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```
#####ask yourself if the h2 suppose to be constant or change?
cc <- read.csv("DATA/ppj220030-sup-0002-tables1.csv")
table(cc$date)
```

```
##
## 1-Sep 12-Aug 14-Aug 16-Aug 20-Aug 22-Aug 23-Aug 26-Aug 3-Sep 30-Aug 5-Sep
##      878      878      878      878      878      878      877      878      878      878
## 6-Jul
##      878
```

```
### adding replication information
```

```
cc$Rep <- "Rep2"
cc[cc$Row< 3000,] $Rep <- "Rep1"
```

```
###Checking our data
```

```
head(cc)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 1 1001 1001 and 1002      A641 Ames 19311  Nitrogen  6-Jul      29.747379 Rep1
## 2 1001 1001 and 1002      A641 Ames 19311  Nitrogen 23-Aug      15.537977 Rep1
## 3 1001 1001 and 1002      A641 Ames 19311  Nitrogen  1-Sep      15.189726 Rep1
## 4 1001 1001 and 1002      A641 Ames 19311  Nitrogen 22-Aug      19.037283 Rep1
## 5 1001 1001 and 1002      A641 Ames 19311  Nitrogen 12-Aug       9.793073 Rep1
## 6 1001 1001 and 1002      A641 Ames 19311  Nitrogen 26-Aug      21.378869 Rep1
```

```
tail(cc)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10530 4583 4583 and 4584      NC262 PI 531085  Nitrogen 23-Aug      17.90676
## 10531 4583 4583 and 4584      NC262 PI 531085  Nitrogen 22-Aug      18.60264
## 10532 4583 4583 and 4584      NC262 PI 531085  Nitrogen 30-Aug      15.17244
## 10533 4583 4583 and 4584      NC262 PI 531085  Nitrogen  5-Sep       9.91415
## 10534 4583 4583 and 4584      NC262 PI 531085  Nitrogen  1-Sep      15.37183
## 10535 4583 4583 and 4584      NC262 PI 531085  Nitrogen 12-Aug      20.12580
##      Rep
## 10530 Rep2
## 10531 Rep2
## 10532 Rep2
## 10533 Rep2
## 10534 Rep2
## 10535 Rep2
```

```
dim(cc)
```

```
## [1] 10535      8
```

```
table(cc$Treatment)
```

```
##  
##      Nitrogen No Nitrogen  
##      5279      5256
```

```
table(cc$Rep)
```

```
##  
## Rep1 Rep2  
## 5364 5171
```

```
table(cc$date)
```

```
##  
## 1-Sep 12-Aug 14-Aug 16-Aug 20-Aug 22-Aug 23-Aug 26-Aug 3-Sep 30-Aug 5-Sep  
## 878 878 878 878 878 878 878 877 878 878 878  
## 6-Jul  
## 878
```

```
#### sub-setting for date 6 Jul
```

```
d1 <- subset(cc, date %in% "6-Jul")  
head(d1)
```

```
##      Row  Row.Numbers Genotype Pedigree Treatment date Canopy_Coverage Rep  
## 1 1001 1001 and 1002 A641 Ames 19311 Nitrogen 6-Jul 29.747379 Rep1  
## 20 1003 1003 and 1004 B76 PI 550483 Nitrogen 6-Jul 47.451452 Rep1  
## 33 1005 1005 and 1006 A654 PI 587141 Nitrogen 6-Jul 43.296481 Rep1  
## 45 1007 1007 and 1008 A239 Ames 23405 Nitrogen 6-Jul 36.716503 Rep1  
## 60 1009 1009 and 1010 A682 PI 587143 Nitrogen 6-Jul 37.797254 Rep1  
## 70 1011 1011 and 1012 C123 Ames 19313 Nitrogen 6-Jul 8.849936 Rep1
```

```
tail(d1)
```

```
##      Row  Row.Numbers Genotype Pedigree Treatment date Canopy_Coverage  
## 10472 4569 4569 and 4570 C0255 Ames 27107 Nitrogen 6-Jul 34.59631  
## 10482 4573 4573 and 4574 N192 PI 550566 Nitrogen 6-Jul 37.25802  
## 10489 4575 4575 and 4576 NC312 Ames 27152 Nitrogen 6-Jul 29.30559  
## 10507 4577 4577 and 4578 NC292 Ames 27142 Nitrogen 6-Jul 27.58182  
## 10516 4579 4579 and 4580 SD44 PI 524969 Nitrogen 6-Jul 36.25121  
## 10526 4583 4583 and 4584 NC262 PI 531085 Nitrogen 6-Jul 12.03667  
##      Rep  
## 10472 Rep2  
## 10482 Rep2  
## 10489 Rep2  
## 10507 Rep2  
## 10516 Rep2  
## 10526 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit1 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
          data=d1)
```

