(hr. (et 10-1)

Show That in The ANOVA decomposition, The "cross-term" between the 2nd and 3rd terms is zero. I.e. show that  $\xi(Y_i, -Y_i)(Y_i, -Y_i, +Y_i) = 0$ 

Hint: This requires only 2 or 3 lines of algebra; and don't forget That  $\geq (Y_{ij} - Y_{..}) = 0$ .

$$= \underbrace{\xi_{i}}_{ij} (Y_{i}; -Y_{i.}) Y_{ij} - \underbrace{\xi_{i}}_{ij} (Y_{i}; -Y_{i.}) (Y_{ii}) - \underbrace{\xi_{i}}_{ij} (Y_{i}; -Y_{i.}) Y_{ij} + \underbrace{\xi_{i}}_{ij} (Y_{i}; -Y_{i.}) Y_{ii}$$

$$= \underbrace{\xi_{i}}_{ij} (Y_{i}; -Y_{i.}) \underbrace{\xi_{i}}_{ij} (Y_{$$

## hur-lectio-2

For The model  $\forall ij = M + di + \beta j + Eij$  (i = 1 ... a, j = 1, ..., b)Show That  $E[MS_{TV}] = \sigma_e^2 + b = \sum_{i=1}^{n} (a_i - \overline{a}_i)^2$ Hint: As always, use The model as a p, so That you can use results That we have proven already. For example, you may use, without prosf. That  $E[(Ei, -E, )^2] = \frac{1}{5}(1 - \frac{1}{4})\sigma_e^2$ 

 $= b \stackrel{>}{>} (\alpha_{1} - \overline{\alpha})^{2} + b a \stackrel{1}{>} (1 - \frac{1}{a}) \sigma_{e}^{2} = b \stackrel{>}{>} (\alpha_{1} - \overline{\alpha})^{2} + (a - i) \sigma_{e}^{2}.$ (a-i)

 $: E[MSTV] = E\left[\frac{SSTV}{\alpha-1}\right] = O_e^2 + \frac{b}{\alpha-1} \cdot \left[\alpha_i - \overline{\alpha}\right]^2$ 

```
hw-let 10-3
# In problem 4.3,
# In an RCBD design with one treatment and one block factor, we know how to test
# the treatment effect (and the block effect). The tests are based on an F-test,
# where F is MS_treatment/MSE (or MS_block/MSE). However, I have also mentioned that
# those tests are equivalent to another set of F tests which are based on the SSE's
# obtained from a full model and two reduced models. Let's confirm !
# For the data in problem 4.3, do the following (all by R):
# a) Write code to produce the anova table for the full model.
# Record/save the value of SSE.
\# Allphao, save the F-ratios for the treatment and block effects in the full anova
table.
# b) Write code to develop the reduced model that excludes the treatment factor.
# Record/save the value of SSE.
# c) Write code to develop the reduced model that excludes the block factor.
# Record/save the value of SSE.
# d) Show that by computing the appropriate ratios of SSE in parts a, b, and c,
# you can compute the same F ratios you saved in the anova table of the full model
# in part a.
y.matrix = matrix(c(
                                              # rows = treatment, col = block
   73, 68, 74, 71, 67,
   73, 67, 75, 72, 70,
   75, 68, 78, 73, 68,
   73, 71, 75, 75, 69), nrow=4,byrow=T)
 y = t(y.matrix)
y = as.vector(y)
  a = 4
  b = 5
  A = as.factor(rep(c(1:a),each=b))
  B = as.factor(rep(c(1:b),a))
# a)
  lm.1 = lm(y \sim A + B)
  temp = summary.aov(lm.1)
  temp
  SSE_mu_tau_beta = temp[[1]][3,2]
#
              Df Sum Sq Mean Sq F value
                                           Pr(>F)
# A
                  12.95
                           4.32
                                  2.376
                                            0.121
# B
               4 157.00
                          39.25
                                  21.606 2.06e-05 ***
# Residuals
                  21.80
                           1.82
              12
```

```
# b)
  lm.reducedB = lm(y \sim B)
  tempB = summary.aov(lm.reducedB)
  tempB
  SSE_mu_beta = tempB[[1]][2,2]
#
              Df Sum Sq Mean Sq F value
                                          Pr(>F)
# B
               4 157.00
                          39.25
                                  16.94 1.95e-05 ***
# Residuals
              15 34.75
                           2.32
# c)
  lm.reducedA = lm(y \sim A)
  tempA = summary.aov(lm.reducedA)
  tempA
  SSE_mu_tau = tempA[[1]][2,2]
              Df Sum Sq Mean Sq F value Pr(>F)
                         4.317
               3 12.95
                                 0.386 0.764
# A
              16 178.80
                         11.175
# Residuals
# d)
  FA_numer = (SSE_mu_beta - SSE_mu_tau_beta) / (a-1)
 FA_denom = SSE_mu_tau_beta/((a-1)*(b-1))
 FA = FA_numer/FA_denom
                                                            # 2.376147 = in part a
 FB_numer = (SSE_mu_tau - SSE_mu_tau_beta) / (b-1)
 FB_denom = SSE_mu_tau_beta/((a-1)*(b-1))
 FB = FB_numer/FB_denom
                                                            # 21.6055 = in part a
# Although I did not ask you to do this, you can see that the p-values are
# the same too:
 pf(FA, df1=a-1, df2=(a-1)*(b-1), 0, lower.tail = FALSE) # 0.1211445 = in part a
 pf(FB, df1=b-1, df2=(a-1)*(b-1), 0, lower.tail = FALSE) # 2.059181e-05 = in part a
```

