

Repeated measures analysis

Repeated measures by profile analysis

- More than one response *measurement* for each subject. Might be
 - measurements of the same thing at different times
 - measurements of different but related things
- Generalization of matched pairs (“matched triples”, etc.).
- Variation: each subject does several different treatments at different times (called *crossover design*).
- Expect measurements on same subject to be correlated, so assumptions of independence will fail.
- Called *repeated measures*. Different approaches, but *profile analysis* uses **Manova** (set up right way).
- Another approach uses *mixed models* (random effects).

Packages

```
library(car)
library(tidyverse)
library(lme4) # for mixed models later
```

Example: histamine in dogs

- 8 dogs take part in experiment.
- Dogs randomized to one of 2 different drugs.
- Response: log of blood concentration of histamine 0, 1, 3 and 5 minutes after taking drug. (Repeated measures.)
- Data in `dogs.txt`, column-aligned.

Read in data

```
my_url <- "http://ritsokiguess.site/datafiles/dogs.txt"
dogs <- read_table(my_url)
```

Setting things up

```
dogs
```

```
# A tibble: 8 x 7
  dog  drug      x    lh0    lh1    lh3    lh5
  <chr> <chr>    <chr> <dbl> <dbl> <dbl> <dbl>
1 A    Morphine N    -3.22 -1.61 -2.3  -2.53
2 B    Morphine N    -3.91 -2.81 -3.91 -3.91
3 C    Morphine N    -2.66  0.34 -0.73 -1.43
4 D    Morphine N    -1.77 -0.56 -1.05 -1.43
5 E    Trimethaphan N    -3.51 -0.48 -1.17 -1.51
6 F    Trimethaphan N    -3.51  0.05 -0.31 -0.51
7 G    Trimethaphan N    -2.66 -0.19  0.07 -0.22
8 H    Trimethaphan N    -2.41  1.14  0.72  0.21
```

```
response <- with(dogs, cbind(lh0, lh1, lh3, lh5))
dogs.1 <- lm(response ~ drug, data = dogs)
response
```

```
      lh0    lh1    lh3    lh5
[1,] -3.22 -1.61 -2.30 -2.53
[2,] -3.91 -2.81 -3.91 -3.91
```

```
[3,] -2.66  0.34 -0.73 -1.43
[4,] -1.77 -0.56 -1.05 -1.43
[5,] -3.51 -0.48 -1.17 -1.51
[6,] -3.51  0.05 -0.31 -0.51
[7,] -2.66 -0.19  0.07 -0.22
[8,] -2.41  1.14  0.72  0.21
```

The repeated measures MANOVA

Get list of response variable names; we call them `times`. Save in data frame.

```
times <- colnames(response)
times
```

```
[1] "lh0" "lh1" "lh3" "lh5"
```

```
times.df <- data.frame(times=factor(times))
times.df
```

```
times
1  lh0
2  lh1
3  lh3
4  lh5
```

Fitting the model

```
dogs.2 <- Manova(dogs.1,
  idata = times.df,
  idesign = ~times
)
```

The output (some; there is a lot)

```
summary(dogs.2)
```

Type II Repeated Measures MANOVA Tests:

Term: (Intercept)

Response transformation matrix:
(Intercept)

lh0	1
lh1	1
lh3	1
lh5	1

Sum of squares and products for the hypothesis:

(Intercept) 285.366

Multivariate Tests: (Intercept)

	Df	test	stat	approx	F	num	Df	den	Df	Pr(>F)
Pillai	1	0.763467	19.36642	1	6	0.0045648	**			
Wilks	1	0.236533	19.36642	1	6	0.0045648	**			
Hotelling-Lawley	1	3.227738	19.36642	1	6	0.0045648	**			
Roy	1	3.227738	19.36642	1	6	0.0045648	**			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Term: drug

Response transformation matrix:
(Intercept)

lh0	1
lh1	1
lh3	1
lh5	1

Sum of squares and products for the hypothesis:

(Intercept) 46.08

Multivariate Tests: drug

	Df	test	stat	approx	F	num	Df	den	Df	Pr(>F)
Pillai	1	0.3426263	3.127229	1	6	0.12741				
Wilks	1	0.6573737	3.127229	1	6	0.12741				
Hotelling-Lawley	1	0.5212048	3.127229	1	6	0.12741				
Roy	1	0.5212048	3.127229	1	6	0.12741				