```
#generating ANOVA table
summary(aov(fit1))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232  63718      275   8.667 < 2e-16 ***
## Treatment       1   7892     7892 249.031 < 2e-16 ***
## Rep            1   1604     1604  50.615 4.9e-12 ***
## Genotype:Treatment 224   6987       31   0.984   0.549
## Residuals      419  13278       32
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the treatments.

```
##Calculating Vprogeny
out1 <- summary(aov(fit1))[[1]]
vprogeny1 <- (out1[1,3] - out1[4,3])/(2*2)
```

```
#Calculating H2
h2_1 = vprogeny1/((vprogeny1 + out1[4,3]/(2*2)))
```

H2 is very high 0.89. This means that canopy coverage is highly influenced by genetic effects more than by the environment.

```
####For in the date 12 Aug
d2 <- subset(cc, date %in% "12-Aug")
head(d2)
```

##	Row	Row.Numbers	Genotype	Pedigree	Treatment	date	Canopy_Coverage	Rep
##	5	1001	1001 and 1002	A641	Ames 19311	Nitrogen 12-Aug	9.793073	Rep1
##	24	1003	1003 and 1004	B76	PI 550483	Nitrogen 12-Aug	72.860293	Rep1
##	35	1005	1005 and 1006	A654	PI 587141	Nitrogen 12-Aug	49.738835	Rep1
##	39	1007	1007 and 1008	A239	Ames 23405	Nitrogen 12-Aug	70.890916	Rep1
##	50	1009	1009 and 1010	A682	PI 587143	Nitrogen 12-Aug	71.139594	Rep1
##	66	1011	1011 and 1012	C123	Ames 19313	Nitrogen 12-Aug	5.203955	Rep1

```
tail(d2)
```

##	Row	Row.Numbers	Genotype	Pedigree	Treatment	date	Canopy_Coverage
##	10468	4569	4569 and 4570	C0255	Ames 27107	Nitrogen 12-Aug	31.74641
##	10484	4573	4573 and 4574	N192	PI 550566	Nitrogen 12-Aug	35.34276
##	10499	4575	4575 and 4576	NC312	Ames 27152	Nitrogen 12-Aug	69.51793
##	10502	4577	4577 and 4578	NC292	Ames 27142	Nitrogen 12-Aug	25.29349
##	10519	4579	4579 and 4580	SD44	PI 524969	Nitrogen 12-Aug	70.22684
##	10535	4583	4583 and 4584	NC262	PI 531085	Nitrogen 12-Aug	20.12580
##		Rep					
##	10468	Rep2					
##	10484	Rep2					

```
## 10499 Rep2
## 10502 Rep2
## 10519 Rep2
## 10535 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit2 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d2)

#generating ANOVA table
summary(aov(fit2))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 336570   1450.7   16.618 < 2e-16 ***
## Treatment      1    1581    1580.8   18.108 2.58e-05 ***
## Rep            1    2096    2096.2   24.012 1.37e-06 ***
## Genotype:Treatment 224  20498     91.5    1.048    0.339
## Residuals     419   36578     87.3
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the treatments.

```
##Calculating Vprogeny
out2 <- summary(aov(fit2))[[1]]
vprogeny2 <- (out2[1,3] - out2[4,3])/(2*2)
```

```
#Calculating H2
h2_2 = vprogeny2/((vprogeny2 + out2[4,3]/(2*2)))
```

H2 is very high 0.94. This means that canopy coverage is highly influenced by genetic effects more than by environmental effects.

```
####For in the date 14 Aug
d3 <- subset(cc, date %in% "14-Aug")
head(d3)
```

##	Row	Row.Numbers	Genotype	Pedigree	Treatment	date	Canopy_Coverage	Rep
## 8	1001	1001 and 1002	A641	Ames 19311	Nitrogen	14-Aug	8.330186	Rep1
## 18	1003	1003 and 1004	B76	PI 550483	Nitrogen	14-Aug	68.743223	Rep1
## 30	1005	1005 and 1006	A654	PI 587141	Nitrogen	14-Aug	48.227840	Rep1
## 40	1007	1007 and 1008	A239	Ames 23405	Nitrogen	14-Aug	69.157072	Rep1
## 52	1009	1009 and 1010	A682	PI 587143	Nitrogen	14-Aug	60.686364	Rep1
## 71	1011	1011 and 1012	C123	Ames 19313	Nitrogen	14-Aug	4.383795	Rep1

```
tail(d3)
```

##	Row	Row.Numbers	Genotype	Pedigree	Treatment	date	Canopy_Coverage
## 10474	4569	4569 and 4570	C0255	Ames 27107	Nitrogen	14-Aug	20.99052
## 10487	4573	4573 and 4574	N192	PI 550566	Nitrogen	14-Aug	22.83648
## 10490	4575	4575 and 4576	NC312	Ames 27152	Nitrogen	14-Aug	55.09773
## 10501	4577	4577 and 4578	NC292	Ames 27142	Nitrogen	14-Aug	18.50045

