### Contents

1	Introduction	1
	1.1 nlmeU R package	
	1.2 ARMD	1
2	Back Ground Materials	2
	2.1 Cook's Distance	2
3	Influence Diagnostics for lme objects.	2
	3.1 Preparatory Steps	2
4	Influence Diagnostics	6
5	20.3.2.3. Cook's Distance for the $\beta$ estimates	13
6	Simulation of the Dependent Variable	16

### 1 Introduction

In this section we aim to replicate the methods proposed by Galecki et al and transfer to them to the models fitted by Roy's methodology fitted to the Blood data, as used in Bland and Altman's papers.

### 1.1 nlmeU R package

This monograph will use functions from the nlmeU package (Galecki and Burzykowski). This package consists of training datasets and utility functions enhancing functionality of nlme package.

logLik1.lme Calculates contribution of one subject to the log-likelihood for lme object

Pwr Calculates power based on a model fit

sigma Extract scale parameter sigma from a model fit

simulateY Simulates values of the dependent variable based on a model fit

#### 1.2 ARMD

Galecki and Burzykowski use in their examples the ARMD data set, sata from Age-Related Macular Degeneration (ARMD) clinical trial. The ARMD data arise from a randomized multicenter clinical trial comparing an experimental treatment (interferon-alpha) versus placebo for patients diagnosed with ARMD.

### 2 Back Ground Materials

Galecki and Burzykowski revise several core topics before proceeding to introduce their proposed methodology.

#### 2.1 Cook's Distance

# 3 Influence Diagnostics for lme objects.

- Cook's Distance
- Likelihood Distance

#### 3.1 Preparatory Steps

```
> summary(fm16.5)
Linear mixed-effects model fit by REML
Data: armd
AIC BIC logLik
6444.9 6483 -3214.5
Random effects:
Formula: "time | subject
Structure: Diagonal
(Intercept)
              time Residual
StdDev:
            7.2357 0.28102
                              5.0391
Variance function:
Structure: Power of variance covariate
Formula: ~time
Parameter estimates:
power
0.11052
Fixed effects: list(lm3.form)
Value Std.Error DF t-value p-value
              5.4416
                        2.26187 632
(Intercept)
                                      2.4058 0.0164
                        0.03822 231 23.5464 0.0000
visual0
               0.8998
             -0.2416
                        0.02392 632 -10.0997 0.0000
treat.fActive -2.6553
                        1.12868 231 -2.3525 0.0195
Correlation:
(Intr) visul0 time
             -0.934
visual0
             -0.071
time
                     0.002
treat.fActive -0.270 0.026 -0.002
Standardized Within-Group Residuals:
Min
           Q1
                   Med
                               Q3
                                        Max
```

### -4.148427 -0.329544 0.051823 0.444189 2.930748

Number of Observations: 867

Number of Groups: 234

- The formula() can recall the definition of the model defining the mean structure.
- Auxillary Function logLik1() which is designed to calculate a contribution of a given subject to the overall likelihood for a given model.
- The number of degrees of freedom reported by loglik is equal to 8. This corresponds to the total number of parameters in the model.

#### Part A

```
<- update(fm16.5)
### code chunk: Chap20.3influence_init
options(width = 65, digits = 5, show.signif.stars = FALSE)
date()
packageVersion("nlmeU")
packageVersion("nlme")
packageVersion("lattice")
sessionInfo()
require(nlme)
require(lattice)
data(armd, package="nlmeU")
## Model M16.5
lm3.form <-</pre>
                       # (12.9, 16.17)
formula(visual ~ visual0 + time + treat.f)
fm16.5 <-
                       # R16.13
lme(lm3.form,
random = list(subject = pdDiag(~time)),
weights =varPower(form=~time),
data = armd)
logLik()
```

```
beta0 <- fixef(fm16.5ml)
beta0

colnames(vcovb) <- names(beta0)
vcovb</pre>
```

We extract the  $\beta$  estimates and their estimated variance covariance matrix.