Term: times

Response transformation matrix:
times1 times2 times3

lh0	1	0	0
lh1	0	1	0
lh3	0	0	1
lh5	-1	-1	-1

Sum of squares and products for the hypothesis:

	times1	times2	times3
times1	18.9728	-11.103400	-4.0810000
times2	-11.1034	6.498012	2.3883125
times3	-4.0810	2.388313	0.8778125

Multivariate Tests: times

	Df	test	stat	approx	F	num	Df	den	Df	Pr(>F)
Pillai	1	0.949879	25.26898	3	4	0.0046308	**			
Wilks	1	0.050121	25.26898	3	4	0.0046308	**			
Hotelling-Lawley	1	18.951738	25.26898	3	4	0.0046308	**			
Roy	1	18.951738	25.26898	3	4	0.0046308	**			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Term: drug:times

Response transformation matrix:
times1 times2 times3

lh0	1	0	0
-----	---	---	---

```

lh1      0      1      0
lh3      0      0      1
lh5     -1     -1     -1

Sum of squares and products for the hypothesis:
      times1      times2      times3
times1  7.60500  2.0572500 -0.0292500
times2  2.05725  0.5565125 -0.0079125
times3 -0.02925 -0.0079125  0.0001125

Multivariate Tests: drug:times
      Df test stat approx F num Df den Df  Pr(>F)
Pillai      1  0.894761 11.33619      3      4 0.020023 *
Wilks      1  0.105239 11.33619      3      4 0.020023 *
Hotelling-Lawley  1  8.502141 11.33619      3      4 0.020023 *
Roy      1  8.502141 11.33619      3      4 0.020023 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Univariate Type II Repeated-Measures ANOVA Assuming Sphericity

      Sum Sq num Df Error SS den Df F value  Pr(>F)
(Intercept) 71.342      1  22.1026      6 19.3664  0.004565 **
drug      11.520      1  22.1026      6  3.1272  0.127406
times     26.160      3   2.2534     18 69.6546 4.215e-10 ***
drug:times   5.111      3   2.2534     18 13.6095 7.050e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Mauchly Tests for Sphericity

      Test statistic p-value
times      0.12334 0.084567
drug:times  0.12334 0.084567

Greenhouse-Geisser and Huynh-Feldt Corrections
for Departure from Sphericity

      GG eps Pr(>F[GG])
times      0.52618  3.745e-06 ***
drug:times 0.52618  0.002349 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

      HF eps  Pr(>F[HF])
times      0.6822614 1.843418e-07
drug:times 0.6822614 7.307096e-04

```

What there is here

- three sets of tests, for
 - times
 - drug
 - their interaction
- two *types* of test for each of these:
 - multivariate
 - univariate
- multivariate is the same as MANOVA
- univariate is more powerful *if* it applies

Sphericity

- The thing that decides whether the univariate tests apply is called “sphericity”.
- This holds if the outcomes have equal variance (to each other) and have the same (positive) correlation across subjects.
- Tested using Mauchly’s test (part of output)
- If sphericity rejected, there are adjustments to the univariate P-values due to Huynh-Feldt and Greenhouse-Geisser. Huynh-Feldt better if responses not actually normal (safer).

Univariate tests

```
summary(dogs.2)$sphericity.tests
```

	Test statistic	p-value
times	0.12334	0.084567
drug:times	0.12334	0.084567

```
summary(dogs.2)$pval.adjustments
```

	GG eps	Pr(>F[GG])	HF eps	Pr(>F[HF])
times	0.5261798	3.744618e-06	0.6822614	1.843418e-07
drug:times	0.5261798	2.348896e-03	0.6822614	7.307096e-04

```
attr("na.action")  
(Intercept)      drug  
      1          2  
attr("class")  
[1] "omit"
```

```
summary(dogs.2)$univariate.tests
```