```
## 10517 4579 4579 and 4580      SD44  PI 524969  Nitrogen 14-Aug      62.04407
## 10528 4583 4583 and 4584      NC262  PI 531085  Nitrogen 14-Aug      13.25547
##      Rep
## 10474 Rep2
## 10487 Rep2
## 10490 Rep2
## 10501 Rep2
## 10517 Rep2
## 10528 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit3 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d3)
```

```
#generating ANOVA table
summary(aov(fit3))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 378248    1630  15.958 < 2e-16 ***
## Treatment      1      0         0   0.001   0.9766
## Rep           1    4371    4371  42.779 1.79e-10 ***
## Genotype:Treatment 224  26921     120   1.176   0.0794 .
## Residuals     419  42807     102
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the genotypes.

```
##Calculating Vprogeny
out3 <- summary(aov(fit3))[[1]]
vprogeny3 <- (out3[1,3] - out3[4,3])/(2*2)
```

```
#Calculating H2
h2_3 = vprogeny3/((vprogeny3 + out3[4,3]/(2*2)))
```

H2 is very high 0.93 This means that canopy coverage is highly influenced by genetic effects more than by environmental effects.

```
####For in the date 16 Aug
d4 <- subset(cc, date %in% "16-Aug")
head(d4)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 12 1001 1001 and 1002      A641 Ames 19311  Nitrogen 16-Aug      11.573201 Rep1
## 15 1003 1003 and 1004      B76  PI 550483  Nitrogen 16-Aug      71.856793 Rep1
## 28 1005 1005 and 1006      A654 PI 587141  Nitrogen 16-Aug      44.140153 Rep1
## 43 1007 1007 and 1008      A239 Ames 23405  Nitrogen 16-Aug      64.226668 Rep1
## 58 1009 1009 and 1010      A682 PI 587143  Nitrogen 16-Aug      59.013240 Rep1
## 68 1011 1011 and 1012      C123 Ames 19313  Nitrogen 16-Aug      4.432205 Rep1
```

```
tail(d4)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10467 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 16-Aug      17.18911
## 10485 4573 4573 and 4574      N192  PI 550566  Nitrogen 16-Aug      21.17202
## 10488 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 16-Aug      57.69139
## 10508 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 16-Aug      19.02845
## 10514 4579 4579 and 4580      SD44  PI 524969  Nitrogen 16-Aug      57.30935
## 10529 4583 4583 and 4584    NC262  PI 531085  Nitrogen 16-Aug      11.09715
##      Rep
## 10467 Rep2
## 10485 Rep2
## 10488 Rep2
## 10508 Rep2
## 10514 Rep2
## 10529 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit4 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d4)

#generating ANOVA table
summary(aov(fit4))
```

```
##              Df Sum Sq Mean Sq F value Pr(>F)
## Genotype      232 345614      1490  17.863 < 2e-16 ***
## Treatment      1    755        755   9.053 0.00278 **
## Rep            1    8709      8709 104.430 < 2e-16 ***
## Genotype:Treatment 224  25079      112   1.342 0.00524 **
## Residuals      419  34944         83
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the treatments.

```
##Calculating Vprogeny
out4 <- summary(aov(fit4))[[1]]
vprogeny4 <- (out4[1,3] - out4[4,3])/(2*2)
```

```
#Calculating H2
h2_4 = vprogeny4/((vprogeny4 + out4[4,3]/(2*2)))
```

##H2 is very high 0.92 This means that canopy coverage is highly influenced by genetic effects more than by environmental effects.

```
####For in the date 20 Aug
d5 <- subset(cc, date %in% "20-Aug")
head(d5)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 9  1001 1001 and 1002    A641 Ames 19311  Nitrogen 20-Aug      10.179700 Rep1
## 14 1003 1003 and 1004      B76  PI 550483  Nitrogen 20-Aug      74.095592 Rep1
## 27 1005 1005 and 1006    A654  PI 587141  Nitrogen 20-Aug      54.376656 Rep1
```

```
## 41 1007 1007 and 1008      A239 Ames 23405  Nitrogen 20-Aug      73.009362 Rep1
## 51 1009 1009 and 1010      A682  PI 587143  Nitrogen 20-Aug      68.290128 Rep1
## 65 1011 1011 and 1012      C123 Ames 19313  Nitrogen 20-Aug      6.406647 Rep1
```

```
tail(d5)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10464 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 20-Aug      15.60959
## 10478 4573 4573 and 4574      N192  PI 550566  Nitrogen 20-Aug      29.47295
## 10492 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 20-Aug      64.22311
## 10503 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 20-Aug      23.73602
## 10513 4579 4579 and 4580      SD44  PI 524969  Nitrogen 20-Aug      64.44475
## 10524 4583 4583 and 4584    NC262  PI 531085  Nitrogen 20-Aug      16.01849
##      Rep
## 10464 Rep2
## 10478 Rep2
## 10492 Rep2
## 10503 Rep2
## 10513 Rep2
## 10524 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit5 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d5)

