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# Chapter 1

## Other Material

### 1.1 Extension of Roy's methodology

Roy's methodology is constructed to compare two methods in the presence of replicate measurements. Necessarily it is worth examining whether this methodology can be adapted for different circumstances.

An implementation of Roy's methodology, whereby three or more methods are used, is not feasible due to computational restrictions. Specifically there is a failure to reach convergence before the iteration limit is reached. This may be due to the presence of additional variables, causing the problem of non-identifiability. In the case of two variables, it is required to estimate two variance terms and four correlation terms, six in all. For the case of three variabilities, three variance terms must be estimated as well as nine correlation terms, twelve in all. In general for  $n$  methods has  $2 \times T_n$  variance terms, where  $T_n$  is the triangular number for  $n$ , i.e. the addition analogue of the factorial. Hence the computational complexity quite increases substantially for every increase in  $n$ .

Should an implementation be feasible, further difficulty arises when interpreting the results. The fundamental question is whether two methods have close agreement so as to be interchangeable. When three methods are present in the model, the null

hypothesis is that all three methods have the same variability relevant to the respective tests. The outcome of the analysis will either be that all three are interchangeable or that all three are not interchangeable.

The tests would not be informative as to whether any two of those three were interchangeable, or equivalently if one method in particular disagreed with the other two. Indeed it is easier to perform three pair-wise comparisons separately and then to combine the results.

## 1.2 Roy's methodology for single measurements

Roy's methodology follows from the decomposition for the covariance matrix of the response vector  $y_i$ , as presented in Hamlett et al. (2004). The decomposition depends on the estimation of correlation terms, which would be absent in the single measurement case. Indeed there can be no within-subject variability if there are no repeated terms for it to describe. There would only be the covariance matrix of the measurements by both methods, which doesn't require the use of LME models. To conclude, simple existing methodologies would be the correct approach where there only one measurements by each method. Roy's methodology is not suitable for the case of single measurements because it follows from the decomposition for the covariance matrix of the response vector  $y_i$ , as presented in Hamlett et al. (2004). The decomposition depends on the estimation of correlation terms, which would be absent in the single measurement case. Indeed there can be no within-subject variability if there are no repeated terms for it to describe. There would only be the covariance matrix of the measurements by both methods, which doesn't require the use of LME models. To conclude, simpler existing methodologies, such as Deming regression, would be the correct approach where there only one measurements by each method.

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### 1.3 Correlation

Bivariate correlation coefficients have been shown to be of limited use in method comparison studies (Bland and Altman, 1986). However, recently correlation analysis has been developed to cope with repeated measurements, enhancing their potential usefulness. Roy incorporates the use of correlation into his methodology.

In addition to the variability tests, Roy advises that it is preferable that a correlation of greater than 0.82 exist for two methods to be considered interchangeable. However if two methods fulfil all the other conditions for agreement, failure to comply with this one can be overlooked. Indeed Roy demonstrates that placing undue importance to it can lead to incorrect conclusions. Roy (2009b) remarks that current computer implementations only gives overall correlation coefficients, but not their variances. Consequently it is not possible to carry out inferences based on all overall correlation coefficients.

### 1.4 Correlation terms

Hamlett et al. (2004) demonstrated how the between-subject and within subject variabilities can be expressed in terms of correlation terms.

$$D = \begin{pmatrix} \sigma_A^2 \rho_A & \sigma_A \sigma_b \rho_{AB} \delta \\ \sigma_A \sigma_b \rho_{AB} \delta & \sigma_B^2 \rho_B \end{pmatrix}$$

$$\mathbf{\Lambda} = \begin{pmatrix} \sigma_A^2(1 - \rho_A) & \sigma_{AB}(1 - \delta) \\ \sigma_{AB}(1 - \delta) & \sigma_B^2(1 - \rho_B) \end{pmatrix}.$$

$\rho_A$  describe the correlations of measurements made by the method  $A$  at different times. Similarly  $\rho_B$  describe the correlation of measurements made by the method  $B$  at different times. Correlations among repeated measures within the same method are known as intra-class correlation coefficients.  $\rho_{AB}$  describes the correlation of measurements taken at the same same time by both methods. The coefficient  $\delta$  is added for when the measurements are taken at different times, and is a constant of less than 1 for linked replicates. This is based on the assumption that linked replicates measurements taken at the same time would have greater correlation than those taken at different times. For unlinked replicates  $\delta$  is simply 1. Hamlett et al. (2004) provides a useful graphical depiction of the role of each correlation coefficients.



## 1.5 Hamlett and Lam

The methodology proposed by ? is largely based on Hamlett et al. (2004), which in turn follows on from Lam et al. (1999).