	Sum Sq	num Df	Error SS	den Df	F value	Pr(>F)
(Intercept)	71.342	1	22.1026	6	19.3664	0.004565 **
drug	11.520	1	22.1026	6	3.1272	0.127406
times	26.160	3	2.2534	18	69.6546	4.215e-10 ***
drug:times	5.111	3	2.2534	18	13.6095	7.050e-05 ***

```
---  
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Comments

- The sphericity test for the interaction is almost significant
- The H-F adjusted P-value for the interaction is a bit bigger than the univariate one, but still strongly significant.
- Therefore any lack of sphericity does not affect our conclusion: there is an interaction between drug and time
- ie that the effect of time on log-histamine is different for the two drugs.

Comments

- Here, univariate test with Huynh-Feldt correction to P-value for interaction was 0.00073.
- Significant interaction *is* the conclusion here.
- If the interaction had not been significant:
 - cannot remove interaction with time
 - so look at univariate (better, especially if adjusted for sphericity) tests of main effects in *this* model

Next

- Interaction significant. Pattern of response over time different for the two drugs.
- Want to investigate interaction.

The wrong shape

- But data frame has several observations per line (“wide format”):

```
dogs %>% slice(1:6)
```

```
# A tibble: 6 x 7
  dog  drug      x    lh0    lh1    lh3    lh5
<chr> <chr>    <chr> <dbl> <dbl> <dbl> <dbl>
1 A   Morphine N   -3.22 -1.61 -2.3  -2.53
2 B   Morphine N   -3.91 -2.81 -3.91 -3.91
3 C   Morphine N   -2.66  0.34 -0.73 -1.43
4 D   Morphine N   -1.77 -0.56 -1.05 -1.43
5 E   Trimethaphan N -3.51 -0.48 -1.17 -1.51
6 F   Trimethaphan N -3.51  0.05 -0.31 -0.51
```

- Plotting works with data in “long format”: one response per line.
- The responses are log-histamine at different times, labelled **lh**-something. Call them all **lh** and put them in one column, with the time they belong to labelled.

Running pivot_longer, try 1

```
dogs %>% pivot_longer(starts_with("lh"),
                      names_to = "time", values_to = "lh")
```

```
# A tibble: 32 x 5
  dog  drug      x    time    lh
  <chr> <chr>   <chr> <chr> <dbl>
1 A    Morphine N    lh0  -3.22
2 A    Morphine N    lh1  -1.61
3 A    Morphine N    lh3   -2.3
4 A    Morphine N    lh5  -2.53
5 B    Morphine N    lh0  -3.91
6 B    Morphine N    lh1  -2.81
7 B    Morphine N    lh3  -3.91
8 B    Morphine N    lh5  -3.91
9 C    Morphine N    lh0  -2.66
10 C   Morphine N    lh1   0.34
# i 22 more rows
```

Getting the times

Not quite right: for the times, we want just the numbers, not the letters lh every time. Want new variable containing just number in time: `parse_number`.

```
dogs %>%
  pivot_longer(starts_with("lh"),
              names_to = "timex", values_to = "lh") %>%
  mutate(time = parse_number(timex))
```

```
# A tibble: 32 x 6
  dog  drug      x    timex    lh    time
  <chr> <chr>   <chr> <chr> <dbl> <dbl>
1 A    Morphine N    lh0  -3.22     0
2 A    Morphine N    lh1  -1.61     1
3 A    Morphine N    lh3   -2.3     3
4 A    Morphine N    lh5  -2.53     5
5 B    Morphine N    lh0  -3.91     0
6 B    Morphine N    lh1  -2.81     1
7 B    Morphine N    lh3  -3.91     3
8 B    Morphine N    lh5  -3.91     5
9 C    Morphine N    lh0  -2.66     0
10 C   Morphine N    lh1   0.34     1
# i 22 more rows
```


What I did differently

- I realized that `pivot_longer` was going to produce something like `lh1`, which I needed to do something further with, so this time I gave it a temporary name `timex`.
- This enabled me to use the name `time` for the actual numeric time.
- This works now, so next save into a new data frame `dogs.long`.

Saving the pipelined results

```
dogs %>%
  pivot_longer(starts_with("lh"),
               names_to = "timex", values_to = "lh") %>%
  mutate(time = parse_number(timex)) -> dogs.long
dogs.long
```