## hr-lect 11-1

## In problem 4.3,

- a) perform anova to compute a p-value from an F test of whether the 4 chemicals have an effect on strength, treating the data as if they were collected in a CRD. by hand
- b) Repeat part a, but with an RCBD, with Bolts as the block factor. by hand.
- c) Produce two residual plots: residuals versus y\_hat, and residuals vs. y, (just to see why the latter is not recommended), for the residuals from the CRD design.
- d) Repeat part c, but for RCBD.
- e) Make qq plots for the CRD and RCBD residuals. Comment

```
y = matrix(c(
                            # rows = treatment, col = block
73, 68, 74, 71, 67,
73, 67, 75, 72, 70,
75, 68, 78, 73, 68,
73, 71, 75, 75, 69), nrow=4,byrow=T)
# a) CRD
a = 4
n = 5
N = n*a
means = apply(y, 1, mean)
                             # 70.6 71.4 72.4 72.6
 vars = apply(y, 1, var)
                          # 9.3 9.3 19.3 6.8 Note: unequal vars!
 grand.mean = mean(means)
SS_{treatment} = n*sum((means - grand.mean)^2)
SS_E = (n-1)*sum(vars)
MS treatment = SS treatment/(a-1)
                                                # 4.316667
MS_E = SS_E/(N-a)
                                           # 11.175
```

 $F_{\text{obs}} = MS_{\text{treatment/MS}} = \#0.3862789$ 

 $pf(F\_obs,\,df1{=}a{-}1,\,df2{=}N{-}a,\,0,\,lower.tail=FALSE)\quad \#\ 0.764377$ 

# Cannot reject H0 (that all 4 tau's are zero) in favor of H1.

```
#b) RCBD
```

a = 4

b = 5

row.means = apply(y,1,mean) # 70.6 71.4 72.4 72.6 col.means = apply(y,2,mean) # 73.50 68.50 75.50 72.75 68.50

 $SS_{treatment} = b*sum((row.means - grand.mean)^2)$ 

 $SS_block = a*sum((col.means - grand.mean)^2)$ 

resid2 = (t(t(y - row.means) - col.means + grand.mean)) # Note t(t())

 $SS_E = sum( resid2^2)$ 

MS\_treatment = SS\_treatment/(a-1) # 4.316667 MS\_block = SS\_block/(b-1) # 39.25 MS\_E = SS\_E/((a-1)\*(b-1)) # 1.816667 F\_obs = MS\_treatment/MS\_E # 2.376147