Hamlett re-analyses the data of Lam et al. (1999) to generalize their model to cover other settings not covered by the Lam method.

In many cases, repeated observation are collected from each subject in sequence and/or longitudinally.

$$y_i = \alpha + \mu_i + \epsilon$$

## 1.6 Limits of agreement in LME models

Limits of agreement are used extensively for assessing agreement, because they are intuitive and easy to use. Necessaire their prevalence in literature has meant that they are now the best known measurement for agreement, and therefore any newer methodology would benefit by making reference to them.

Carstensen et al. (2008) uses LME models to determine the limits of agreement. Between-subject variation for method  $m$  is given by  $d_m^2$  and within-subject variation is given by  $\lambda_m^2$ . Carstensen et al. (2008) remarks that for two methods  $A$  and  $B$ , separate values of  $d_A^2$  and  $d_B^2$  cannot be estimated, only their average. Hence the assumption that  $d_x = d_y = d$  is necessary. The between-subject variability  $\mathbf{D}$  and within-subject variability  $\mathbf{\Lambda}$  can be presented in matrix form,

$$\mathbf{D} = \begin{pmatrix} d_A^2 & 0 \\ 0 & d_B^2 \end{pmatrix} = \begin{pmatrix} d^2 & 0 \\ 0 & d^2 \end{pmatrix}, \quad \mathbf{\Lambda} = \begin{pmatrix} \lambda_A^2 & 0 \\ 0 & \lambda_B^2 \end{pmatrix}.$$

The variance for method  $m$  is  $d_m^2 + \lambda_m^2$ . Limits of agreement are determined using the standard deviation of the case-wise differences between the sets of measurements by two methods  $A$  and  $B$ , given by

$$\text{var}(y_A - y_B) = 2d^2 + \lambda_A^2 + \lambda_B^2. \quad (1.1)$$

Importantly the covariance terms in both variability matrices are zero, and no covariance component is present.

Carstensen et al. (2008) presents a data set ‘fat’, which is a comparison of measurements of subcutaneous fat by two observers at the Steno Diabetes Center, Copenhagen. Measurements are in millimeters (mm). Each person is measured three times by each observer. The observations are considered to be ‘true’ replicates.

A linear mixed effects model is formulated, and implementation through several software packages is demonstrated. All of the necessary terms are presented in the computer output. The limits of agreement are therefore,

$$0.0449 \pm 1.96 \times \sqrt{2 \times 0.0596^2 + 0.0772^2 + 0.0724^2} = (-0.220, 0.309). \quad (1.2)$$

Roy (2009a) has demonstrated a methodology whereby  $d_A^2$  and  $d_B^2$  can be estimated separately. Also covariance terms are present in both  $\mathbf{D}$  and  $\mathbf{\Lambda}$ . Using Roy’s methodology, the variance of the differences is

$$\text{var}(y_{iA} - y_{iB}) = d_A^2 + \lambda_B^2 + d_A^2 + \lambda_B^2 - 2(d_{AB} + \lambda_{AB}) \quad (1.3)$$

All of these terms are given or determinable in computer output. The limits of agreement can therefore be evaluated using

$$\bar{y}_A - \bar{y}_B \pm 1.96 \times \sqrt{\sigma_A^2 + \sigma_B^2 - 2(\sigma_{AB})}. \quad (1.4)$$

For Carstensen’s ‘fat’ data, the limits of agreement computed using Roy’s method are consistent with the estimates given by Carstensen et al. (2008);  $0.044884 \pm 1.96 \times 0.1373979 = (-0.224, 0.314)$ .

### 1.6.1 Variance Ratios

**Variance Ratios** The approach proposed by Roy deals with the question of agreement, and indeed interchangeability, as developed by Bland and Altman’s corpus of work. In the view of Dunn, a question relevant to many practitioners is which of the two methods is more precise.

The relationship between precision and the within-item and between-item variability must be established. Roy establishes the equivalence of repeatability and within-item variability, and hence precision. The method with the smaller within-item variability can be deemed to be the more precise. A useful approach is to compute the confidence intervals for the ratio of within-item standard deviations (equivalent to the ratio of repeatability coefficients), which can be interpreted in the usual manner. In fact, the ratio of within-item standard deviations, with the attendant confidence interval, can be determined using a single R command: `intervals()`. Pinheiro and Bates (pg 93-95) give a description of how confidence intervals for the variance components are computed. Furthermore a complete set of confidence intervals can be computed to complement the variance component estimates. What is required is the computation of the variance ratios of within-item and between-item standard deviations. A naive approach would be to compute the variance ratios by relevant F distribution quantiles. However, the question arises as to the appropriate degrees of freedom. Limits of agreement are easily computable using the LME framework. While we will not be considering this analysis, a demonstration will be provided in the example.