```
# A tibble: 32 x 6
   dog  drug    x    timex    lh  time
  <chr> <chr>  <chr> <chr> <dbl> <dbl>
1 A     Morphine N    lh0   -3.22    0
2 A     Morphine N    lh1   -1.61    1
3 A     Morphine N    lh3    -2.3    3
4 A     Morphine N    lh5   -2.53    5
5 B     Morphine N    lh0   -3.91    0
6 B     Morphine N    lh1   -2.81    1
7 B     Morphine N    lh3   -3.91    3
8 B     Morphine N    lh5   -3.91    5
9 C     Morphine N    lh0   -2.66    0
10 C    Morphine N    lh1    0.34    1
# i 22 more rows
```

Comments

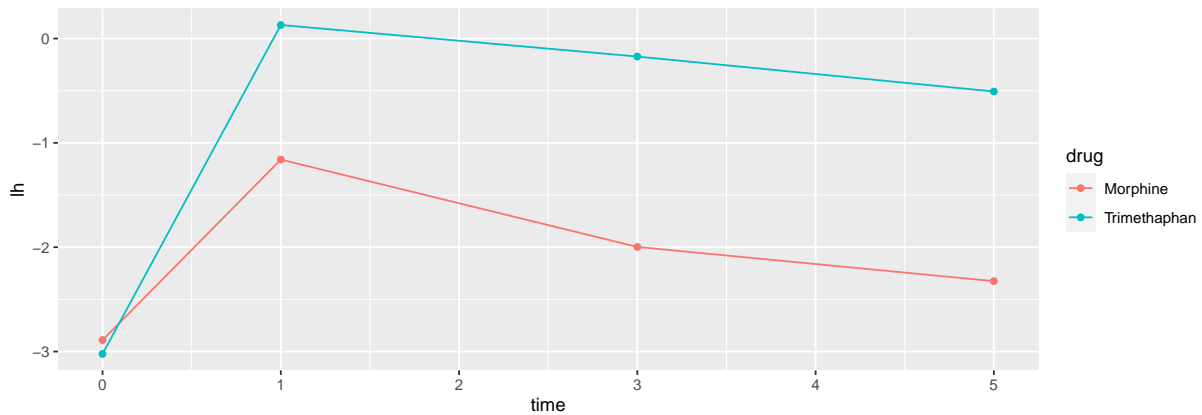
This says:

- Take data frame `dogs`, and then:
- Combine the columns `lh0` through `lh5` into one column called `lh`, with the column that each `lh` value originally came from labelled by `timex`, and then:
- Pull out numeric values in `timex`, saving in `time` and then:

- save the result in a data frame `dogs.long`.

Interaction plot

```
ggplot(dogs.long, aes(x = time, y = lh,
                      colour = drug, group = drug)) +
  stat_summary(fun = mean, geom = "point") +
  stat_summary(fun = mean, geom = "line")
```



Comments

- Plot mean `lh` value at each time, joining points on same drug by lines.
- drugs same at time 0
- after that, Trimethaphan higher than Morphine.
- Effect of drug not consistent over time: significant interaction.

Take out time zero

- Lines on interaction plot would then be parallel, and so interaction should no longer be significant.
- Go back to original “wide” `dogs` data frame.

```
response <- with(dogs, cbind(lh1, lh3, lh5)) # excl time 0
dogs.1 <- lm(response ~ drug, data = dogs)
times <- colnames(response)
```

```
times.df <- data.frame(times=factor(times))
dogs.2 <- Manova(dogs.1,
  idata = times.df,
  idesign = ~times
)
```

Results (univariate)

```
summary(dogs.2)$sphericity.tests
```

```

      Test statistic p-value
times           0.57597 0.25176
drug:times      0.57597 0.25176

```

```
summary(dogs.2)$pval.adjustments
```

```

      GG eps  Pr(>F[GG])  HF eps  Pr(>F[HF])
times      0.7022305 0.0003752847 0.8520467 0.0001117394
drug:times 0.7022305 0.1078608639 0.8520467 0.0942573437
attr(,"na.action")
(Intercept)      drug
              1      2
attr(,"class")
[1] "omit"

```

```
summary(dogs.2)$univariate.tests
```

```

      Sum Sq num Df Error SS den Df F value  Pr(>F)
(Intercept) 24.2607      1 20.1874      6 7.2106 0.03628 *
drug         16.2197      1 20.1874      6 4.8207 0.07053 .
times        3.3250      2  0.7301     12 27.3251 3.406e-05 ***
drug:times   0.3764      2  0.7301     12 3.0929 0.08254 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Comments

