

Lecture 8

Repeated Measures Analysis

DSA 8070 Multivariate Analysis

October 10 - October 14, 2022

Whitney Huang
Clemson University

A completely randomized block design was carried out to determine the effects of 4 surgical treatments on coronary potassium in a group of 36 dogs. There are 9, 8, 9, and 10 dogs in each treatment group, respectively. Each dog was **measured at four different time points** (1, 5, 9, and 13 minutes) following one of four experimental treatments:

- Control - no surgical treatment is applied
- Extrinsic cardiac denervation immediately prior to treatment
- Bilateral thoracic sympathectomy and stellectomy 3 weeks prior to treatment
- Extrinsic cardiac denervation 3 weeks prior to treatment

We are looking at the treatment effect on the coronary sinus potassium levels

Let Y_{ijk} be the potassium level for treatment i in dog j at time k :

- there are $a = 4$ treatments (i.e., $i = 1, 2, 3, 4$)
- n_i dogs received treatment i (therefore, there are $n_1 + \dots + n_a = 9 + 8 + 9 + 10 = 36$ dogs in total)
- $t = 4$, the number of observations over time (i.e., $k = 1, 2, 3, 4$)

Approaches

- Split-plot ANOVA
- MANOVA
- Mixed Models

Approach 1: Split-plot ANOVA

Model: $Y_{ijk} = \mu + \alpha_i + \delta_{j(i)} + \beta_k + (\alpha\beta)_{ik} + \varepsilon_{ijk}$,
where

- α_i : effect of treatment i
- $\delta_{j(i)}$: random effect of dog j receiving treatment i
- β_k : effect of time k
- $(\alpha\beta)_{ik}$: treatment by time interaction
- ε_{ijk} : random error

Assumptions:

- $\varepsilon_{ijk} \stackrel{i.i.d.}{\sim} N(0, \sigma_\varepsilon^2)$
- $\delta_{j(i)} \stackrel{i.i.d.}{\sim} N(0, \sigma_\delta^2)$
- β_k does not depend on the dog \Rightarrow no time by dog interaction

Split-plot ANOVA Table

Source	df	MS	F
Trt	$a - 1$	$MS_{trt} = \frac{SS_{trt}}{a-1}$	$F = \frac{MS_{trt}}{MS_{error_1}}$
Error 1	$N - a$	$MS_{error_1} = \frac{SS_{error_1}}{N-a}$	
Time	$t - 1$	$MS_{time} = \frac{SS_{time}}{t-1}$	$F = \frac{MS_{time}}{MS_{error_2}}$
Trt \times Time	$(a - 1)(t - 1)$	$MS_{trt \times time} = \frac{SS_{trt \times time}}{(a-1)(t-1)}$	$F = \frac{MS_{trt \times time}}{MS_{error_2}}$
Error 2	$(N - a)(t - 1)$	$MS_{error_2} = \frac{SS_{error_2}}{(N-a)(t-1)}$	
Total	$Nt - 1$		

Dog Experiment Split-plot Analysis

```
> library(lmerTest)
> fit <- lmer(Response ~ Treatment * Time + (1 | Dog_id), data = dat)
> anova(fit)
```

Type III Analysis of Variance Table with Satterthwaite's method

	Sum Sq	Mean Sq	NumDF	DenDF	F value	Pr(>F)
Treatment	3.3396	1.11319	3	32	6.0038	0.002297 **
Time	6.2043	2.06811	3	96	11.1540	2.404e-06 ***
Treatment:Time	3.4397	0.38219	9	96	2.0613	0.040573 *

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

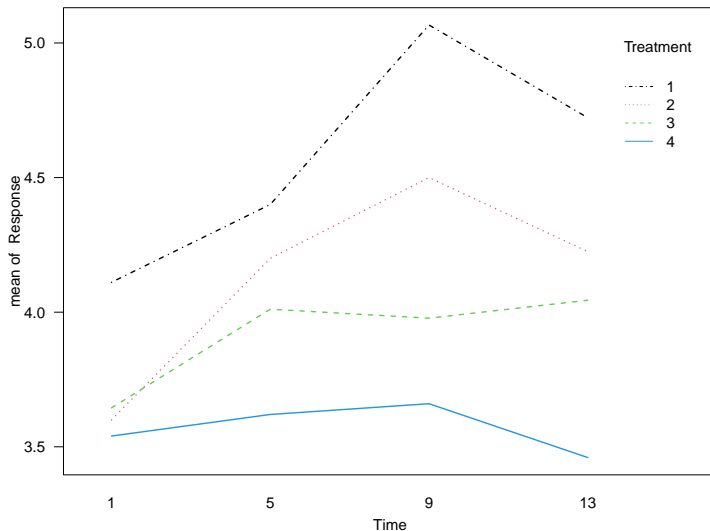
Hypothesis Tests:

