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)</pre>
```

Setting things up

dogs

```
# A tibble: 8 x 7
 dog
       drug
                    X
                           1h0
                                 lh1
                                       1h3
                                            1h5
  <chr> <chr>
                    <chr> <dbl> <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
                         -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
                         -2.41 1.14 0.72 0.21
8 H
       Trimethaphan N
response <- with(dogs, cbind(lh0, lh1, lh3, lh5))
response
```

1h0 1h1 1h3 1h5 [1,] -3.22 -1.61 -2.30 -2.53

The repeated measures MANOVA

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

```
times <- colnames(response)</pre>
times
[1] "lh0" "lh1" "lh3" "lh5"
times.df <- data.frame(times=factor(times))</pre>
times.df
  times
    1h0
  lh1
3
  1h3
4
    1h5
```

Fitting the model

```
dogs.1 <- lm(response ~ drug, data = dogs)
dogs.2 <- Manova(dogs.1,
  idata = times.df,
  idesign = ~times
)</pre>
```

The output (come there is a lot)

```
Type II Repeated Measures MANOVA Tests:
Term: (Intercept)
 Response transformation matrix:
   (Intercept)
1h0
1h1
1h3
1h5
Sum of squares and products for the hypothesis:
          (Intercept)
(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 **
              1 3.227738 19.36642 1 6 0.0045648 **
Roy
---
Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
------
Term: drug
Response transformation matrix:
   (Intercept)
```

1h0

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

times

summary(dogs.2)\$sphericity.tests

drug:times 0.12334 0.084567

Test statistic p-value 0.12334 0.084567

```
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
                               1h1 1h3
 dog
       drug
                          1h0
                                          1h5
                 X
 <chr> <chr> <chr> <chr> <dbl> <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
                  N -2.66 0.34 -0.73 -1.43
3 C
    Morphine
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 1h-something. Call them all 1h 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
                               1h
  dog
        drug x
                      time
  <chr> <chr> <chr> <chr> <chr> <chr> <chr> <dbl>
 1 A
        Morphine N
                      1h0 -3.22
        Morphine N
2 A
                      lh1 -1.61
3 A
        Morphine N
                      1h3 -2.3
4 A Morphine N
                      1h5 -2.53
5 B
        Morphine N
                      lh0 -3.91
6 B
        Morphine N
                      lh1 -2.81
7 B
        Morphine N
                      1h3
                           -3.91
8 B
        Morphine N
                      lh5
                           -3.91
9 C
        Morphine N
                      lh0
                           -2.66
10 C
        Morphine N
                             0.34
                      lh1
# i 22 more rows
```

Getting the times

Not quite right: for the times, we want just the numbers, not the letters 1h every time. Want new variable containing just number in time: parse_number.

```
# A tibble: 32 x 6
  dog
                            1h
       drug x
                    timex
                               time
  <chr> <chr> <chr> <chr> <chr> <dbl> <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
                                  3
8 B
       Morphine N
                    lh5 -3.91
                                  5
9 C
       Morphine N
                    lh0 -2.66
10 C
       Morphine N
                    lh1
                          0.34
```

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

Morphine N

Morphine N

Morphine N

7 B

8 B

9 C

```
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
                     timex lh
  dog
       drug x
                                time
  <chr> <chr> <chr> <chr> <chr> <dbl> <dbl>
 1 A
       Morphine N
                     1h0 -3.22
                     lh1 -1.61
2 A
       Morphine N
                     lh3 -2.3
                                   3
3 A
       Morphine N
                     lh5 -2.53
4 A
       Morphine N
5 B
       Morphine N
                    lh0 -3.91
       Morphine N
6 B
                    lh1 -2.81
```

lh3 -3.91

lh5 -3.91

-2.66

lh0

3

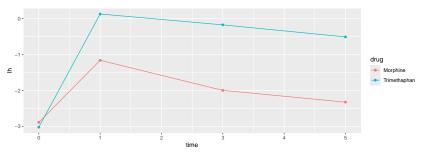
5

Comments

This says:

- Take data frame dogs, and then:
- Combine the columns 1h0 through 1h5 into one column called 1h, with the column that each 1h 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



Comments

- Plot mean 1h 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
)</pre>
```

Results (univariate)

summary(dogs.2)\$sphericity.tests

Test statistic p-value 0.57597 0.25176

(Intercept) drug 1 2 attr(,"class")

[1] "omit"

times

summary(dogs.2)\$univariate.tests

Comments

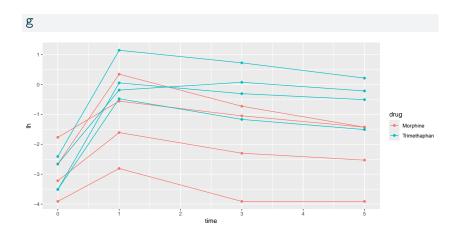
- > sphericity: no problem (P-value 0.25)
- univariate test for interaction no longer significant (P-value 0.082)
- look at main effects:
 - strong significance of time, even after taking out time 0
 - actually not significant drug effect, despite interaction plot

Is the non-significant drug effect reasonable?

- Plot actual data: 1h 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 1me4)

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

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?

Model:

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

```
Single term deletions
```

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

Re-fit without interaction

```
dogs.4 <- update(dogs.3,.~.-drug:time)</pre>
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 '
```

- ▶ This time neither drug nor (surprisingly) time is significant.
- MANOVA and 1mer 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

4

5

6

8

9

10

Separated by tabs:

```
url <- "http://ritsokiguess.site/datafiles/exercise2.txt"</pre>
exercise.long <- read_tsv(url)
exercise.long
# A tibble: 90 x 5
     id diet exertype pulse time
   <dbl> <chr>
                 <chr> <dbl> <chr>
      1 nonlowfat atrest
                             85 min01
      1 nonlowfat atrest
 2
                             85 min15
3
      1 nonlowfat atrest
                             88 min30
```

92 min15

93 min30

97 min01

97 min15

94 min30

80 min01

2 nonlowfat atrest 90 min01

2 nonlowfat atrest

2 nonlowfat atrest

3 nonlowfat atrest

3 nonlowfat atrest

3 nonlowfat atrest

4 nonlowfat atrest

Making wide format

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

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))</pre>
```

Predict that from diet and exertype and interaction using lm:

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

Run this through Manova:

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

Signif. codes:

summary(exercise.2)\$univariate.tests

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

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

0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Making some graphs

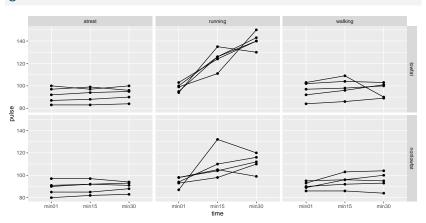
- Three-way interactions are difficult to understand. To make an attempt, look at some graphs.
- ▶ Plot time trace of pulse rates for each individual, joined by lines, and make separate plots for each diet-exertype combo.
- ggplot again. Using *long* data frame:

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

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)





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
)</pre>
```

Sphericity tests

summary(runners.2)\$sphericity.tests

```
Test statistic p-value times 0.81647 \quad 0.4918 diet:times 0.81647 \quad 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
                      339.2
                               8 9045.3333 1.668e-13 ***
diet
         1920 1 339.2 8 45.2830 0.0001482 ***
        4714 2 1242.0 16 30.3644 3.575e-06 ***
times
                  2 1242.0
diet:times
           789
                              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:

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> <chr> <chr> chr> chr> chr> 3.70

2 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:

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.