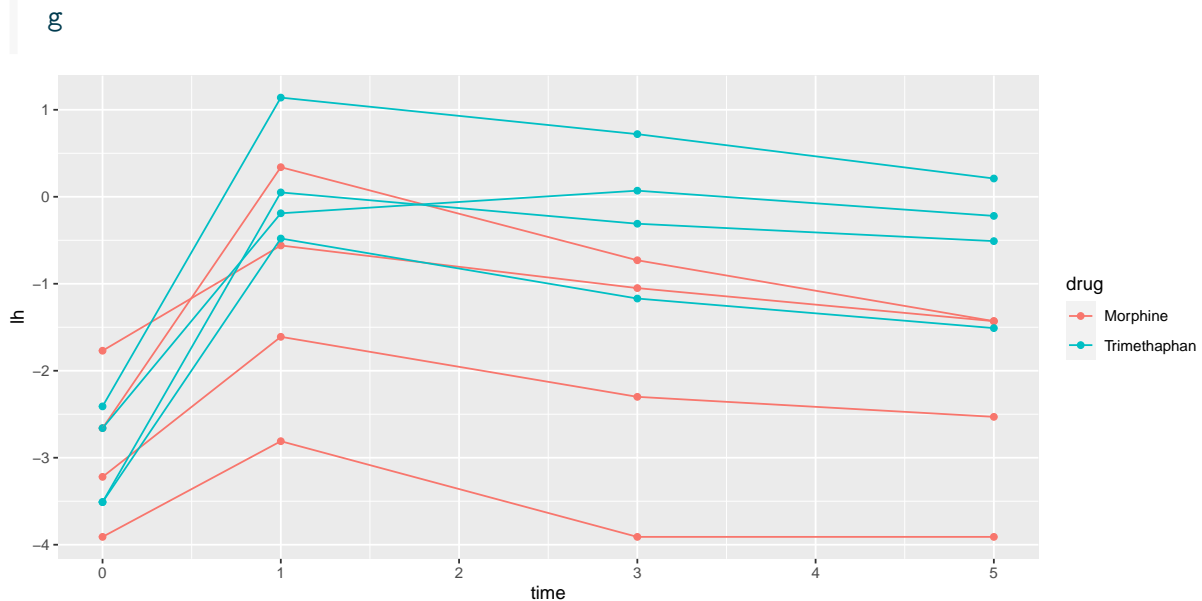
- sphericity: no problem (P-value 0.25)
- univariate test for interaction no longer significant (P-value 0.082)
- look at main effects:
 - strong significance of time, even after taking out time 0
 - actually *not* significant drug effect, despite interaction plot

Is the non-significant drug effect reasonable?

- Plot *actual data*: lh against days, labelling observations by drug: “spaghetti plot”.
- Uses long data frame (confusing, yes I know):
- Plot (time, lh) points coloured by drug
- and connecting measurements for each *dog* by lines.
- This time, we want `group = dog` (want the measurements for each *dog* joined by lines), but `colour = drug`:

```
ggplot(dogs.long, aes(x = time, y = lh,  
  colour = drug, group = dog)) +  
  geom_point() + geom_line() -> g
```

The spaghetti plot



Comments

- For each dog over time, there is a strong increase and gradual decrease in log-histamine. The gradual decrease explains the significant time effect after we took out time 0.

- The pattern is more or less the same for each dog, regardless of drug. This explains the non-significant interaction.
- Most of the trimethaphan dogs (blue) have higher log-histamine throughout (time 1 and after), and some of the morphine dogs have lower.
- *But* two of the morphine dogs have log-histamine profiles like the trimethaphan dogs. This ambiguity is probably why the **drug** effect is not quite significant.

Mixed models

- Another way to fit repeated measures
- Subjects (on whom repeated measures taken) are *random sample of all possible subjects* (random effects)
- Times and treatments are *the only ones we care about* (fixed effects)
- Use package **lme4** function **lmer** (like **lm** in some ways)
- Uses long-format “tidy” data

Fitting the model (uses lme4)

```
# dogs.long including time zero
dogs.3 <- lmer(lh~drug*time+(1|dog), data=dogs.long)
```

- note specification of random effect: each dog has “random intercept” that moves log-histamine up or down for that dog over all times

What can we drop?