 $pf(F_obs, df1=a-1, df2=(a-1)*(b-1), 0, lower.tail = FALSE) # 0.1211445$ 

# Cannot reject H0 (that all 4 tau's are zero) in favor of H1.

```
# The conclusion to part a is this: assuming the design is CRD
# (when we know it's not), and applying an anova F-test, does NOT
# allow us to find the treatment effect.
# part b tells us that doing an F test that acknowledges
# the existing RCBD design is still not capable of finding the
# treatement effect, but it does have a lower p-value.
# c) residual for CRD
 resid.crd = y - row.means
 # residuals vs. y_hat:
 plot(rep(row.means,5), resid.crd, xlab="y_hat", ylab="residual")
 abline(h=0)
 # residuals vs. y: (not recommended)
 plot(as.vector(y), resid.crd, xlab="y", ylab="residual")
 abline(h=0)
 # The residual plots look adequate, except for the 2nd one
 # which is why it is not recommended.
# d) residual for RCBD
 resid.rcbd = t(t(y - row.means) - col.means + grand.mean)
 plot(rep(row.means,5), as.vector(resid.rcbd), xlab="row.means",ylab="residual")
 abline(h=0)
 plot(as.vector(y), as.vector(resid.rcbd), xlab="row.means", ylab="residual")
 abline(h=0)
 # This time both residual plots look adequate.
# e) qq plots fo CRD and RCBD
 qqnorm(resid.crd)
                           # consistent with normal
 ggnorm(resid.rcbd)
                           # consistent with normal.
```

Consider The following data from an LSD: Above, we proved That more qually. (is The middle 51' Yim = 3 ym ( 51' Vin = PYm) Viju = / Y111

Y221 Y232 Y331 Y312 Y323/ Y3. Similarly, it can be shown That Y .. 1 Y .. 2 Y .. 3 E Y.j. = & Y... = 3 Y ... Here, Show That & Yish = & Yi. unvestricted restricted and Sightish = & Yith  $\frac{2}{3} \frac{1}{10} \frac{1}{10} = \frac{1}{10} + \frac{1}{122} + \frac{1}{133} = \frac{1}{10} = \frac{3}{10} = \frac$ Yn Yn Yn => 5 Ynk = 3 Ynk FYI: Similarly, Six Yuk = 5 Yig. . So, Y = 5 Yin Y ... = = = > Yi..... The previous of previous problem

harled 11-2

## hn-leit 11-4

To estimate The params of The LSD model, we are supposed to differentiate SSE wirt. M. Xi, Ti, Bh, and set The result to posses as Eight = 51 ( Yi)h-M-Xi-Ti-Bh)2

Because of The restricted sum, we can't quite differentiate this.

a) First, write out The 5' in SSE

For The specific LSD 

Viju = Y221 Y232 Y213

Y331 Y312 Y323

 $SSE = (\gamma_{111} - \mu - \alpha_1 - \gamma_1 - \beta_1)^2 + (\gamma_{122} - \mu - \alpha_1 - \gamma_2 - \beta_2)^2 + (\gamma_{123} - \mu - \alpha_1 - \gamma_3 - \beta_3)^2$   $(\gamma_{221} - \mu - \alpha_2 - \gamma_2 - \beta_1)^2 + (\gamma_{232} - \mu - \alpha_2 - \gamma_3 - \beta_2)^2 + (\gamma_{213} - \mu - \alpha_2 - \gamma_1 - \beta_3)^2$   $(\gamma_{231} - \mu - \alpha_3 - \gamma_3 - \beta_1)^2 + (\gamma_{312} - \mu - \alpha_3 - \gamma_1 - \beta_2)^2 + (\gamma_{312} - \mu - \alpha_3 - \gamma_2 - \beta_3)^2$ 

b) Find 2/20, 1/2,2,7 =0

 $\frac{\partial SSE}{\partial T_{1}}: \left(Y_{111} - M - \alpha_{1} - T_{1} - \beta_{1}\right) + \left(Y_{213} - M - \alpha_{2} - T_{1} - \beta_{3}\right) + \left(Y_{312} - M - \alpha_{3} - T_{1} - \beta_{2}\right)$   $\left(Y_{111} + Y_{213} + Y_{312}\right) - 3M - \left(\alpha_{1} + \alpha_{2} + \alpha_{3}\right) - 3T_{1} - \left(\beta_{1} + \beta_{2} + \beta_{3}\right) = 0.$ 

 $2 \left( \frac{2}{ijk} \right) \left( \frac{1}{1} - 3\hat{\mu} - 3\hat{\tau}_1 - \hat{\alpha}_1 - \hat{\beta}_1 \right) = 0$ 

hur le + 12-1

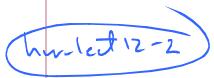
Show that SSrow as defined by 50 (\(\frac{1}{1...-\frac{1}{1...}}\)2 is equal to \$ 2 11. - fr 12. You may use The specific LS shown ]