## 1.7 Testing Procedures

Roy's methodology requires the construction of four candidate models. The first candidate model is compared to each of the three other models successively. It is the alternative model in each of the three tests, with the other three models acting as the respective null models.

The probability distribution of the test statistic can be approximated by a chi-square distribution with  $(\nu_1 - \nu_2)$  degrees of freedom, where  $\nu_1$  and  $\nu_2$  are the degrees of freedom of models 1 and 2 respectively.

Likelihood ratio tests are very simple to implement in R, simply use the `'anova()'` commands. Sample output will be given for each variability test. The likelihood ratio test is the procedure used to compare the fit of two models. For each candidate model,

the ‘-2 log likelihood’ ( $M2LL$ ) is computed. The test statistic for each of the three hypothesis tests is the difference of the  $M2LL$  for each pair of models. If the  $p$ -value in each of the respective tests exceed as significance level chosen by the analyst, then the null model must be rejected.

$$-2 \ln \Lambda_d = [M2LL \text{ under } H_0 \text{ model}] - [M2LL \text{ under } H_A \text{ model}] \quad (1.5)$$

These test statistics follow a chi-square distribution with the degrees of freedom computed as the difference of the LRT degrees of freedom.

$$\nu = [\text{LRT df under } H_0 \text{ model}] - [\text{LRT df under } H_A \text{ model}] \quad (1.6)$$

```
> anova(MCS1,MCS2)
>
>
Model df      AIC      BIC logLik  Test L.Ratio p-value
MCS1    1  8 4077.5 4111.3 -2030.7
MCS2    2  7 4075.6 4105.3 -2030.8 1 vs 2 0.15291 0.6958
```

### 1.7.1 Roy's Reference Model

Conventionally LME models can be tested using Likelihood Ratio Tests, wherein a reference model is compared to a nested model.

```
> Ref.Fit = lme(y ~ meth-1, data = dat, #Symm , Symm#
+ random = list(item=pdSymm(~ meth-1)),
+ weights=varIdent(form=~1|meth),
+ correlation = corSymm(form=~1 | item/repl),
+ method="ML")
```

Roy(2009) presents two nested models that specify the condition of equality as required, with a third nested model for an additional test. There three formulations share the same structure, and can be specified by making slight alterations of the code for the Reference Model.

### 1.7.2 Nested Model (Between-Item Variability)

```
> NMB.fit = lme(y ~ meth-1, data = dat, #CS , Symm#
+ random = list(item=pdCompSymm(~ meth-1)),
+ correlation = corSymm(form=~1 | item/repl),
```

```
+     method="ML")
```

## 1.8 Classical model for single measurements

The classical model is based on measurements  $y_{mi}$  by method  $m = 1, 2$  on item  $i = 1, 2, \dots$

$$y_{mi} = \alpha_m + \mu_i + e_{mi}$$

$$e_{mi} \sim \mathcal{N}(0, \sigma_m^2)$$

Even though the separate variances can not be identified, their sum can be estimated by the empirical variance of the differences.

Like wise the separate  $\alpha$  can not be estimated, only their difference can be estimated as  $\bar{D}$

In the first instance, we require a simple model to describe a measurement by method  $m$ . We use the term *item* to denote an individual, subject or sample, to be measured, being randomly sampled from a population. Let  $y_{mi}$  be the measurement for item  $i$  made by method  $m$ .

$$y_{mi} = \alpha_m + \mu_i + e_{mi}$$

- $\alpha_m$  is the fixed effect associated with method  $m$ ,
- $\mu_i$  is the true value for subject  $i$  (fixed effect),
- $e_{mi}$  is a random effect term for errors with  $e_{mi} \sim \mathcal{N}(0, \sigma_m^2)$ .

.

This model implies that the difference between the paired measurements can be expressed as

$$d_i = y_{1i} - y_{2i} \sim \mathcal{N}(\alpha_1 - \alpha_2, \sigma_1^2 + \sigma_2^2).$$

Importantly, this is independent of the item levels  $\mu_i$ . As the case-wise differences are of interest, the parameters of interest are the fixed effects for methods  $\alpha_m$ .

$$y_{mi} = \alpha_m + \mu_i + e_{mi}$$



Importantly these variance covariance structures are central to Roy methodology.

? proposes a series of hypothesis tests based on these matrices as part of her methodology. These tests shall be reverted to in due course.