- using **drop1**:

```
drop1(dogs.3, test="Chisq")
```

Single term deletions

Model:

```
lh ~ drug * time + (1 | dog)
      npar    AIC    LRT Pr(Chi)
<none>      113.26
drug:time    1 114.21 2.9534 0.0857 .
---
```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

- Interaction again not significant, but P-value smaller than before

Re-fit without interaction

```
dogs.4 <- update(dogs.3, ~.-drug:time)
drop1(dogs.4, test="Chisq")
```

Single term deletions

Model:

```
lh ~ drug + time + (1 | dog)
      npar    AIC    LRT Pr(Chi)
<none>      114.21
drug       1 115.57 3.3560 0.06696 .
time       1 114.96 2.7501 0.09725 .
---
```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

- This time neither drug nor (surprisingly) time is significant.
- MANOVA and `lmer` methods won't agree, but both valid ways to approach problem.

The exercise data

- 30 people took part in an exercise study.
- Each subject was randomly assigned to one of two diets ("low fat" or "non-low fat") and to one of three exercise programs ("at rest", "walking", "running").
- There are $2 \times 3 = 6$ experimental treatments, and thus each one is replicated $30/6 = 5$ times.
- Nothing unusual so far.
- However, each subject had their pulse rate measured at three different times (1, 15 and 30 minutes after starting their exercise), so have repeated measures.

Reading the data

Separated by *tabs*:

```
url <- "http://ritsokiguess.site/datafiles/exercise2.txt"
exercise.long <- read_tsv(url)
exercise.long
```

```
# A tibble: 90 x 5
```

	id	diet	exertype	pulse	time
	<dbl>	<chr>	<chr>	<dbl>	<chr>
1	1	nonlowfat	atrest	85	min01
2	1	nonlowfat	atrest	85	min15
3	1	nonlowfat	atrest	88	min30
4	2	nonlowfat	atrest	90	min01
5	2	nonlowfat	atrest	92	min15
6	2	nonlowfat	atrest	93	min30
7	3	nonlowfat	atrest	97	min01
8	3	nonlowfat	atrest	97	min15
9	3	nonlowfat	atrest	94	min30
10	4	nonlowfat	atrest	80	min01

```
# i 80 more rows
```

- This is “long format”, which is usually what we want.
- But for repeated measures analysis, we want *wide* format!
- `pivot_wider`.

Making wide format

- `pivot_wider` needs: a column that is going to be split, and the column to make the values out of:

```
exercise.long %>% pivot_wider(names_from=time,
                             values_from=pulse) -> exercise.wide
exercise.wide %>% sample_n(5)
```

```
# A tibble: 5 x 6
```

	id	diet	exertype	min01	min15	min30
	<dbl>	<chr>	<chr>	<dbl>	<dbl>	<dbl>
1	2	nonlowfat	atrest	90	92	93
2	26	lowfat	running	95	126	143
3	3	nonlowfat	atrest	97	97	94
4	30	lowfat	running	99	111	150
5	10	lowfat	atrest	100	97	100

- Normally `pivot_longer` min01, min15, min30 into one column called `pulse` labelled by the number of minutes. But `Manova` needs it the other way.

Setting up the repeated-measures analysis

- Make a response variable consisting of min01, min15, min30:

```
response <- with(exercise.wide, cbind(min01, min15, min30))
```

- Predict that from diet and exertype and interaction using lm:

```
exercise.1 <- lm(response ~ diet * exertype,  
  data = exercise.wide  
)
```

- Run this through Manova:

```
times <- colnames(response)  
times.df <- data.frame(times=factor(times))  
exercise.2 <- Manova(exercise.1,  
  idata = times.df,  
  idesign = ~times)
```

Sphericity tests

```
summary(exercise.2)$sphericity.tests
```

	Test statistic	p-value
times	0.92416	0.40372
diet:times	0.92416	0.40372
exertype:times	0.92416	0.40372
diet:exertype:times	0.92416	0.40372

No problem with sphericity; go to univariate tests.

