = \frac{\sigma^2_{RIL}}{\sigma^2_{RIL} + \frac{\sigma^2_{RIL\times LOC}}{n_l} + \frac{\sigma^2_e}{n_r n_l}}$$

- $\sigma^2_{RIL\times LOC} = \frac{MS(RIL\times LOC) - MS(Error)}{n_r}$,
  $\sigma^2_{RIL} = \frac{MS(RIL) - MS(RIL\times LOC)}{n_r n_l}$, and
  $\sigma^2_e = MS(Error)$

Estimating heritability from ANOVA/OLS

```
anova.res <- as.data.frame(anova(mod1))

sigma_err <- anova.res[5,3]
sigma_G.E <- (anova.res[3,3] - sigma_err) / 2
sigma_G <- (anova.res[2,3] - anova.res[3,3]) / 8

H2.OLS <- sigma_G / (sigma_G + sigma_G.E/4 + sigma_err/8)
print(H2.OLS)
```

```
## [1] 0.9548218
```

Obtaining genetic values (BLUEs) with a mixed model

- We will fit a mixed model to estimate line values for each RIL
  - *RIL* as a fixed effect, and *Loc* and *Rep* as random effects
  - $Var(Loc) \sim N(0, \mathbf{I}\sigma^2_{LOC})$, $Var(rep) \sim N(0, \mathbf{I}\sigma^2_{rep})$, and $Var(e) \sim N(0, \mathbf{I}\sigma^2_e)$

Obtaining genetic values (BLUEs) with a mixed model in lme4

- Random terms are specified by '(1|some term)'.
  - '(1|location/rep)' is the random effect of rep nested within location
  - '(1|location:RIL)' is the random effect of location x RIL interaction

```
library(lme4)
mod2 <- lmer(height ~ RIL + (1|location/rep) + (1|location:RIL), maize)

#List the estimates for the fixed effects
summary(mod2)$coefficients[1,1] + summary(mod2)$coefficients[2,1]
```

```
## [1] 182.875
```

Estimating heritability with a mixed model in lme4

- Here, all terms with the exception of μ will be considered random.

```
mod3 <- lmer(height ~ 1 + (1|RIL) +
               (1|location/rep) +
               (1|location:RIL), maize)

#extract the variance components
MM.varcomps <- as.data.frame(VarCorr(mod3))

sigma_err.MM <- MM.varcomps[5,4]
sigma_G.E.MM <- MM.varcomps[1,4]
sigma_G.MM <- MM.varcomps[2,4]

H2.MM <- sigma_G.MM /
  (sigma_G.MM + sigma_G.E.MM/4 + sigma_err.MM/8)

print(H2.MM)
```

```
## [1] 0.9548218
```

BLUPs for maize height

- When we want to make a prediction on a random term in the model the predicted value is called BLUP
- In lme4:

```
mod3 <- lmer(height ~ 1 + (1|RIL) + (1|location/rep)
             + (1|location:RIL), maize)

#extract the blups for RILs
blups_m3 <- ranef(mod3)$RIL
```

- More on BLUPs later!

On your own

- Run a similar analyis with the unbalanced data and compare OLS and MM approaches
- Which is more trustworthy?

Reference

- Isik, F., Holland, J. & Maltecca, C. Genetic data analysis for plant and animal breeding. (Springer, 2017).