The standard deviation of the differences of variables  $a$  and  $b$  is computed as

$$\text{var}(a - b) = \text{var}(a) + \text{var}(b) - 2\text{cov}(a, b)$$

Hence the variance of the difference of two methods, that allows for the calculation of the limits of agreement, can be calculated as

$$\text{var}(d) = \omega_1^2 + \omega_2^2 - 2 \times \omega_1 \omega_2$$

### 1.8.1 Sampling

*One important feature of replicate observations is that they should be independent of each other. In essence, this is achieved by ensuring that the observer makes each measurement independent of knowledge of the previous value(s). This may be difficult to achieve in practice. (Check who said this )*

### 1.8.2 Remarks on the Multivariate Normal Distribution

Diligence is required when considering the models. Carstensen specifies his models in terms of the univariate normal distribution. Roy's model is specified using the bivariate normal distribution. This gives rise to a key difference between the two models, in that a bivariate model accounts for covariance between the variables of interest. The multivariate normal distribution of a  $k$ -dimensional random vector  $X = [X_1, X_2, \dots, X_k]$  can be written in the following notation:

$$X \sim \mathcal{N}(\mu, \Sigma),$$

or to make it explicitly known that  $X$  is  $k$ -dimensional,

$$X \sim \mathcal{N}_k(\mu, \Sigma).$$

with  $k$ -dimensional mean vector

$$\mu = [E[X_1], E[X_2], \dots, E[X_k]]$$

and  $k \times k$  covariance matrix

$$\Sigma = [\text{Cov}[X_i, X_j]], \quad i = 1, 2, \dots, k; \quad j = 1, 2, \dots, k$$

#### 1. Univariate Normal Distribution

$$X \sim \mathcal{N}(\mu, \sigma^2),$$

#### 2. Bivariate Normal Distribution

(a)

$$X \sim \mathcal{N}_2(\mu, \Sigma),$$

(b)

$$\mu = \begin{pmatrix} \mu_x \\ \mu_y \end{pmatrix}, \quad \Sigma = \begin{pmatrix} \sigma_x^2 & \rho\sigma_x\sigma_y \\ \rho\sigma_x\sigma_y & \sigma_y^2 \end{pmatrix}.$$

## 1.9 Demonstration of Roy's testing

Roy provides three case studies, using data sets well known in method comparison studies, to demonstrate how the methodology should be used. The first two examples used are from the 'blood pressure' data set introduced by Bland and Altman (1999). The data set is a tabulation of simultaneous measurements of systolic blood pressure were made by each of two experienced observers (denoted 'J' and 'R') using a sphygmomanometer and by a semi-automatic blood pressure monitor (denoted 'S'). Three sets of readings were made in quick succession. Roy compares the 'J' and 'S' methods in his first example, and the 'R' and 'S' methods in his second.

### 1.9.1 Matrix structures

Before discussing the tests, it is useful to point out the difference between symmetric form and compound symmetry form. Consider a generic matrix  $A$ ,

$$A = \begin{pmatrix} a_{11} & a_{12} \\ a_{21} & a_{22} \end{pmatrix}. \quad (1.7)$$

A symmetric matrix allows the diagonal terms  $a_{11}$  and  $a_{22}$  to differ. The compound symmetry structure requires that both of these terms be equal, i.e  $a_{11} = a_{22}$ .

### 1.9.2 Variability test 1

This is a test on whether both methods  $A$  and  $B$  have the same between-subject variability or not.

$$H_0 : d_A = d_B \quad (1.8)$$

$$H_A : d_A \neq d_B \quad (1.9)$$

When implemented using **R**, this test is facilitated by constructing a model specifying a symmetric form for  $D$  (i.e. the alternative model) and comparing it with a model

that has compound symmetric form for  $D$  (i.e. the null model). For this test  $\hat{\Lambda}$  has a symmetric form for both models, and will be the same for both.

### Bland-Altman's blood data

With the alternative model, the MLE of the between-subject variance covariance matrix is given by

$$\hat{D}_{Symm} = \begin{pmatrix} 923.98 & 785.24 \\ 785.24 & 971.30 \end{pmatrix} \quad (1.10)$$

With the null model the MLE is as follows:

$$\hat{D}_{CS} = \begin{pmatrix} 946.50 & 784.32 \\ 784.32 & 946.50 \end{pmatrix} \quad (1.11)$$

A likelihood ratio test is perform to determine which model is more suitable. The outcome of this test is presented in the following **R** code.

```
> anova(MCS1,MCS2)
>
>
Model df      AIC      BIC  logLik   Test L.Ratio p-value
MCS1    1  8 4077.5 4111.3 -2030.7
MCS2    2  7 4075.6 4105.3 -2030.8 1 vs 2 0.15291 0.6958
```

The test statistic is the difference of the  $-2$  log likelihoods; 0.15291. The  $p$ -value is 0.6958. Therefore we fail to reject the hypothesis that both have the same between-subject variabilities.