Univariate tests

```
summary(exercise.2)$univariate.tests
```


	Sum Sq	num Df	Error SS	den Df	F value	Pr(>F)
(Intercept)	894608	1	2085.2	24	10296.6595	< 2.2e-16 ***
diet	1262	1	2085.2	24	14.5238	0.0008483 ***
exertype	8326	2	2085.2	24	47.9152	4.166e-09 ***
diet:exertype	816	2	2085.2	24	4.6945	0.0190230 *
times	2067	2	1563.6	48	31.7206	1.662e-09 ***
diet:times	193	2	1563.6	48	2.9597	0.0613651 .
exertype:times	2723	4	1563.6	48	20.9005	4.992e-10 ***
diet:exertype:times	614	4	1563.6	48	4.7095	0.0027501 **

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

- The three-way interaction is significant
 - the effect of diet on pulse rate over time is different for the different exercise types

Making some graphs

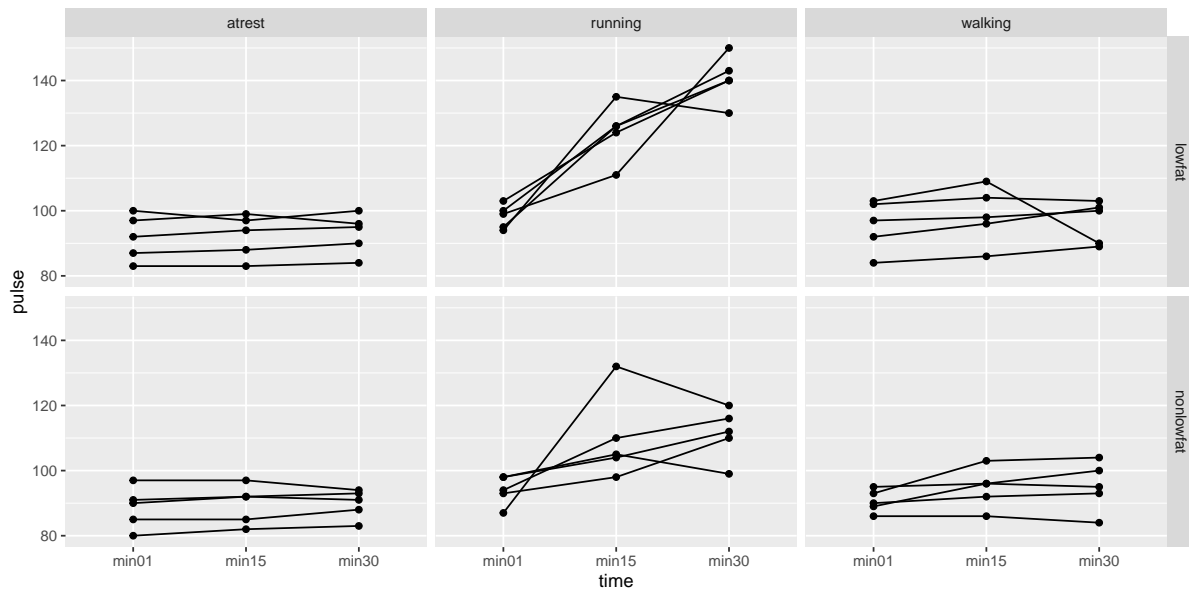
- Three-way interactions are difficult to understand. To make an attempt, look at some graphs.
- Plot time trace of pulse rates for each individual, joined by lines, and make *separate* plots for each diet-exertype combo.
- ggplot again. Using *long* data frame:

```
g <- ggplot(exercise.long, aes(
  x = time, y = pulse,
  group = id
)) + geom_point() + geom_line() +
  facet_grid(diet ~ exertype)
```

- `facet_grid(diet~exertype)`: do a separate plot for each combination of diet and exercise type, with diets going down the page and exercise types going across. (Graphs are usually landscape, so have the factor `exertype` with more levels going across.)

The graph(s)

```
g
```



Comments on graphs

- For subjects who were at rest, no change in pulse rate over time, for both diet groups.
- For walking subjects, not much change in pulse rates over time. Maybe a small increase on average between 1 and 15 minutes.
- For both running groups, an overall increase in pulse rate over time, but the increase is stronger for the **lowfat** group.
- No consistent effect of diet over all exercise groups.
- No consistent effect of exercise type over both diet groups.
- No consistent effect of time over all diet-exercise type combos.

