

Experimental Design: Blocking

- A study is conducted to measure effect of drug on locomotor activity in hyperactive children
- Between-subjects design:
 - 3 groups differ in drug dosage: zero, low, & high
- Dependent variable: locomotor activity
 - measured for fixed interval after drug administration
- Before study, measure baseline locomotor activity in each subject
 - baseline measure used as a **blocking variable**

Randomized Block Design

- 4 blocks of 12 Ss created using baseline locomotor activity measure
- subjects in each block assigned randomly to drug dose condition

		Drug Dose		
		zero	low	high
Block (Baseline Locomotor Activity)	low	4	4	4
	medium	4	4	4
	high	4	4	4
	very high	4	4	4

Do blocks differ from each other?

```
> with(theData, tapply(activity, block, mean));

      low      med      high      very
5.153699  8.007714 10.287328 13.818771

> summary(aov(activity ~ block, data=theData) );

              Df Sum Sq Mean Sq F value Pr(>F)
block          3  483.1  161.02  106.3 <2e-16 ***
Residuals    44   66.6    1.51
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Effect of blocking factor on $SS_{\text{residuals}}$

```
> aov.1<-aov(y~drug,data=theData)
> aov.2<-aov(y~block+drug+block:drug,data=theData)
> summary(aov.1)
```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
drug	2	37.1	18.539	2.173	0.126
Residuals	45	384.0	8.532		

$SS_{\text{total}} = 421$

```
> summary(aov.2)
```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
block	3	129.84	43.28	6.964	0.000818 ***
drug	2	37.08	18.54	2.983	0.063294 .
block:drug	6	30.37	5.06	0.815	0.565798
Residuals	36	223.75	6.22		

$SS_{\text{total}} = 421$

Analysis of Covariance (ANCOVA)

- blocking allows variation in dependent variable (Y) that is associated with blocking variable to be removed from residuals
- in hyperactivity example, Y was linearly related to baseline activity
 - but blocking variable was a qualitative factor
 - did not fully take advantage of quantitative relation between Y and baseline locomotor activity
- ANCOVA quantitatively models association between dependent variable and **covariate** (baseline activity) using each subject's activity measure rather than dividing subjects into 4 factor groups

ANCOVA

```
> lm.1<-lm(y~activity+drug,data=theData)
> lm.2<-lm(y~activity,data=theData)
> anova(lm.2,lm.1)
```

Analysis of Variance Table

Model	1: y ~ activity	2: y ~ activity + drug				
	Res.Df	RSS	Df	Sum of Sq	F	Pr(>F)
1	46	278.92				
2	44	239.64	2	39.286	3.6066	0.03544 *

```
> anova(lm.1)
```

Analysis of Variance Table

Response: y	Df	Sum Sq	Mean Sq	F value	Pr(>F)
activity	1	142.116	142.116	26.0939	6.739e-06 ***
drug	2	39.286	19.643	3.6066	0.03544 *
Residuals	44	239.638	5.446		

Difference between models shows SS_{drug} after controlling for the linear association between Y and baseline activity

Same result obtained with sequential sums-of-squares ANOVA table for full model

Order of terms does matter

but in this case we probably should put covariate first (why?)

```
> lm.1<-lm(y~activity+drug,data=theData)
> anova(lm.1)
```

Analysis of Variance Table

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
activity	1	142.116	142.116	26.0939	6.739e-06 ***
drug	2	39.286	19.643	3.6066	0.03544 *
Residuals	44	239.638	5.446		

```
> lm.1b<-lm(y~drug+activity,data=theData)
> anova(lm.1b)
```

Analysis of Variance Table

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
drug	2	37.078	18.539	3.4039	0.04222 *
activity	1	144.324	144.324	26.4993	5.91e-06 ***
Residuals	44	239.638	5.446		

```
> library(car)
> Anova(lm.1,type=2)
```

Anova Table (Type II tests)

	Sum Sq	Df	F value	Pr(>F)
activity	144.324	1	26.4993	5.91e-06 ***
drug	39.286	2	3.6066	0.03544 *
Residuals	239.638	44		

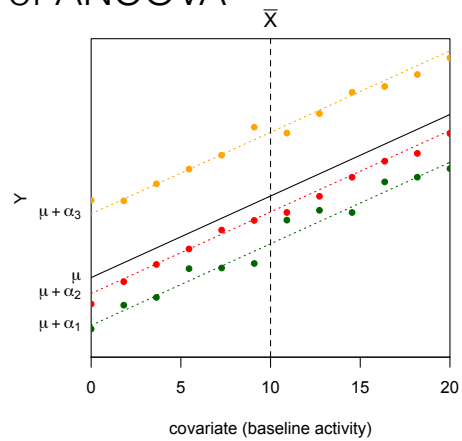
```
> Anova(lm.1b,type=2)
```

Anova Table (Type II tests)

	Sum Sq	Df	F value	Pr(>F)
drug	39.286	2	3.6066	0.03544 *
activity	144.324	1	26.4993	5.91e-06 ***
Residuals	239.638	44		

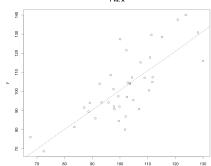
Graphical illustration of ANCOVA

- ANCOVA computes regression lines for each group
 - equal slopes
 - variable intercepts
- group effect (α_j) corresponds to vertical shift of regression intercept

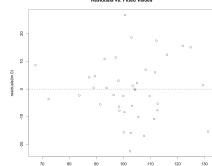


checking linear regression

Y vs X

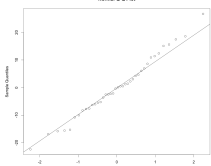


Residuals vs Fitted Values

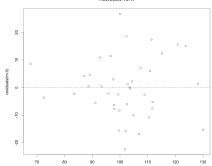