#generating ANOVA table
summary(aov(fit5))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 356944   1538.6   17.443 < 2e-16 ***
## Treatment      1    1180    1180.1   13.380 0.000287 ***
## Rep            1     777     776.9    8.808 0.003171 **
## Genotype:Treatment 224  26990    120.5    1.366 0.003343 **
## Residuals     419  36957     88.2
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the genotypes.

```
##Calculating Vprogeny
out5 <- summary(aov(fit5))[[1]]
vprogeny5 <- (out5[1,3] - out5[4,3])/(2*2)

#Calculating H2
h2_5 = vprogeny5/((vprogeny5 + out5[4,3]/(2*2)))
```

H2 is very high 0.92 This means that canopy coverage is highly influenced by genetic effects more than by environmental effects.

```
####For in the date 22 Aug
d6 <- subset(cc, date %in% "22-Aug")
head(d6)
```

```
##      Row   Row.Numbers Genotype   Pedigree Treatment   date Canopy_Coverage Rep
## 4   1001 1001 and 1002      A641 Ames 19311   Nitrogen 22-Aug      19.03728 Rep1
## 16  1003 1003 and 1004       B76  PI 550483   Nitrogen 22-Aug      75.43454 Rep1
## 29  1005 1005 and 1006      A654  PI 587141   Nitrogen 22-Aug      53.39257 Rep1
## 38  1007 1007 and 1008      A239 Ames 23405   Nitrogen 22-Aug      69.77542 Rep1
## 56  1009 1009 and 1010      A682  PI 587143   Nitrogen 22-Aug      62.87168 Rep1
## 67  1011 1011 and 1012      C123 Ames 19313   Nitrogen 22-Aug      11.63524 Rep1
```

```
tail(d6)
```

```
##      Row   Row.Numbers Genotype   Pedigree Treatment   date Canopy_Coverage
## 10470 4569 4569 and 4570      C0255 Ames 27107   Nitrogen 22-Aug      18.43629
## 10479 4573 4573 and 4574       N192  PI 550566   Nitrogen 22-Aug      32.23088
## 10493 4575 4575 and 4576      NC312 Ames 27152   Nitrogen 22-Aug      68.20305
## 10505 4577 4577 and 4578      NC292 Ames 27142   Nitrogen 22-Aug      25.38502
## 10521 4579 4579 and 4580       SD44  PI 524969   Nitrogen 22-Aug      65.04087
## 10531 4583 4583 and 4584      NC262  PI 531085   Nitrogen 22-Aug      18.60264
##      Rep
## 10470 Rep2
## 10479 Rep2
## 10493 Rep2
## 10505 Rep2
## 10521 Rep2
## 10531 Rep2
```

```
### Defining the model to get multiple linear regression for date 6 Jul
fit6 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d6)

#generating ANOVA table
summary(aov(fit6))
```

```
##      Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 319387   1376.7   18.021 < 2e-16 ***
## Treatment       1   2643   2643.2   34.599 8.29e-09 ***
## Rep             1    676    676.0    8.848 0.0031 **
## Genotype:Treatment 224  21771    97.2    1.272 0.0183 *
## Residuals      419  32009    76.4
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the genotypes.

```
##Calculating Vprogeny
out6 <- summary(aov(fit6))[[1]]
vprogeny6 <- (out6[1,3] - out6[4,3])/(2*2)

#Calculating H2
h2_6 = vprogeny6/((vprogeny6 + out6[4,3]/(2*2)))
```

##H2 is very high 0.93 This means that canopy coverage is highly influenced by genetic effects more than by environmental effects.


```
####For in the date 23 Aug
d7 <- subset(cc, date %in% "23-Aug")
head(d7)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 2  1001 1001 and 1002    A641 Ames 19311  Nitrogen 23-Aug      15.53798 Rep1
## 13 1003 1003 and 1004     B76  PI 550483  Nitrogen 23-Aug      75.10952 Rep1
## 32 1005 1005 and 1006    A654  PI 587141  Nitrogen 23-Aug      52.16748 Rep1
## 48 1007 1007 and 1008    A239 Ames 23405  Nitrogen 23-Aug      71.81781 Rep1
## 55 1009 1009 and 1010    A682  PI 587143  Nitrogen 23-Aug      74.07453 Rep1
## 69 1011 1011 and 1012    C123 Ames 19313  Nitrogen 23-Aug      10.83411 Rep1
```

```
tail(d7)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10473 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 23-Aug      20.26747
## 10476 4573 4573 and 4574     N192  PI 550566  Nitrogen 23-Aug      31.52568
## 10498 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 23-Aug      65.63632
## 10511 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 23-Aug      26.28921
## 10518 4579 4579 and 4580     SD44  PI 524969  Nitrogen 23-Aug      67.77127
## 10530 4583 4583 and 4584    NC262  PI 531085  Nitrogen 23-Aug      17.90676
##      Rep
## 10473 Rep2
## 10476 Rep2
## 10498 Rep2
## 10511 Rep2
## 10518 Rep2
## 10530 Rep2
```

```
#### Defining the model to get multiple linear regression for date 6 Jul
fit7 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d7)
```