“Simple effects” of diet for the subjects who ran

- Looks as if there is only any substantial time effect for the runners. For them, does diet have an effect?
- Pull out only the runners from the wide data:

```
exercise.wide %>%
  filter(exertype == "running") -> runners.wide
```

- Create response variable and do MANOVA. Some of this looks like before, but I have different data now:

```
response <- with(runners.wide, cbind(min01, min15, min30))
runners.1 <- lm(response ~ diet, data = runners.wide)
times <- colnames(response)
times.df <- data.frame(times=factor(times))
runners.2 <- Manova(runners.1,
  idata = times.df,
  idesign = ~times
)
```

Sphericity tests

```
summary(runners.2)$sphericity.tests
```

	Test statistic	p-value
times	0.81647	0.4918
diet:times	0.81647	0.4918

- No problem, look at univariate tests.

Univariate tests

```
summary(runners.2)$univariate.tests
```

	Sum Sq	num Df	Error SS	den Df	F value	Pr(>F)
(Intercept)	383522	1	339.2	8	9045.3333	1.668e-13 ***
diet	1920	1	339.2	8	45.2830	0.0001482 ***
times	4714	2	1242.0	16	30.3644	3.575e-06 ***
diet:times	789	2	1242.0	16	5.0795	0.0195874 *

 Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

- Interaction still significant
 - dependence of pulse rate on time still different for the two diets

How is the effect of diet different over time?

- Table of means. Only I need long data for this:

```
runners.wide %>%
  pivot_longer(starts_with("min"),
               names_to = "time", values_to = "pulse") %>%
  group_by(time, diet) %>%
  summarize(
    mean = mean(pulse),
    sd = sd(pulse)
  ) -> summ
```

- Result of `summarize` is data frame, so can save it (and do more with it if needed).

Understanding diet-time interaction

- The summary:

```
summ
```

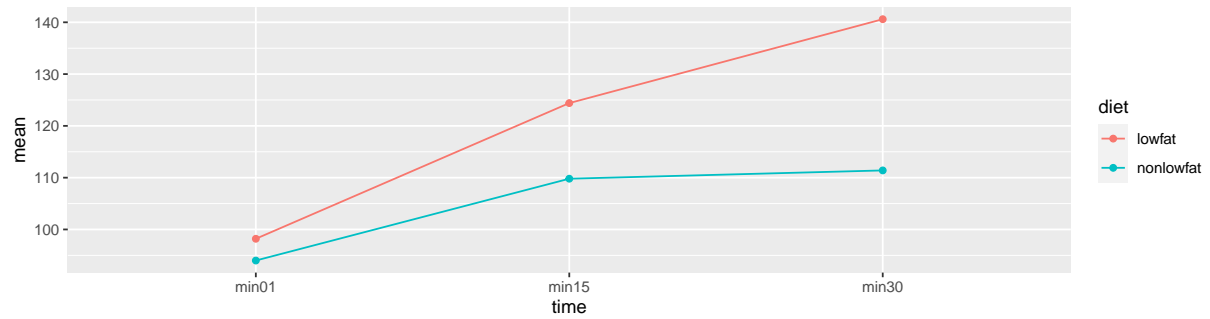
```
# A tibble: 6 x 4
# Groups:   time [3]
  time  diet    mean    sd
  <chr> <chr>   <dbl> <dbl>
1 min01 lowfat    98.2  3.70
2 min01 nonlowfat  94    4.53
3 min15 lowfat   124.   8.62
4 min15 nonlowfat 110.  13.1
5 min30 lowfat   141.   7.20
6 min30 nonlowfat 111.   7.92
```

- Pulse rates at any given time higher for `lowfat` (diet effect),
- Pulse rates increase over time of exercise (time effect),
- but the *amount by which pulse rate higher* for a diet depends on time: diet by time interaction.

Interaction plot

- We went to trouble of finding means by group, so making interaction plot is now mainly easy:

```
ggplot(summ, aes(x = time, y = mean, colour = diet,
                 group = diet)) + geom_point() + geom_line()
```



Comment on interaction plot

- The lines are not parallel, so there is interaction between diet and time for the runners.
- The effect of time on pulse rate is different for the two diets, even though all the subjects here were running.