

**A Handbook of Statistical Analyses
Using R — 2nd Edition**

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Analysing Longitudinal Data I: Computerised Delivery of Cognitive Behavioural Therapy – Beat the Blues

12.1 Introduction

12.2 Analysing Longitudinal Data

12.3 Analysis Using R

We shall fit both random intercept and random intercept and slope models to the data including the baseline BDI values (`pre.bdi`), `treatment` group, `drug` and `length` as fixed effect covariates. Linear mixed effects models are fitted in R by using the `lmer` function contained in the `lme4` package (Bates and Sarkar, 2008, Pinheiro and Bates, 2000, Bates, 2005), but an essential first step is to rearrange the data from the ‘wide form’ in which they appear in the `BtheB` data frame into the ‘long form’ in which each separate repeated measurement and associated covariate values appear as a separate row in a *data.frame*. This rearrangement can be made using the following code:

```
R> data("BtheB", package = "HSAUR2")
R> BtheB$subject <- factor(rownames(BtheB))
R> nobs <- nrow(BtheB)
R> BtheB_long <- reshape(BtheB, idvar = "subject",
+   varying = c("bdi.2m", "bdi.3m", "bdi.5m", "bdi.8m"),
+   direction = "long")
R> BtheB_long$time <- rep(c(2, 3, 5, 8), rep(nobs, 4))
```

such that the data are now in the form (here shown for the first three subjects)

```
R> subset(BtheB_long, subject %in% c("1", "2", "3"))
```

	<i>drug</i>	<i>length</i>	<i>treatment</i>	<i>bdi.pre</i>	<i>subject</i>	<i>time</i>	<i>bdi</i>
1.2m	No	>6m	TAU	29	1	2	2
2.2m	Yes	>6m	BtheB	32	2	2	16
3.2m	Yes	<6m	TAU	25	3	2	20
1.3m	No	>6m	TAU	29	1	3	2
2.3m	Yes	>6m	BtheB	32	2	3	24
3.3m	Yes	<6m	TAU	25	3	3	NA
1.5m	No	>6m	TAU	29	1	5	NA
2.5m	Yes	>6m	BtheB	32	2	5	17
3.5m	Yes	<6m	TAU	25	3	5	NA
1.8m	No	>6m	TAU	29	1	8	NA

```

R> data("BtheB", package = "HSAUR2")
R> layout(matrix(1:2, nrow = 1))
R> ylim <- range(BtheB[,grep("bdi", names(BtheB))],
+               na.rm = TRUE)
R> tau <- subset(BtheB, treatment == "TAU")[,
+             grep("bdi", names(BtheB))]
R> boxplot(tau, main = "Treated as Usual", ylab = "BDI",
+          xlab = "Time (in months)", names = c(0, 2, 3, 5, 8),
+          ylim = ylim)
R> btheb <- subset(BtheB, treatment == "BtheB")[,
+             grep("bdi", names(BtheB))]
R> boxplot(btheb, main = "Beat the Blues", ylab = "BDI",
+          xlab = "Time (in months)", names = c(0, 2, 3, 5, 8),
+          ylim = ylim)

```

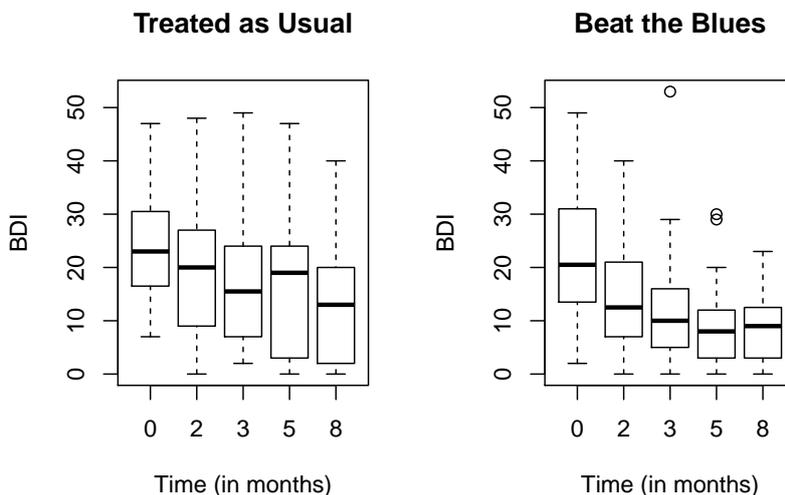


Figure 12.1 Boxplots for the repeated measures by treatment group for the `BtheB` data.