```
#generating ANOVA table
summary(aov(fit7))
```

```
##      Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232  329716      1421  16.308 < 2e-16 ***
## Treatment       1    3164      3164  36.311 3.69e-09 ***
## Rep            1    3270      3270  37.522 2.08e-09 ***
## Genotype:Treatment 224  23591      105   1.209  0.0502 .
## Residuals      419   36514         87
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the genotypes.

```
##Calculating Vprogeny
out7 <- summary(aov(fit7))[[1]]
vprogeny7 <- (out7[1,3] - out7[4,3])/(2*2)
```

```

#Calculating H2
h2_7 = vprogeny7/((vprogeny7 + out7[4,3]/(2*2)))

##H2 is very high 0.93 This means that canopy coverage is highly influenced by genetic effects more than
environmental effects

####For in the date 26 Aug
d8 <- subset(cc, date %in% "26-Aug")
head(d8)

```

```

##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 6   1001 1001 and 1002    A641 Ames 19311  Nitrogen 26-Aug      21.37887 Rep1
## 23  1003 1003 and 1004     B76  PI 550483  Nitrogen 26-Aug      72.06732 Rep1
## 25  1005 1005 and 1006    A654  PI 587141  Nitrogen 26-Aug      55.76904 Rep1
## 37  1007 1007 and 1008    A239 Ames 23405  Nitrogen 26-Aug      74.37770 Rep1
## 57  1009 1009 and 1010    A682  PI 587143  Nitrogen 26-Aug      70.56410 Rep1
## 64  1011 1011 and 1012    C123 Ames 19313  Nitrogen 26-Aug      16.30953 Rep1

```

```
tail(d8)
```

```

##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10471 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 26-Aug      15.53898
## 10483 4573 4573 and 4574     N192  PI 550566  Nitrogen 26-Aug      36.37043
## 10491 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 26-Aug      76.05566
## 10506 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 26-Aug      32.15958
## 10520 4579 4579 and 4580     SD44  PI 524969  Nitrogen 26-Aug      73.31089
## 10527 4583 4583 and 4584    NC262  PI 531085  Nitrogen 26-Aug      21.15264
##      Rep
## 10471 Rep2
## 10483 Rep2
## 10491 Rep2
## 10506 Rep2
## 10520 Rep2
## 10527 Rep2

```

```

#### Defining the model to get multiple linear regression for date 6 Jul
fit8 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d8)

```

```

#generating ANOVA table
summary(aov(fit8))

```

```

##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 314911    1357   15.905 < 2e-16 ***
## Treatment      1    4076    4076   47.756 1.81e-11 ***
## Rep            1    1273    1273   14.919 0.00013 ***
## Genotype:Treatment 224  23682     106    1.239 0.03166 *
## Residuals     418  35674      85
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

#As the p-value is less than the significance level 0.05, we can conclude that there are significant di

##Calculating Vprogeny

```
out8 <- summary(aov(fit8))[[1]]
vprogeny8 <- (out8[1,3] - out8[4,3])/(2*2)
```

#Calculating H2

```
h2_8 = vprogeny8/((vprogeny8 + out8[4,3]/(2*2)))
```

##H2 is very high 0.92 This means that canopy coverage is highly influenced by genetic effects more tha

####For in the date 30 Aug

```
d9 <- subset(cc, date %in% "30-Aug")
head(d9)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 10 1001 1001 and 1002    A641 Ames 19311  Nitrogen 30-Aug      9.945542 Rep1
## 22 1003 1003 and 1004     B76  PI 550483  Nitrogen 30-Aug     75.113953 Rep1
## 31 1005 1005 and 1006    A654  PI 587141  Nitrogen 30-Aug     50.838188 Rep1
## 47 1007 1007 and 1008    A239 Ames 23405  Nitrogen 30-Aug     79.089834 Rep1
## 54 1009 1009 and 1010    A682  PI 587143  Nitrogen 30-Aug     90.333319 Rep1
## 62 1011 1011 and 1012    C123 Ames 19313  Nitrogen 30-Aug     10.285431 Rep1
```

```
tail(d9)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10469 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 30-Aug      5.547609
## 10477 4573 4573 and 4574     N192  PI 550566  Nitrogen 30-Aug     26.156860
## 10497 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 30-Aug     63.367000
## 10500 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 30-Aug     23.513413
## 10512 4579 4579 and 4580     SD44  PI 524969  Nitrogen 30-Aug     62.546048
## 10532 4583 4583 and 4584    NC262  PI 531085  Nitrogen 30-Aug     15.172438
##      Rep
## 10469 Rep2
## 10477 Rep2
## 10497 Rep2
## 10500 Rep2
## 10512 Rep2
## 10532 Rep2
```

Defining the model to get multiple linear regression for date 6 Jul

```
fit9 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
           data=d9)
```

#generating ANOVA table