Towards this end, we use the functions fixef() and vcov() respectively, we can save these estimates and the matrix as objects beta0 and vcovb respectively.

With the help of the abbreviate function the names of the beta estimates are shortened to simplify the display of content.

An auxillary function logLik1()

we used the logLik() function to obtain the value of the log-likelihood for the fitted model. It should be noted that the function returns the log-likelihood evaluated at the set of the

The auxillary function logLik1() has been included in the package nlmeU.

estimated fixed effects and variance-covariance parameters and for the data set, to which the model is fitted. In the context of influence diagnostics, we need a more general function that allows

modfit an object of class *lme* representing the lme model fited to a given dataset using ML estimation.

dt1 a dataframe with data for one subject, for whom the likelihood function to be evaluated

dtInit an optional auxillary data frame

The dataframe provided in the argument <code>dt1</code> is typically created by choosing a subset with one subject from the data used to obtain the model fit object specified in the <code>modfit</code> argument. However, in general, any pluasible fata for one subject , not necessarily from the dataset used to fit the model, can be used.

The auxillary data provided in the argument dtInit is temporarily appended to the dt1 data. during the logLik() function execution.

The logLik1() function returns the numeric contributions of the single subject, with the data specified in the dt1 argument, to the log likelihood for the model specified in the modfit argument.

Contributions of Individual Subjects to the log-likelihood for fitted model

```
lLik.i <- as.vector(lLik.i)

lLik.i[1:5]
sum(lLik.i)</pre>
```

Plot of individual contributions to the log-likelihood (traditional graphics)

```
subject.c <-
subject.x <-
plot()
points()
text()</pre>
```

next we use the function logLik1() to compute the loglikelihood contributions for all subjects.

```
### code chunk: R20.7a
require(nlmeU)
df1 <- subset(armd, subject %in% "1")  # Data for subject "1"
logLik1(fm16.5ml, df1)  # logLik_i for subject</pre>
logLik1(fm16.5ml, df1)
                                    # logLik_i for subject "1"
lLik.i <- by(armd, armd$subject,</pre>
FUN = function(dfi) logLik1(fm16.5ml, dfi))
lLik.i <- as.vector(lLik.i) # Coerse array to vector</pre>
                     # logLik_i for the first five subjects
# Sum logLik_i; compare to Panel 20.6a
lLik.i[1:5]
sum(lLik.i)
### code chunk: R20.7b
nx <- by(armd, armd$subject, nrow)</pre>
lLik.n <- lLik.i/as.vector(nx)</pre>
                                    # logLiki
outL \leftarrow llik.n \leftarrow -6
                                    # TRUE for values < -6
lLik.n[outL]
                                    # logLiki/ni < -6
subject.c <- levels(armd$subject)</pre>
subject.x <- as.numeric(subject.c)</pre>
plot(lLik.n ~ subject.x, type = "h") # Fig. 20.1
points(subject.x[outL], lLik.n[outL], type = "p", pch = 16)
text(subject.x[outL], lLik.n[outL], subject.c[outL])
### code chunk: R20.8a
lmeU <- function(cx) {</pre>
dfU <- subset(armd, subject != cx)  # LOO data</pre>
update(fm16.5ml, data = dfU)
                                  # L00 fit
}
```

We present the syntax to plot the per-observation individual log-likelihood contributions. First, with the help of the by() function, we create the array nx, which contains the number of observations

## 4 Influence Diagnostics

We use the results of the preparatory steps to perform influence-diagnostic calculations for the model. More specifically we evaluate the influence of every subject included in the data set.

We create a list containing the results of fitted model the "leave-one-subject-out" (LOO) datasets and explore its contents.

We define the function lmeU(), which fits the model to the data from the armd

when the function lmeU() is executed, and LOO data frame, named dfU, is created with the subject, indicated by the cx argument.

Subsequently model is fitted by dfI by applying the

next, with the help of the function lapply(), we apply the lmeU() to the consecutive elements of the character vector subject.c.