2.8m	Yes	>6m	<i>BtheB</i>	32	2	8	20
3.8m	Yes	<6m	<i>TAU</i>	25	3	8	NA

The resulting `data.frame` `BtheB_long` contains a number of missing values and in applying the `lmer` function these will be dropped. But notice it is only the missing values that are removed, *not* participants that have at least one missing value. All the available data is used in the model fitting process. The `lmer` function is used in a similar way to the `lm` function met in Chapter~6 with the addition of a random term to identify the source of the repeated

measurements, here `subject`. We can fit the two models (??) and (??) and test which is most appropriate using

```
R> library("lme4")
R> BtheB_lmer1 <- lmer(bdi ~ bdi.pre + time + treatment + drug +
+   length + (1 | subject), data = BtheB_long,
+   REML = FALSE, na.action = na.omit)
R> BtheB_lmer2 <- lmer(bdi ~ bdi.pre + time + treatment + drug +
+   length + (time | subject), data = BtheB_long,
+   REML = FALSE, na.action = na.omit)
R> anova(BtheB_lmer1, BtheB_lmer2)

Data: BtheB_long
Models:
BtheB_lmer1: bdi ~ bdi.pre + time + treatment + drug + length + (1 | subject)
BtheB_lmer2: bdi ~ bdi.pre + time + treatment + drug + length + (time | subject)
      Df    AIC    BIC  logLik  Chisq Chi Df Pr(>Chisq)
BtheB_lmer1  8 1887.5 1916.6 -935.75
BtheB_lmer2 10 1891.0 1927.4 -935.52 0.4542      2    0.7969
```

The `summary` method for `lmer` objects doesn't print p -values for Gaussian mixed models because the degrees of freedom of the t reference distribution are not obvious. However, one can rely on the asymptotic normal distribution for computing univariate p -values for the fixed effects using the `cfest` function from package **multcomp**. The asymptotic p -values are given in Figure~12.3.

We can check the assumptions of the final model fitted to the `BtheB` data, i.e., the normality of the random effect terms and the residuals, by first using the `ranef` method to *predict* the former and the `residuals` method to calculate the differences between the observed data values and the fitted values, and then using normal probability plots on each. How the random effects are predicted is explained briefly in Section~??. The necessary R code to obtain the effects, residuals and plots is shown with Figure~12.4. There appear to be no large departures from linearity in either plot.

```
R> summary(BtheB_lmer1)
```

```
Linear mixed model fit by maximum likelihood
Formula: bdi ~ bdi.pre + time + treatment + drug + length + (1 | subject)
Data: BtheB_long
   AIC   BIC logLik deviance REMLdev
1887 1917 -935.7   1871    1867
Random effects:
Groups   Name          Variance Std.Dev.
subject (Intercept) 48.777    6.9841
Residual                25.140    5.0140
Number of obs: 280, groups: subject, 97

Fixed effects:
              Estimate Std. Error t value
(Intercept)    5.59244    2.24232    2.494
bdi.pre         0.63967    0.07789    8.213
time           -0.70477    0.14639   -4.814
treatmentBtheB -2.32912    1.67026   -1.394
drugYes        -2.82497    1.72674   -1.636
length>6m      0.19712    1.63823    0.120

Correlation of Fixed Effects:
              (Intr) bdi.pr time  trtmBB drugYs
bdi.pre       -0.682
time          -0.238  0.020
tretmntBthB  -0.390  0.121  0.018
drugYes       -0.073 -0.237 -0.022 -0.323
length>6m    -0.243 -0.242 -0.036  0.002  0.157
```

Figure 12.2 R output of the linear mixed-effects model fit for the BtheB data.

```
R> cftest(BtheB_lmer1)
```

Simultaneous Tests for General Linear Hypotheses

```
Fit: lmer(formula = bdi ~ bdi.pre + time + treatment + drug + length +
          (1 | subject), data = BtheB_long, REML = FALSE, na.action = na.omit)
```

Linear Hypotheses:

	<i>Estimate</i>	<i>Std. Error</i>	<i>z value</i>	<i>Pr(> z)</i>
<i>(Intercept) == 0</i>	5.59244	2.24232	2.494	0.0126
<i>bdi.pre == 0</i>	0.63967	0.07789	8.213	2.22e-16
<i>time == 0</i>	-0.70477	0.14639	-4.814	1.48e-06
<i>treatmentBtheB == 0</i>	-2.32912	1.67026	-1.394	0.1632
<i>drugYes == 0</i>	-2.82497	1.72674	-1.636	0.1018
<i>length>6m == 0</i>	0.19712	1.63823	0.120	0.9042

(Univariate p values reported)