```
summary(aov(fit9))
```

```
##      Df Sum Sq Mean Sq F value  Pr(>F)
## Genotype      232 364874    1573  14.113 < 2e-16 ***
## Treatment       1    937     937   8.404 0.00394 **
## Rep             1   9057   9057  81.272 < 2e-16 ***
## Genotype:Treatment 224  31197     139   1.250 0.02650 *
```

```
## Residuals          419  46692      111
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

##As the p-value is less than the significance level 0.05, we can conclude that there are significant di

##Calculating Vprogeny

```
out9 <- summary(aov(fit9))[[1]]
vprogeny9 <- (out9[1,3] - out9[4,3])/(2*2)
```

##Calculating H2

```
h2_9 = vprogeny9/((vprogeny9 + out9[4,3]/(2*2)))
```

##H2 is very high 0.91 This means that canopy coverage is highly influenced by genetic effects more tha

####For in the date 1 Sep

```
d10 <- subset(cc, date %in% "1-Sep")
head(d10)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 3   1001 1001 and 1002    A641 Ames 19311  Nitrogen 1-Sep      15.189726 Rep1
## 17  1003 1003 and 1004     B76  PI 550483  Nitrogen 1-Sep      74.382934 Rep1
## 26  1005 1005 and 1006    A654  PI 587141  Nitrogen 1-Sep      48.696939 Rep1
## 44  1007 1007 and 1008    A239 Ames 23405  Nitrogen 1-Sep      71.570800 Rep1
## 59  1009 1009 and 1010    A682  PI 587143  Nitrogen 1-Sep      76.387863 Rep1
## 63  1011 1011 and 1012    C123 Ames 19313  Nitrogen 1-Sep       7.711603 Rep1
```

```
tail(d10)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10466 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 1-Sep        6.362997
## 10480 4573 4573 and 4574     N192  PI 550566  Nitrogen 1-Sep       25.053398
## 10495 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 1-Sep       67.485713
## 10510 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 1-Sep       21.563958
## 10523 4579 4579 and 4580     SD44  PI 524969  Nitrogen 1-Sep       70.742389
## 10534 4583 4583 and 4584    NC262  PI 531085  Nitrogen 1-Sep       15.371834
##      Rep
## 10466 Rep2
## 10480 Rep2
## 10495 Rep2
## 10510 Rep2
## 10523 Rep2
## 10534 Rep2
```

Defining the model to get multiple linear regression for date 6 Jul

```
fit10 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
            data=d10)
```

#generating ANOVA table

```
summary(aov(fit10))
```

```
##      Df Sum Sq Mean Sq F value    Pr(>F)
```

```
## Genotype          232 441587      1903 14.332 < 2e-16 ***
## Treatment          1   2971      2971 22.372 3.07e-06 ***
## Rep                1   3309      3309 24.919 8.79e-07 ***
## Genotype:Treatment 224 32509      145  1.093    0.22
## Residuals          419 55645      133
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

##As the p-value is less than the significance level 0.05, we can conclude that there are significant di.

##Calculating Vprogeny

```
out10 <- summary(aov(fit10))[[1]]
vprogeny10 <- (out10[1,3] - out10[4,3])/(2*2)
```

##Calculating H2

```
h2_10 = vprogeny10/((vprogeny10 + out10[4,3]/(2*2)))
```

##H2 is very high 0.93 This means that canopy coverage is highly influenced by genetic effects more than

####For in the date 3 Sep

```
d11 <- subset(cc, date %in% "3-Sep")
head(d11)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 7   1001 1001 and 1002    A641 Ames 19311  Nitrogen 3-Sep      13.53673 Rep1
## 19  1003 1003 and 1004     B76  PI 550483  Nitrogen 3-Sep      61.73542 Rep1
## 36  1005 1005 and 1006    A654  PI 587141  Nitrogen 3-Sep      41.31357 Rep1
## 42  1007 1007 and 1008    A239 Ames 23405  Nitrogen 3-Sep      51.48036 Rep1
## 53  1009 1009 and 1010    A682  PI 587143  Nitrogen 3-Sep      49.44327 Rep1
## 72  1011 1011 and 1012    C123 Ames 19313  Nitrogen 3-Sep      10.07274 Rep1
```

```
tail(d11)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10465 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 3-Sep       2.541246
## 10481 4573 4573 and 4574     N192  PI 550566  Nitrogen 3-Sep      20.475784
## 10494 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 3-Sep      50.602426
## 10504 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 3-Sep      21.216328
## 10522 4579 4579 and 4580     SD44  PI 524969  Nitrogen 3-Sep      55.747890
## 10525 4583 4583 and 4584    NC262  PI 531085  Nitrogen 3-Sep      14.445922
##      Rep
## 10465 Rep2
## 10481 Rep2
## 10494 Rep2
## 10504 Rep2
## 10522 Rep2
## 10525 Rep2
```

Defining the model to get multiple linear regression for date 6 Jul

```
fit11 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
            data=d11)
```

#generating ANOVA table

```
summary(aov(fit11))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 260177     1121  15.232 < 2e-16 ***
## Treatment      1    801      801  10.882 0.00105 **
## Rep            1   6455    6455  87.678 < 2e-16 ***
## Genotype:Treatment 224  20558      92   1.247 0.02790 *
## Residuals      419  30848      74
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the treatments.