### 1.9.3 Variability test 2

This is a test on whether both methods  $A$  and  $B$  have the same within-subject variability or not.

$$H_0 : \lambda_A = \lambda_B \quad (1.12)$$

$$H_A : \lambda_A = \lambda_B \quad (1.13)$$

This model is performed in the same manner as the first test, only reversing the roles of  $\hat{\mathbf{D}}$  and  $\hat{\mathbf{\Lambda}}$ . The null model is constructed a symmetric form for  $\hat{\mathbf{\Lambda}}$  while the alternative model uses a compound symmetry form. This time  $\hat{\mathbf{D}}$  has a symmetric form for both models, and will be the same for both.

### Bland-Altman's blood data

For the null model the MLE of the within-subject variance covariance matrix is given below.

$$\hat{\mathbf{\Lambda}}_{\text{Symm}} = \begin{pmatrix} 37.40 & 16.06 \\ 16.06 & 83.14 \end{pmatrix} \quad (1.14)$$

With the alternative model the MLE is as follows:

$$\hat{\mathbf{\Lambda}}_{CS} = \begin{pmatrix} 60.27 & 16.06 \\ 16.06 & 60.27 \end{pmatrix} \quad (1.15)$$

A likelihood ratio test is perform to determine which model is more suitable. The outcome of this test is that it can be assumed that they have equal The test statistic is the difference of the  $-2 \log$  likelihoods; 28.617. The  $p$ -value is less than 0.0001. In this case we reject the null hypothesis that both models have the same within-subject variabilities.

### 1.9.4 Variability test 3

This is a test on whether both methods  $A$  and  $B$  have the same overall variability or not.

$$H_0 : \sigma_A = \sigma_B \quad (1.16)$$

$$H_A : \sigma_A = \sigma_B \quad (1.17)$$

The null model is constructed a symmetric form for both  $\hat{\mathbf{D}}$  and  $\hat{\mathbf{\Lambda}}$  while the alternative model uses a compound symmetry form for both.

#### Bland-Altman's blood data

With the null model the MLE of the within-subject variance covariance matrix is given below.

$$\hat{\Sigma}_{Symm} = \begin{pmatrix} 961.38 & 801.40 \\ 801.40 & 1054.43 \end{pmatrix} \quad (1.18)$$

With the alternative model the MLE is as follows:

$$\hat{\Sigma}_{CS} = \begin{pmatrix} 1007.92 & 801.65 \\ 801.65 & 1007.92 \end{pmatrix} \quad (1.19)$$

Again a likelihood ratio test is used to determine the most suitable of the two candidate models. The test statistic is the difference of the  $-2 \log$  likelihoods; 28.884. The  $p$ -value is less than 0.0001. We again reject the null hypothesis. Each model has a different overall variability, a foregone conclusion from the second variability test.

### 1.9.5 Test for inter-method bias

The inter-method bias between the two method is found to be 15.62 , with a  $p$ -value of

### 1.9.6 Correlation Test

$$\hat{\mathbf{r}}_{\Omega_{ii}} = \begin{pmatrix} 1 & 0.7959 \\ 0.7959 & 1 \end{pmatrix} \quad (1.20)$$

The diagonal blocks  $\hat{\mathbf{r}}_{\Omega_{ii}}$  of the correlation matrix indicate an overall coefficient of 0.7959. This is less than the threshold of 0.82 that Roy recommends.

The off diagonal blocks of the overall correlation matrix  $\hat{\mathbf{r}}_{\Omega_{ii'}}$  are

$$\hat{\mathbf{r}}_{\Omega_{ii'}} = \begin{pmatrix} 0.9611 & 0.7799 \\ 0.7799 & 0.9212 \end{pmatrix}. \quad (1.21)$$

### 1.9.7 Conclusion of procedure

The overall conclusion of the procedure is that the two methods are not in agreement, specifically due to the within-subject variability, and the inter-method bias. The repeatability coefficients are substantially different, one being 49% larger than the other. Additionally the overall correlation coefficient did not exceed the recommended threshold of 0.82.



## 1.10 Basic Models Fits

Further to Pinheiro and Bates (1994), several simple LME models are constructed for the blood pressure data. This data set is the subject of a method comparison study in Bland and Altman (1999).

### 1.10.1 Implementing the Mixed Models Fits

They are implemented using the following R code, utilising the ‘nlme’ package. An analysis of variance is used to compare the model fits.