```
par(cex=1.5)
plot(residuals(lm.0)~fitted(lm.0),
     main="Residuals vs. Fitted Values")
abline(h=0, lty=2)
```

qqnorm(residuals)

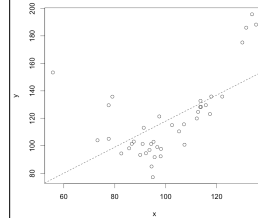


Residuals vs X

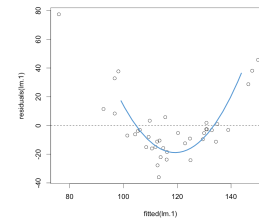


```
plot(residuals(lm.0)~df0$x,
     main="Residuals vs. X")
abline(h=0, lty=2)
```

Y vs. X

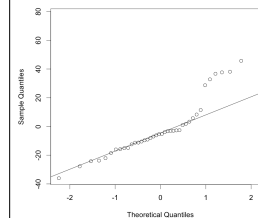


Residuals vs. Fitted Values

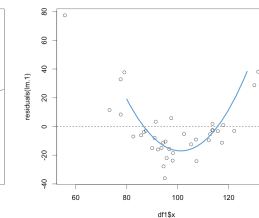


```
par(cex=1.5)
plot(residuals(lm.0)~fitted(lm.0),
     + main="Residuals vs. Fitted Values")
abline(h=0, lty=2)
```

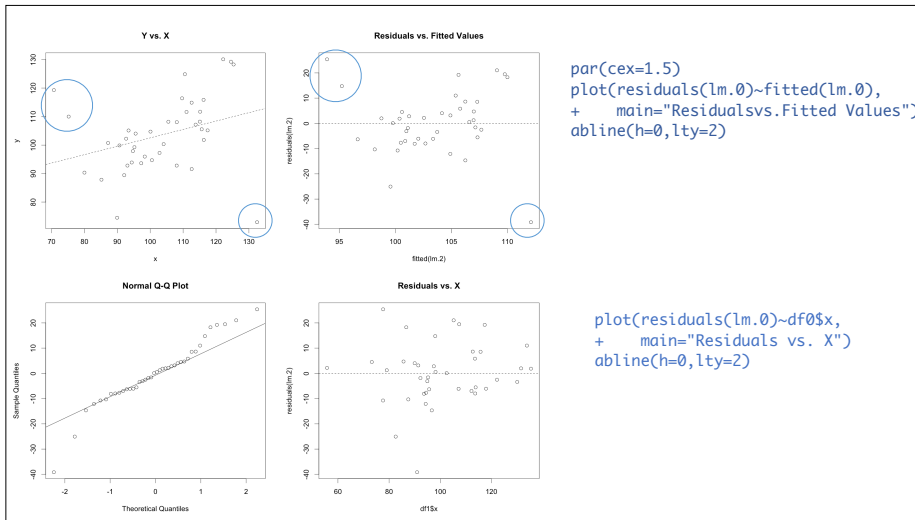
Normal Q-Q Plot



Residuals vs. X

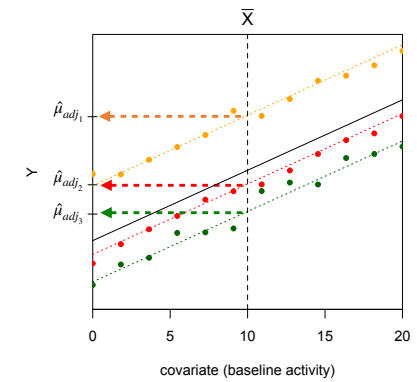


```
plot(residuals(lm.0)~df0$x,
     + main="Residuals vs. X")
abline(h=0, lty=2)
```



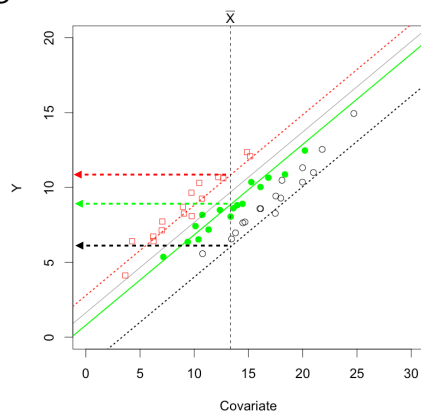
Adjusted Group Means

- point where regression line crosses grand mean of covariate
- provides estimates of group differences after removing effects of covariate



Adjusted Group Means

- point where regression line crosses grand mean of covariate
- provides estimates of group differences after removing effects of covariate
- most useful when groups differ considerably in terms of mean covariate



Computing Adjusted Means