55 = \(\frac{1}{1}\left(\frac{1}{1}\ldots\frac{1}\ldots\frac{1}{1}\ldots\frac{1}{1}\ldots\frac{1}\ldots\frac{1}{1}\ldots\frac{1}{1}\ldots\frac{1}{1}\ldots\frac{1}{1}\ldots\frac

$$= (\overline{Y_{1}}, -\overline{Y_{1}}, -\overline{Y_{1$$

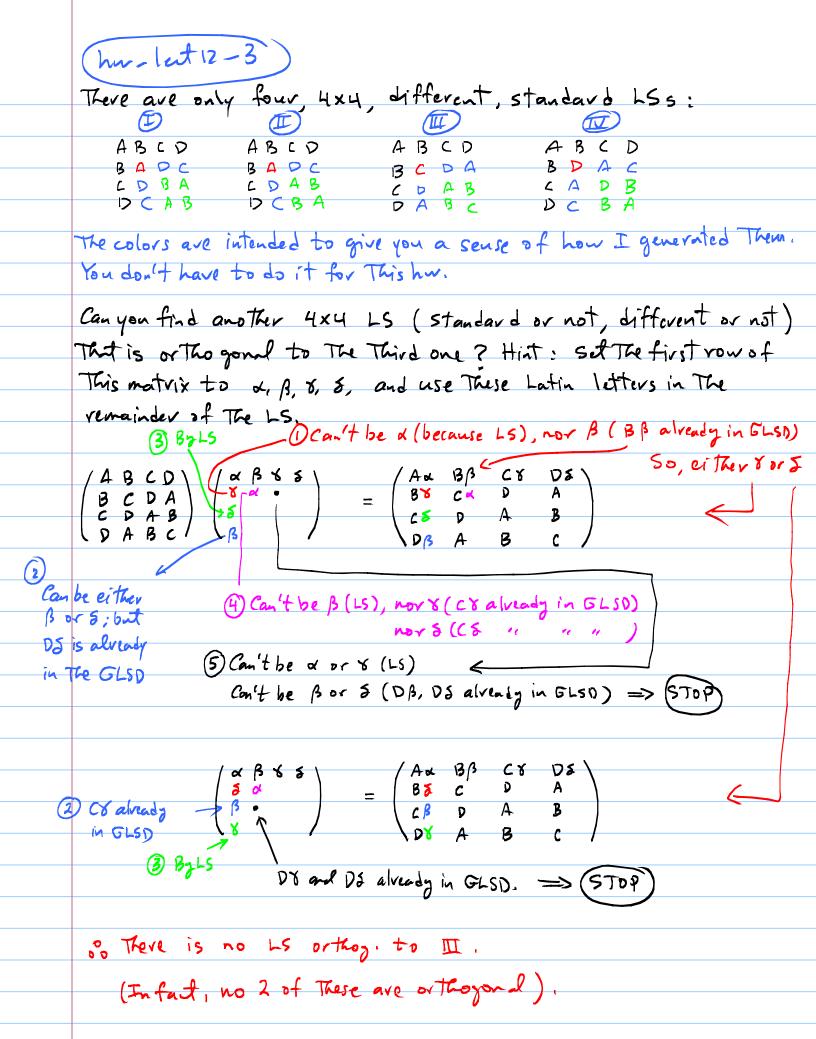
 $=3\frac{3}{5}\left(\frac{2}{4i..}+\frac{2}{4i..}-2\frac{1}{4i..}\right)=3\left(\frac{3}{5}\frac{2}{4i..}+3\frac{2}{4i..}-2\frac{2}{4i..}\right)$ =3 = 7 in +3 7 ... -6 7 ... =3 [ 5 ( 1 /i-)2-3 ( 4 /···)2]