##Calculating Vprogeny

```
out11 <- summary(aov(fit11))[[1]]
vprogeny11 <- (out11[1,3] - out11[4,3])/(2*2)
```

##Calculating H2

```
h2_11 = vprogeny11/((vprogeny11 + out11[4,3]/(2*2)))
```

##H2 is very high 0.92 This means that canopy coverage is highly influenced by genetic effects more than by the environment.

####For in the date 5 Sep

```
d12 <- subset(cc, date %in% "5-Sep")
head(d12)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage  Rep
## 11 1001 1001 and 1002    A641 Ames 19311  Nitrogen 5-Sep      8.183619 Rep1
## 21 1003 1003 and 1004    B76  PI 550483  Nitrogen 5-Sep     67.182663 Rep1
## 34 1005 1005 and 1006    A654  PI 587141  Nitrogen 5-Sep     37.853154 Rep1
## 46 1007 1007 and 1008    A239 Ames 23405  Nitrogen 5-Sep     61.086586 Rep1
## 49 1009 1009 and 1010    A682  PI 587143  Nitrogen 5-Sep     61.464023 Rep1
## 61 1011 1011 and 1012    C123 Ames 19313  Nitrogen 5-Sep     10.114035 Rep1
```

```
tail(d12)
```

```
##      Row  Row.Numbers Genotype  Pedigree Treatment  date Canopy_Coverage
## 10475 4569 4569 and 4570    C0255 Ames 27107  Nitrogen 5-Sep      3.814582
## 10486 4573 4573 and 4574     N192  PI 550566  Nitrogen 5-Sep     15.824509
## 10496 4575 4575 and 4576    NC312 Ames 27152  Nitrogen 5-Sep     50.597114
## 10509 4577 4577 and 4578    NC292 Ames 27142  Nitrogen 5-Sep     16.482220
## 10515 4579 4579 and 4580     SD44  PI 524969  Nitrogen 5-Sep     51.163657
## 10533 4583 4583 and 4584    NC262  PI 531085  Nitrogen 5-Sep      9.914150
##      Rep
## 10475 Rep2
## 10486 Rep2
## 10496 Rep2
## 10509 Rep2
## 10515 Rep2
## 10533 Rep2
```

Defining the model to get multiple linear regression for date 6 Jul

```
fit12 <- lm(Canopy_Coverage ~ Genotype + Treatment + Genotype:Treatment + Rep,
            data=d12)
```

```
#generating ANOVA table
summary(aov(fit12))
```

```
##              Df Sum Sq Mean Sq F value    Pr(>F)
## Genotype      232 343152     1479   17.036 < 2e-16 ***
## Treatment      1    369       369    4.251  0.0399 *
## Rep            1   5681     5681   65.430 6.56e-15 ***
## Genotype:Treatment 224  21211        95    1.091  0.2250
## Residuals     419  36379        87
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#As the p-value is less than the significance level 0.05, we can conclude that there are significant differences between the treatments.

```
##Calculating Vprogeny
out12 <- summary(aov(fit12))[[1]]
vprogeny12 <- (out12[1,3] - out12[4,3])/(2*2)
```

```
##Calculating H2
h2_12 = vprogeny12/((vprogeny12 + out12[4,3]/(2*2)))
```

##H2 is very high 0.94 This means that canopy coverage is highly influenced by genetic effects more than the environment.

```
#Creating a table with dates and h2 values
datest=table(cc$date)
datest
```

```
##
## 1-Sep 12-Aug 14-Aug 16-Aug 20-Aug 22-Aug 23-Aug 26-Aug 3-Sep 30-Aug 5-Sep
##   878   878   878   878   878   878   878   877   878   878   878
## 6-Jul
##   878
```

```
typeof(datest)
```

```
## [1] "integer"
```

```
mytable = data.frame(date=c("06-Jul","12-Aug","14-Aug","16-Aug","20-Aug","22-Aug","23-Aug","26-Aug","30-Aug","05-Sep"),
                      h2=c(h2_1,h2_2,h2_3,h2_4,h2_5,h2_6,h2_7,h2_8,h2_9,h2_10,h2_11,h2_12))
mytable[order(mytable$date),]
```

```
##      date      h2
## 10 01-Sep 0.9237515
## 11 03-Sep 0.9181626
## 12 05-Sep 0.9359811
## 1  06-Jul 0.8864210
## 2  12-Aug 0.9369237
## 3  14-Aug 0.9262842
## 4  16-Aug 0.9248455
```

```
## 5 20-Aug 0.9216848
## 6 22-Aug 0.9294018
## 7 23-Aug 0.9258939
## 8 26-Aug 0.9221132
## 9 30-Aug 0.9114451
```