```
> lm.1 <- lm(y~activity+drug, data=theData)
> dummy.coef(lm.1);
```

Full coefficients are

```
(Intercept):    1.28519
activity:        0.5124572
drug:            zero      low      high
                -0.8216185 -0.4386966  1.2603150
```

dummy coefficients list the parameters for the lines fit to each group

```
> library(effects)
> effect(term="drug", lm.1)
```

```
drug effect
drug      zero      low      high
5.238072  5.620994  7.320006
```

$$\bar{Y}'_j = \mu + \beta \bar{X} + \alpha_j$$

covariate mean = 9.3168, so adjusted group means are:

```
> 1.28519 + 0.5124*9.3168-0.8216
```

```
[1] 5.237518 zero
```

```
> 1.28519 + 0.5124*9.3168-0.4386
```

```
[1] 5.620518 low
```

```
> 1.28519 + 0.5124*9.3168+1.2603
```

```
[1] 7.319418 high
```

```
> emmeans(lm.1, specs=~drug)
```

drug	emmean	SE	df	lower.CL	upper.CL
zero	5.24	0.583	44	4.06	6.41
low	5.62	0.583	44	4.45	6.80
high	7.32	0.583	44	6.14	8.50

linear contrasts on adjusted means

$$F = \frac{\hat{\psi}^2}{s_{\hat{\psi}}^2}$$

$$\hat{\psi} = \sum_{j=1}^a c_j \bar{Y}_j'$$

$$s_{\hat{\psi}}^2 = MS_{residuals} \left[\sum_{j=1}^a \frac{c_j^2}{n_j} + \frac{\left(\sum_{j=1}^a c_j \bar{X}_j \right)^2}{\sum_{j=1}^a \sum_{i=1}^n (X_{ij} - \bar{X}_j)^2} \right]$$

```
> lm.1<-lm(y~activity+drug,data=theData)
> levels(theData$drug)
[1] "zero" "low" "high"
> w1 <- c(-1,1,0)
> w2 <- c(-1,0,1)
> w3 <- c(0,-1,1)
> lm1.emm <- emmeans(lm.1,specs=~drug)
> contrast(lm1.emm,method=list(ZvsL=w1,ZvsH=w2,LvsH=w3))
contrast estimate SE df t.ratio p.value
ZvsL 0.383 0.825 44 0.464 0.6449
ZvsH 2.082 0.825 44 2.523 0.0153
LvsH 1.699 0.825 44 2.059 0.0454
```

Association strength

(ignores covariate)

$$\omega^2 = \frac{df_{effect}(MS_{effect} - MS_{residuals})}{SS_{total} + MS_{residuals}}$$

(removes variation due to covariate)

$$\omega_{partial(group)}^2 = \frac{df_{group}(F_{group} - 1)}{df_{group}(F_{group} - 1) + N}$$

```
> require(effectsize)
> mw97.aov.01 <- aov(post~pre+treatment,data=mw97)
> omega_squared(mw97.aov.01,partial=F)
Parameter | Omega2 | 90% CI
-----|-----|-----
pre | 0.23 | [0.04, 0.44]
treatment | 0.12 | [0.00, 0.30]

> omega_squared(mw97.aov.01,partial=T)
Parameter | Omega2 (partial) | 90% CI
-----|-----|-----
pre | 0.26 | [0.05, 0.47]
treatment | 0.15 | [0.00, 0.34]
```

Homogeneity of slopes assumption

- ANCOVA assumes that slope of regression line is the same in each group
 - implies that there is no covariate x group interaction
- if valid, then group differences are independent of covariate
- if not valid, then group differences vary with covariate

```
> anova(lm.4)

Analysis of Variance Table

Response: y
          Df Sum Sq Mean Sq F value    Pr(>F)
activity.c  1 142.116 142.116 26.4881 6.604e-06
drug        2  39.286  19.643  3.6611  0.03422
activity.c:drug  2  14.298  7.149  1.3324  0.27476
Residuals   42 225.340  5.365
```

```
> lm.4 <- lm(y~activity.c + drug + activity.c:drug,data=theData)
```

Why not difference scores?

```
> theData$diff <- theData$y - theData$activity;
> diff.lm.1 <- lm(diff~drug,data=theData)
> anova(diff.lm.1);
```

Analysis of Variance Table

```
Response: diff
          Df Sum Sq Mean Sq F value    Pr(>F)
drug        2  41.46  20.7318  2.5196 0.09179 .
Residuals  45 370.27  8.2282
```

ANOVA on difference scores (activity - baseline.activity) finds drug is not significant. Why?

Why not difference scores?

$$Y_{ij} - X_{ij} = \mu + \alpha_j + \epsilon_{ij}$$

- ANOVA on difference scores is equivalent to ANCOVA with slope fixed at 1.0

$$Y_{ij} = \mu + \alpha_j + X_{ij} + \epsilon_{ij}$$

- When slope $\neq 1$, then ANCOVA provides better fit (lower MS-residuals)