As a result, we obtain the list lmeUall, with lme-class model fit objects as elements. The model-fit objects contains the result of fitting model M16.5.

Finally, we name the components of the lmeUall() list using the subjects idenifier stored in the vector subject.c.

This technique is computationally expensive, as it required the model to be fitted m number of times, omitting one of the m cases each time.

Execution time can be improved if we decided to perform a reduced number of likelihood iterations, instead of performing iterations until there is convergence.

The values, based on the first few iterations, are expected to give a fairly good approximation of the LOO estimates.

The names of the first six components are printed out using the function names().

To extract the LOO data frame for, e.g., the subject "6", we refer to the "6" component of the lmeUall list.

The extracted data frame is stored in the object dataU6. By using the function dim() we can check the dimensions of the data frame.

Model is fitted to a sequence of "leave-one-subject-out" out data sets.

```
# back - what is cx?
# cx is Case identity
lmeU <- function(cx){
dfU <- subset(myData, subject !=cx)
update(mymodel,data=dfU)
}

# what is lmeU?
lmeUall <- lapply(subject.c, lmeU)</pre>
```

Exploring the contents of the lmeUall object.

```
names(lmeUall)
dataU6 <- lmeUall[["6"]]$data
dim(dataU6)
unique(dataU6$subject)[1:6]</pre>
```

```
### code chunk: R20.6a
fm16.5ml <- update(fm16.5, method = "ML") # ML estimation</pre>
                              # Recall model formula.
formula(fm16.5ml)
fm16.5ml$call$data
                              # Recall data name.
logLik(fm16.5ml)
                              # Log-likelihood value
### code chunk: R20.6b
beta0 <- fixef(fm16.5ml)</pre>
                              # beta
names(beta0)
                              # Long names
names(beta0) <- abbreviate(names(beta0), minlength = 7) # Short names</pre>
                             # beta printed.
vcovb <- vcov(fm16.5ml)
                              # vcovb
colnames(vcovb) <- names(beta0)</pre>
                             # Short names
vcovb
                              # vcovb printed.
```

Galecki presents the code used to calculate and plot individial likelihood displacements.

For an LMM, it is required the computation of the full log-likelihood for  $\hat{\theta}$ , the ML estimate for  $\theta$  obtained by fitting the model to all data, and for  $\hat{\theta}_{(-i)}$ , the ML estimate obtained by fitting the model to the data with the i-th subject excluded.

Note that both values of the log-likelihood, used in the definition of the likelihood displacement, should be calculated taking into account all observations.

Galeck creates an auxiliary function <code>llik()</code> which , for a given subject indicated by the main argument, extracts the lime model fit object for the corresponding LOO data.

The corresponding log-likelihood function is extracted from the lmeU with the help of the logLik()

The returned value <code>lLikU + lLik.s</code> is the log-likelihood evaluated for all observations, using the displaced estimates of the model parameters.

Calculation of the likelihood displacement

```
lLik <- function(cx)
  {
  lmeU <- lmeUall[[cx]]
  lLikU <- logLik(lmeU, REML = FALSE)
  df.s <- subset(armd, subject == cx)

  lLik.s <- logLik1(lmeU,df.s)
  return(lLikU + lLik.s)
  }

lLikUall <- apply(subject.c,lLik)</pre>
```

```
lLik <- function(cx){
lmeU <- lmeUall[[cx]]  # LOO fit extracted
lLikU <- logLik(lmeU, REML = FALSE) # LOO log-likelihood
df.s <-  # Data for subject cx...
subset(armd, subject == cx)
lLik.s <- logLik1(lmeU, df.s)  # ...and log-likelihood.
return(lLikU + lLik.s)  # "Displaced" log-likelihood...
}
lLikUall <- sapply(subject.c, lLik)  # ...for all subjects.

dif.2Lik <- 2*(logLik(fm16.5ml) - lLikUall) # Vector of LDi
summary(dif.2Lik)</pre>
```

```
names(dif.2Lik) <- subject.c #subjects ids assigned
outL <- dif.2Lik > 0.5

dif.2Lik[outL]

## Plot Component

libary(lattice)

subject.f <- factor(subject.c, levels = subject.c)

myPanel <- function(x,y, ...){
    x1 <- as.numeric(x)
    panel.xyplot(x1,y, ...)
    ltext(x1[outL],y[outL], subject.c[outL]) # outlying LDis
    }

dtp <- dotplot(dif.2Lik ~subject.f, panel = myPanel, type= "h")

# ggplot?
lxlims <- length(dtp$x.limits)

update(dtp,xlim=rep(" ", lxlims),grid= "h")</pre>
```