```
mytable
```

```
##      date      h2
## 1 06-Jul 0.8864210
## 2 12-Aug 0.9369237
## 3 14-Aug 0.9262842
## 4 16-Aug 0.9248455
## 5 20-Aug 0.9216848
## 6 22-Aug 0.9294018
## 7 23-Aug 0.9258939
## 8 26-Aug 0.9221132
## 9 30-Aug 0.9114451
## 10 01-Sep 0.9237515
## 11 03-Sep 0.9181626
## 12 05-Sep 0.9359811
```

```
class(mytable)
```

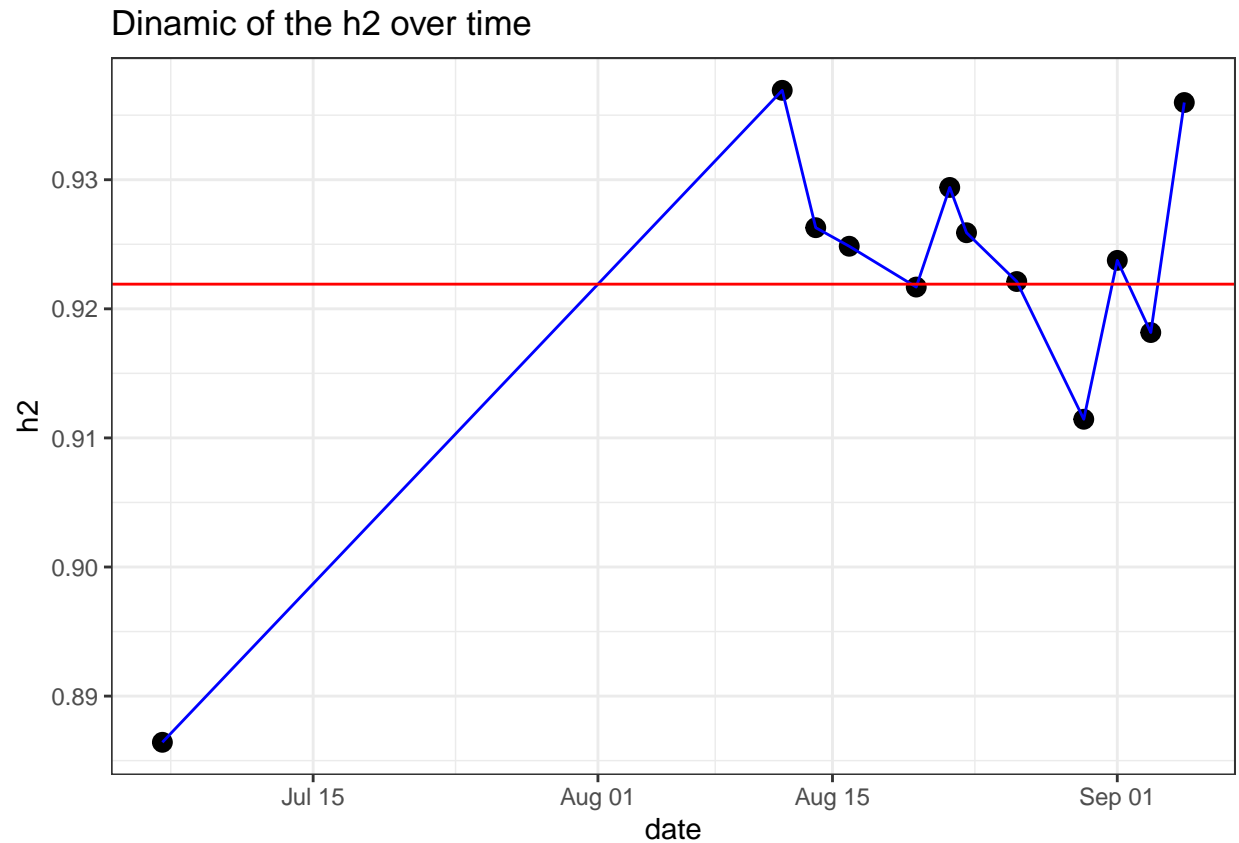
```
## [1] "data.frame"
```

```
mytable$date=as.Date(mytable$date, "%d-%b")
```

```
#Visualize the data
```

```
library(ggplot2)
```

```
ggplot(mytable,aes(date,h2,color="h2 dinamic"))+
  geom_point(colour="black", size=3)+ geom_line(colour="blue")+ labs(title="Dinamic of the h2 over time")
  theme_bw()+geom_hline(yintercept=mean(mytable$h2),color="red")
```

#My Hypothesis is that according with plant physiology, the more older the crop, the more canopy covera

#According to the graph, the h2 value is increasing over time. We can see that the lowest value is in t

#We can see that except for the first data point, the h2 is almost constant and close to the h2 mean ov

#The heritability can change over time because the variance of the genetic values can change due to env

#At the first dates, maybe the plant is in a growing stage in which do not express a lot the genes but

#According to the graph, I reject the null hypothesis. The h2 dynamic over time in the graph is showing