The R script:

```
fit1 = lme( BP ~ method, data = dat, random = ~1 | subject )
fit2 = update(fit1, random = ~1 | subject/method )
fit3 = update(fit1, random = ~method - 1 | subject )
#analysis of variance
anova(fit1,fit2,fit3)
```

1. Simplest workable model, allows differences between methods and incorporates a random intercept for each subject. For subject 1 we have

$$\mathbf{X}_i = \begin{pmatrix} 1 & 0 \\ 1 & 0 \\ 1 & 0 \\ 1 & 1 \\ 1 & 1 \\ 1 & 1 \\ 1 & 1 \end{pmatrix}, \quad \boldsymbol{\beta} = \begin{pmatrix} \beta_0 \\ \beta_1 \end{pmatrix}, \quad \mathbf{Z}_i = \begin{pmatrix} 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \\ 1 \end{pmatrix}, \quad \mathbf{b}_i = b$$

where  $E(b) = 0$  and  $\text{var}(b) = \psi$ .

2.

$$\mathbf{Z}_i = \begin{pmatrix} 1 & 0 \\ 1 & 0 \\ 1 & 0 \\ 0 & 1 \\ 0 & 1 \\ 0 & 1 \end{pmatrix} \quad \mathbf{b}_i = \begin{pmatrix} b_1 & 0 \\ 0 & b_2 \end{pmatrix}$$

where  $E(b_i) = 0$  and  $\text{var}(\mathbf{b}) = \mathbf{\Psi}$ .

The variance of error terms is a  $6 \times 6$  matrix.

### 1.10.2 Model Fit 1

This is a simple model with no interactions. There is a fixed effect for each method and a random effect for each subject.

$$y_{ijk} = \beta_j + b_i + \epsilon_{ijk}, \quad i = 1, \dots, 2, j = 1, \dots, 85, k = 1, \dots, 3$$

$$b_i \sim \mathcal{N}(0, \sigma_b^2), \quad \epsilon_i \sim \mathcal{N}(0, \sigma^2)$$

Linear mixed-effects model fit by REML

Data: dat

Log-restricted-likelihood: -2155.853

Fixed: BP ~ method

(Intercept)      methodS

127.40784      15.61961

Random effects:

Formula: ~1 | subject

(Intercept) Residual

StdDev:      29.39085 12.44454

Number of Observations: 510

Number of Groups: 85

The following output was obtained.

Linear mixed-effects model fit by REML

Data: dat

Log-restricted-likelihood: -2047.582

Fixed: BP ~ method

(Intercept)      methodS

127.40784      15.61961

Random effects:

Formula: ~method - 1 | subject

Structure: General positive-definite, Log-Cholesky parametrization

StdDev      Corr

methodJ   30.455093   methdJ

methodS   31.477237   0.835

Residual   7.763666

Number of Observations: 510

Number of Groups: 85

### 1.10.3 Model Fit 1

This is a simple model with no interactions. There is a fixed effect for each method and a random effect for each subject.

$$y_{ijk} = \beta_j + b_i + \epsilon_{ijk}, \quad i = 1, \dots, 2, j = 1, \dots, 85, k = 1, \dots, 3$$

$$b_i \sim \mathcal{N}(0, \sigma_b^2), \quad \epsilon_i \sim \mathcal{N}(0, \sigma^2)$$

Linear mixed-effects model fit by REML

Data: dat

Log-restricted-likelihood: -2155.853

Fixed: BP ~ method

(Intercept)      methodS

127.40784      15.61961

Random effects:

Formula: ~1 | subject

(Intercept) Residual

StdDev:      29.39085 12.44454

Number of Observations: 510

Number of Groups: 85

The following output was obtained.

Linear mixed-effects model fit by REML

Data: dat

Log-restricted-likelihood: -2047.582

Fixed: BP ~ method

(Intercept)      methodS

127.40784      15.61961

Random effects:

Formula: ~method - 1 | subject

Structure: General positive-definite, Log-Cholesky parametrization

StdDev      Corr

methodJ   30.455093   methdJ

methodS   31.477237   0.835

Residual   7.763666

Number of Observations: 510

Number of Groups: 85

### 1.10.4 Model Fit 2

This is a simple model, this time with an interaction effect. There is a fixed effect for each method. This model has random effects at two levels  $b_i$  for the subject, and another,  $b_{ij}$ , for the respective method within each subject.

$$y_{ijk} = \beta_j + b_i + b_{ij} + \epsilon_{ijk}, \quad i = 1, \dots, 2, j = 1, \dots, 85, k = 1, \dots, 3$$

$$b_i \sim \mathcal{N}(0, \sigma_1^2), \quad b_{ij} \sim \mathcal{N}(0, \sigma_2^2), \quad \epsilon_i \sim \mathcal{N}(0, \sigma^2)$$

In this model, the random interaction terms all have the same variance  $\sigma_2^2$ . These terms are assumed to be independent of each other, even within the same subject.