By applying the **summary()** function to the vector, we obtain summary statistics of the computed likelihood-displacement values.

we create the logical output vector outL which indicates the subjects with the values of the likelihood displacement exceeding say 0.5.

From the printout of the selected elements of the vectir dif.2Lik it follows that there are seven such subjects.

We then use the function dotplot() from the package lattice to plot the likelihood distance value for all subjects.

The x-axis of the plot is constructed using numeric representation of the numeric.f factor, containing values ranging from 1 to 234.

```
### code chunk: R20.9b
names(dif.2Lik) <- subject.c
                                    # Subjects' ids assigned
outL <- dif.2Lik > 0.5
                                    # Outlying LDi's
dif.2Lik[outL]
library(lattice)
subject.f <- factor(subject.c, levels = subject.c)</pre>
myPanel <- function(x, y, ...){</pre>
x1 <- as.numeric(x)</pre>
panel.xyplot(x1, y, ...)
ltext(x1[outL], y[outL], subject.c[outL]) # Label outlying LDi
dtp <-
                                        # Fig. 20.2
dotplot(dif.2Lik ~ subject.f, panel = myPanel, type = "h")
lxlims <- length(dtp$x.limits)</pre>
update(dtp, xlim = rep("", lxlims), grid = "h")
```

# 5 20.3.2.3. Cook's Distance for the $\beta$ estimates

Cook's distance for the  $\beta$  estimates was defined in (4.26) for the classical LM. The definition can be extended to the LMMs in a straightforward manner.

Part 2 plot of cook's distance using traditional graphics

```
CookD.num <- apply(betaUall, 2, CookDfun)

(n.fixeff <- length(betaO))  # Number of fixed effects
rankX <- n.fixeff  # Rank of matrix X

CookD <- CookD.num/rankX  # Cook's distance Di
```

```
### Blood Data Equivalent
betaUall <- sapply(lmeUall,fixef)

vb.inv <- solve(vcovb)

cookDfun <- function(betaU){
   dbetaU <- betaU - betaO
   cookD.value <- t(dbetaU) %*% vb.inv %*% dbetaU
  }</pre>
```

Plot of Cook's Distance using traditional graphics. Outlying values annotated.

```
### Blood Data Equivalent
outD <- cookD > 0.03

#Create the Stick plot
plot(CookD ~ subject.c,
  ylab="Cook's Distance", type="h")
```

text()
points()

Cook's distance for the beta estimates were defined by 4.26 for the classical LM model.

Next we compute the inverse of the variance covariance matrix  $\hat{\beta}$  using the solve() function. We store the resulting matrix in the object vb.inv.

Subsequently we define the function cookDfun() which, for a vector given in the betaU argument, computes the value of the numerator of Cook's Distance, as in (4.26).

The function is then applied sequentially to all columns of the matrix betaUall.

The resultant vector is divided by the number of the fixed effects coefficients, which, under the assumption that the design matrix is of full rank, is equivalent to the rank of the design matrix.

The outcome is stored in the vector **cookD** and contains the values of cooks distanced for all subjects.

We create the logical vector out D which indicates the subjects with cook's distance values that exceed 0.03.

Present the scatterplot matrix of the two-dimensional projections of the differences. for all pairs of the fixed-effects coefficients.

The plot was generated using the splom function. The main argument of the function was obtained by subtracting the beta0 vector from the rows of the transposed betaUall matrix.