We start with the interaction between treatment and time:

$$H_0 : (\alpha\beta)_{ik} = 0 \quad \forall i = 1, \dots, a, k = 1, 2, \dots, t$$

Result: We conclude the effect of treatment depends on time at $\alpha = 0.05$ level

Interaction Plot



Rejecting $H_0 : (\alpha\beta)_{ik} = 0$ means we reject the assumption of “parallelism”

- The Split-plot ANOVA Approach assumes a constant correlation between any two observations from the same dog, that is, $\text{Cor}(Y_{ijk}, Y_{ijk'}) = \frac{\sigma_\delta^2}{\sigma_\delta^2 + \sigma_\varepsilon^2}$, this is the so-called **compound symmetry** correlation structure
- This assumption is unlikely to be valid with repeated measurements over time as the correlation for two nearby time points is likely to be higher than the correlation for two far apart time points
- Next, we are going to take a multivariate approach (MANOVA) as an attempt to address this issue

Here we consider the observations over time from the same dog, dog j receiving treatment i as a single vector of interest

$$\mathbf{Y}_{ij} = (Y_{ij1}, Y_{ij2}, \dots, Y_{ijt})^T,$$

and we will perform a one-way MANOVA

Assumptions:

- Dogs receiving treatment i have common mean vector μ_i
- All dogs have common covariance matrix Σ
- Data from different dogs are independently sampled
- Data are multivariate normally distributed

```
> dat <- read.table("dog1.txt")
> out <- manova(cbind(V3, V4, V5, V6) ~ as.factor(V1), data = dat)
> summary(out, test = "Wilks")
```

	Df	Wilks	approx F	num Df	den Df	Pr(>F)
as.factor(V1)	3	0.48452	2.022	12	77.018	0.03316 *
Residuals	32					

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

Results: There are significant differences between at least one pair of treatments in at least one measurement of time

Criticism: MANOVA makes no assumptions regarding the temporal correlation structure, and hence, may be overparameterized leading to poor parameter estimates

Approach 3: Mixed Model Analysis

Main idea: Split-plot makes a too restrictive assumption while MANOVA makes no assumptions regarding the temporal correlation structure. The mixed model approach allows us to model the temporal correlation involving a limited number of parameters.

Model: $Y_{ijk} = \mu + \alpha_i + \delta_{j(i)} + \beta_k + (\alpha\beta)_{ik} + \varepsilon_{ijk}$.

Assumptions:

- $\varepsilon_{j(ik)} \stackrel{i.i.d.}{\sim} N(0, \sigma_\varepsilon^2)$
- $\delta_{j(i)} \stackrel{i.i.d.}{\sim} N(0, \sigma_\delta^2)$
- The correlation between the errors for the same dog depends only on the difference in observation time points: $|k - k'|$, e.g., $\text{Cor}(Y_{ijk}, Y_{ijk'}) = \rho^{|k-k'|}$ (Autoregressive with order 1)

```
> library(nlme)
> fit1 = gls(Response ~ Treatment * Time,
+           correlation = corCompSymm(form = ~ 1 | Dog_id), data = dat2)
> fit2 = gls(Response ~ Treatment * Time,
+           correlation = corAR1(form = ~ 1 | Dog_id), data = dat2)
> anova(fit1, fit2)
```

	Model	df	AIC	BIC	logLik
fit1	1	18	275.8063	327.1429	-119.9032
fit2	2	18	277.5811	328.9177	-120.7906

Results:

- Based on both AIC/BIC, having an AR(1) does not necessarily improve the model fit (in this data)
- However, having the option of modeling repeated measurement error structure can be useful in general as it provides additional modeling choices