Linear mixed-effects model fit by REML

Data: dat

Log-restricted-likelihood: -2047.714

Fixed: BP ~ method

(Intercept)      methodS

127.40784      15.61961

Random effects:

Formula: ~1 | subject

(Intercept)

StdDev:      28.28452

Formula: ~1 | method %in% subject

(Intercept) Residual

StdDev:      12.61562 7.763666

Number of Observations: 510

Number of Groups:

subject method %in% subject

85

170



### 1.10.5 Model Fit 3

This model is a more general model, compared to 'model fit 2'. This model treats the random interactions for each subject as a vector and allows the variance-covariance matrix for that vector to be estimated from the set of all positive-definite matrices.  $\mathbf{y}_i$  is the entire response vector for the  $i$ th subject.  $\mathbf{X}_i$  and  $\mathbf{Z}_i$  are the fixed- and random-effects design matrices respectively.

$$\mathbf{y}_i = \mathbf{X}_i\boldsymbol{\beta} + \mathbf{Z}_i\mathbf{b}_i + \boldsymbol{\epsilon}_i, \quad i = 1, \dots, 85$$

$$\mathbf{Z}_i \sim \mathcal{N}(\mathbf{0}, \boldsymbol{\Psi}), \quad \boldsymbol{\epsilon}_i \sim \mathcal{N}(\mathbf{0}, \sigma^2 \boldsymbol{\Lambda})$$

For the first subject the response vector,  $\mathbf{y}_1$ , is:

observation	BP	subject	method	replicate
1	100.00	1	J	1
86	106.00	1	J	2
171	107.00	1	J	3
511	122.00	1	S	1
596	128.00	1	S	2
681	124.00	1	S	3

The fixed effects design matrix  $\mathbf{X}_i$  is given by:

(Intercept)	method S
1	0
1	0
1	0
1	1
1	1
1	1

The random effects design matrix  $\mathbf{Z}_i$  is given by:

method J	method S
1	0
1	0
1	0
0	1
0	1
0	1

### 1.10.6 Using LME models to create Prediction Intervals

Carstensen (2004) also advocates the use of linear mixed models in the study of method comparisons. The model is constructed to describe the relationship between a value of measurement and its real value. The non-replicate case is considered first, as it is the context of the Bland Altman plots. This model assumes that inter-method bias is the only difference between the two methods. A measurement  $y_{mi}$  by method  $m$  on individual  $i$  is formulated as follows;

$$y_{mi} = \alpha_m + \mu_i + e_{mi} \quad (e_{mi} \sim N(0, \sigma_m^2)) \quad (1.22)$$

The differences are expressed as  $d_i = y_{1i} - y_{2i}$ . For the replicate case, an interaction term  $c$  is added to the model, with an associated variance component. All the random effects are assumed independent, and that all replicate measurements are assumed to be exchangeable within each method.

$$y_{mir} = \alpha_m + \mu_i + c_{mi} + e_{mir} \quad (e_{mi} \sim N(0, \sigma_m^2), c_{mi} \sim N(0, \tau_m^2)) \quad (1.23)$$

Carstensen et al. (2008) proposes a methodology to calculate prediction intervals in the presence of replicate measurements, overcoming problems associated with Bland-Altman methodology in this regard. It is not possible to estimate the interaction variance components  $\tau_1^2$  and  $\tau_2^2$  separately. Therefore it must be assumed that they are equal. The variance of the difference can be estimated as follows:

$$var(y_{1j} - y_{2j}) \quad (1.24)$$

### 1.10.7 Computation

Modern software packages can be used to fit models accordingly. The best linear unbiased predictor (BLUP) for a specific subject  $i$  measured with method  $m$  has the form  $BLUP_{mir} = \hat{\alpha}_m + \hat{\beta}_m \mu_i + c_{mi}$ , under the assumption that the  $\mu$ s are the true item values.

## 1.11 Limits of agreement in LME models

Limits of agreement are used extensively for assessing agreement, due to they're being intuitive and easy to use. Necessarily their prevalence in literature has meant that they are now the best known measurement for agreement, and that any newer methodology would benefit by making reference to them.