The labels used in the panels located on the diagonal of the figure provide the fixed estimates of fixed effects coefficients of the model and their estimates SEs. The panels above the diagonal include points for all subjects. The points for non-outlying values are plotted using small size open circles. The five outlying values are plotted using different plotting symbols defined in the legend of the figure at the top of the graph.

The effect of removal of this subject on the estimates of the remaining fixed-effects coefficients is relatively small. In contrast, removing the subject "227" affects the estimate of all fixed-effects coefficients to a different degree and in different directions.

More specifically, the intercept is driven towards lower values: the positive slope associated with visual acuity at baseline visual0, is further increased, the negative slope associated with time is brough closer to zero, and the treatment effect is attenuated.

Overall we note that the effect of removing any of the subjects on the fixed effects estimates is very small, as it does not exceed 0.05 of the SE of any of the estimates.

## 6 Simulation of the Dependent Variable

```
### Note: Simulations in Panel R20.11 take a long time
### code chunk: Chap20.4init
options(width = 65, digits = 5, show.signif.stars = FALSE)
date()
sessionInfo()
require(nlme)
data(armd, package="nlmeU")
lm3.form <-</pre>
                                  # (12.9)
formula(visual ~ visual0 + time + treat.f)
fm16.5 <-
            #update(fm16.4,
                                  # M16.5 <- M16.4
lme (lm3.form,
random = list(subject = pdDiag(~time)),
weights = varPower(form = ~time), #
data = armd)
fm16.5ml <- update(fm16.5, method = "ML")</pre>
### code chunk: R20.11
library(nlmeU)
simY <- simulateY(fm16.5ml, nsim = 1000, seed = 1238917) # Simulated y from M16.1</pre>
auxDt <- subset(armd,</pre>
                                   # Auxiliary data
select = c(subject, visual, visual0, time, treat.f))
simYsumm <-
apply(simY,
MARGIN = 2,
                           # Over columns
FUN = function(y){
auxDt$visual <- y</pre>
                         # Dependent variable updated
auxFit <-
                         # Update M16.1 with new y
update(fm16.5ml, data = auxDt)
summ <- summary(auxFit)</pre>
                        # Summary
beta <- fixef(summ)</pre>
list(beta = beta)
})
simYsumm[[1]]
                                   # beta for the 1st simulation
```

We cosnider simulations of the dependent variable, based on the marginal distribution implied by the fitted model. Towards this end, we have developed simulateY(), which can be used for objects of class lme.

We note that the function is different from simulate.lme(), available in the nlme in that the latter returns simulation-based REML and/or ML values and not the value of the dependent variable.

We demonstrate the use of the simulateY() function to create the empirical distribution of the  $\beta$  estimates. As an example, we consider the model which was fitted to the armd data.

Note however that the presented syntax is fairly general can be used for other LMEs as well. We apply the function simulateY() to the object fm16.5ml.

```
library(nlmeU)
simY <- simulateY(fm16.5ml, nsim=1000)

simYsumm <- apply( simY, MARGIN = 2,
  FUN = function(y){
     }
    )

simYsumm[[1]]

# \hat{\beta} for the 1st simulation</pre>
```

There is a second argument nsim that indicated how many simulations. The auxillary function performs the following steps

- the dependent variable visual is the auxDt data frame is replaced with a new set of simulated values contained in the vector y.
- Model M16.5 is fitted to the modified data frame

- the vector beta with the estimates of the fixed effects coefficients is extracted from the summary of the model-fit object with the help of the fixef() function.
- The vector of estimates is returned as a list with one component named beta.

They are drawn from the marginal distribution of the dependent variable implied by the fitted model.

It should be mentioned that the creation of the simYsumm list involves refitting the model many times, and it therefore takes a long time.

Toward this end, with the help of the sapply() function, we extract he vectors with the values of  $\hat{\beta}$  for each simulation for the list -object simYsumm and bind them column-wise into the betaE.

Then we use the function rowMeans() to compute the mean values of the columns (i.e. accross the rows) of betaE.