= 1 2 Yin - 1 7 m



```
\# For the data in problem 4.22, perform the appropriate ANOVA analysis; report SSA,
SSB, $SC, SST, SSE, the F ratios, and the corresponding p-values a) by hand, b) by R.
  rm(list=ls(all=TRUE))
  na = 5
  nb = 5
  nc = 5
  p = na
  y.m = matrix(nrow=na,ncol=nb)
                                      # rows = factor, col = replicates
  y.m[1,]=c(8,7,1,7,3)
  y.m[2,]=c(11,2,7,3,8)
  y.m[3,]=c(4,9,10,1,5)
  y.m[4,]=c(6,8,6,6,10)
  y.m[5,]=c(4,2,3,8,8)
  y = as.vector(t(y.m))
  A = as.factor(c(rep(1:na,each=nb)))
                                                  # row-var = Batch (block)
  B = as.factor(rep(c(1:nb),na))
                                                  # col-var = Day (block)
  C = as.factor(c(1,2,4,3,5,
                                                  # ingredient
                    3,5,1,4,2,
                    2,1,3,5,4,
                    4,3,5,2,1,
                    5,4,2,1,3))
  cbind(A,B,C,y)
                      # you can confirm that the data in R is same as in problem.
  y... = sum(y)
  yi. = c(sum(y[A==1]), sum(y[A==2]), sum(y[A==3]), sum(y[A==4]), sum(y[A==5]))
  y.j. = c(sum(y[B==1]), sum(y[B==2]), sum(y[B==3]), sum(y[B==4]), sum(y[B==5]))
  y..k = c(sum(y[C==1]), sum(y[C==2]), sum(y[C==3]), sum(y[C==4]), sum(y[C==5]))
  SSA = sum((yi..)^2)/p - (y...)^2/p^2
  SSB = sum((y.j.)^2)/p - (y...)^2/p^2
  SSC = sum((y..k)^2)/p - (y...)^2/p^2
  SST = sum(y^2) - (y...)^2/p^2
  SSE = SST - (SSA + SSB + SSC)
```

```
MSA = SSA/(p-1)
   MSB = SSB/(p-1)
   MSC = SSC/(p-1)
   MSE = SSE/((p-2)*(p-1))
   FA = MSA/MSE
   FB = MSB/MSE
   FC = MSC/MSE
   pf(FA,p-1,(p-2)*(p-1),lower.tail=F)
                                           # 0.3476182
   pf(FB,p-1,(p-2)*(p-1),lower.tail=F)
                                           # 0.4550143
   pf(FC,p-1,(p-2)*(p-1),lower.tail=F)
                                           # 0.0004876512
# By R
   lm.1 = lm(y \sim A + B + C)
   summary.aov(lm.1)
#
              Df Sum Sq Mean Sq F value
                                           Pr(>F)
# A
                                   1.235 0.347618
                 15.44
                           3.86
# B
               4
                  12.24
                           3.06
                                   0.979 0.455014
# C
               4 141.44
                           35.36
                                 11.309 0.000488 ***
# Residuals
              12
                  37.52
                           3.13
# All the values in the table are the same as those computed by hand, above.
# The conclusion of the study is that there is no evidence that A and B have an
effect,
# but C (the ingredient) does. Technically, we should look at residual plots to
# check all the assumptions.
```



```
4.36
```

```
#4.29 (7th ed.) = 4.36 (8th ed.)
rm(list=ls(all=TRUE))
p = 4
 y.m = matrix(nrow=p,ncol=p)
 y.m[1,]=c(11, 10, 14, 8)
 y.m[2,]=c(8, 12, 10, 12)
 y.m[3,]=c(9, 11, 7, 15)
 y.m[4,]=c(9, 8, 18, 6)
 y = as.vector(t(y.m))
 A = as.factor(c(rep(1:p,each=p)))
                                       # row-factor (Order of Assembly).
                                   # Greek factor (treatment)
 B = as.factor(c(3, 2, 4, 1,
            2, 3, 1, 4,
            1, 4, 2, 3,
            4, 1, 3, 2))
 C = as.factor(rep(c(1:p),p))
                                    # col-factor (Operator).
 D = as.factor(c(2, 3, 4, 1,
            1, 4, 3, 2,
            4, 1, 2, 3,
                              # Latin factor
            3, 2, 1, 4))
                                    # Always, visually confirm data are correct. summary.aov(lm(y~A+B+C+D))
 cbind(A,B,C,D,y)
#
         Df Sum Sq Mean Sq F value Pr(>F)
           3 0.5 0.17 0.018 0.996
# A
# B
           3 95.5 31.83 3.473 0.167
          3 19.0 6.33 0.691 0.616
# C
# D
          3 7.5 2.50 0.273 0.843
# Residuals 3 27.5 9.17
# All the p-values are relatively large, and so there is no evidence from data
# that any of the 4 factors have an effect on the response. But, it's worth noting
# that all of the p-values are based on only 3 degrees of freedom. That's almost
# like having a sample of size 3! Would you trust the results of that data?!
```

This document was created with Win2PDF available at <a href="http://www.win2pdf.com">http://www.win2pdf.com</a>. The unregistered version of Win2PDF is for evaluation or non-commercial use only. This page will not be added after purchasing Win2PDF.