Figure 12.3 R output of the asymptotic p -values for linear mixed-effects model fit for the BtheB data.

```
R> layout(matrix(1:2, ncol = 2))
R> qint <- ranef(BtheB_lmer1)$subject[["(Intercept)"]]
R> qres <- residuals(BtheB_lmer1)
R> qqnorm(qint, ylab = "Estimated random intercepts",
+        xlim = c(-3, 3), ylim = c(-20, 20),
+        main = "Random intercepts")
R> qqline(qint)
R> qqnorm(qres, xlim = c(-3, 3), ylim = c(-20, 20),
+        ylab = "Estimated residuals",
+        main = "Residuals")
R> qqline(qres)
```

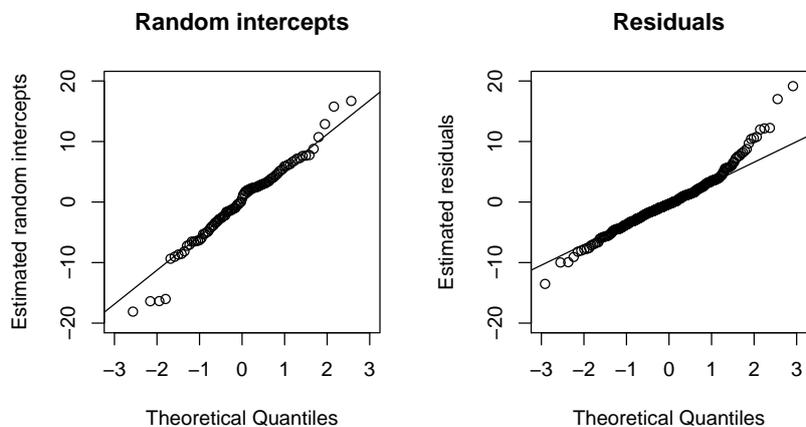


Figure 12.4 Quantile-quantile plots of predicted random intercepts and residuals for the random intercept model `BtheB_lmer1` fitted to the `BtheB` data.

```
R> bdi <- BtheB[, grep("bdi", names(BtheB))]
R> plot(1:4, rep(-0.5, 4), type = "n", axes = FALSE,
+       ylim = c(0, 50), xlab = "Months", ylab = "BDI")
R> axis(1, at = 1:4, labels = c(0, 2, 3, 5))
R> axis(2)
R> for (i in 1:4) {
+   dropout <- is.na(bdi[,i + 1])
+   points(rep(i, nrow(bdi)) + ifelse(dropout, 0.05, -0.05),
+         jitter(bdi[,i]), pch = ifelse(dropout, 20, 1))
+ }
```

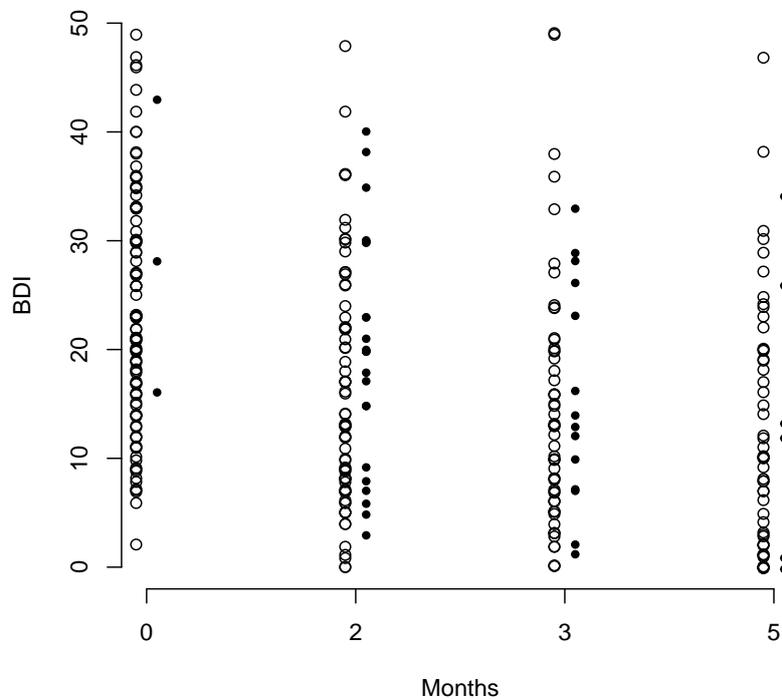


Figure 12.5 Distribution of BDI values for patients that do (circles) and do not (bullets) attend the next scheduled visit.



Bibliography

- Bates, D. (2005), “Fitting linear mixed models in R,” *R News*, 5, 27–30, URL <http://CRAN.R-project.org/doc/Rnews/>.
- Bates, D. and Sarkar, D. (2008), *lme4: Linear Mixed-Effects Models Using Eigen and S4 Classes*, URL <http://CRAN.R-project.org/package=lme4>, R package version 0.999375-28.
- Pinheiro, J. C. and Bates, D. M. (2000), *Mixed-Effects Models in S and S-PLUS*, New York, USA: Springer-Verlag.