Carstensen et al. (2008) uses LME models to determine the limits of agreement. Between-subject variation for method  $m$  is given by  $d_m^2$  and within-subject variation is given by  $\lambda_m^2$ . Carstensen et al. (2008) remarks that for two methods  $A$  and  $B$ , separate values of  $d_A^2$  and  $d_B^2$  cannot be estimated, only their average. Hence the assumption that  $d_x = d_y = d$  is necessary. The between-subject variability  $\mathbf{D}$  and within-subject variability  $\mathbf{\Lambda}$  can be presented in matrix form,

$$\mathbf{D} = \begin{pmatrix} d_A^2 & 0 \\ 0 & d_B^2 \end{pmatrix} = \begin{pmatrix} d^2 & 0 \\ 0 & d^2 \end{pmatrix}, \quad \mathbf{\Lambda} = \begin{pmatrix} \lambda_A^2 & 0 \\ 0 & \lambda_B^2 \end{pmatrix}. \quad (1.25)$$

The variance for method  $m$  is  $d_m^2 + \lambda_m^2$ . Limits of agreement are determined using the standard deviation of the case-wise differences between the sets of measurements by two methods  $A$  and  $B$ , given by

$$\text{var}(y_A - y_B) = 2d^2 + \lambda_A^2 + \lambda_B^2. \quad (1.26)$$

Importantly the covariance terms in both variability matrices are zero, and no covariance component is present.

Carstensen et al. (2008) presents a data set ‘fat’, which is a comparison of measurements of subcutaneous fat by two observers at the Steno Diabetes Center, Copenhagen. Measurements are in millimeters (mm). Each person is measured three times by each observer. The observations are considered to be ‘true’ replicates.

A linear mixed effects model is formulated, and implementation through several software packages is demonstrated. All of the necessary terms are presented in the computer output. The limits of agreement are therefore,

$$0.0449 \pm 1.96 \times \sqrt{2 \times 0.0596^2 + 0.0772^2 + 0.0724^2} = (-0.220, 0.309). \quad (1.27)$$

Roy (2009a) has demonstrated a methodology whereby  $d_A^2$  and  $d_B^2$  can be estimated separately. Also covariance terms are present in both  $\mathbf{D}$  and  $\mathbf{\Lambda}$ . Using Roy's methodology, the variance of the differences is

$$\text{var}(y_{iA} - y_{iB}) = d_A^2 + \lambda_B^2 + d_A^2 + \lambda_B^2 - 2(d_{AB} + \lambda_{AB}) \quad (1.28)$$

All of these terms are given or determinable in computer output. The limits of agreement can therefore be evaluated using

$$\bar{y}_A - \bar{y}_B \pm 1.96 \times \sqrt{\sigma_A^2 + \sigma_B^2 - 2(\sigma_{AB})}. \quad (1.29)$$

For Carstensen's 'fat' data, the limits of agreement computed using Roy's method are consistent with the estimates given by Carstensen et al. (2008);  $0.044884 \pm 1.96 \times 0.1373979 = (-0.224, 0.314)$ .

### 1.11.1 Linked replicates

Carstensen et al. (2008) proposes the addition of an random effects term to their model when the replicates are linked. This term is used to describe the 'item by replicate' interaction, which is independent of the methods. This interaction is a source of variability independent of the methods. Therefore failure to account for it will result in variability being wrongly attributed to the methods.

Carstensen et al. (2008) introduces a second data set; the oximetry study. This study done at the Royal Children's Hospital in Melbourne to assess the agreement between co-oximetry and pulse oximetry in small babies.

In most cases, measurements were taken by both method at three different times. In some cases there are either one or two pairs of measurements, hence the data is unbalanced. Carstensen et al. (2008) describes many of the children as being very sick, and with very low oxygen saturations levels. Therefore it must be assumed that a biological change can occur in interim periods, and measurements are not true replicates.

Carstensen et al. (2008) demonstrate the necessity of accounting for linked replicated by comparing the limits of agreement from the 'oximetry' data set using a model

with the additional term, and one without. When the interaction is accounted for the limits of agreement are (-9.62,14.56). When the interaction is not accounted for, the limits of agreement are (-11.88,16.83). It is shown that the failure to include this additional term results in an over-estimation of the standard deviations of differences.

Limits of agreement are determined using Roy's methodology, without adding any additional terms, are found to be consistent with the 'interaction' model; (-9.562, 14.504). Roy's methodology assumes that replicates are linked. However, following Carstensen's example, an additional interaction term is added to the implementation of Roy's model to assess the effect, the limits of agreement estimates do not change. However there is a conspicuous difference in within-subject matrices of both models (denoted 1 and 2 respectively);

$$\begin{aligned} \hat{\boldsymbol{\Lambda}}_1 &= \begin{pmatrix} 16.61 & 11.67 \\ 11.67 & 27.65 \end{pmatrix} \\ \hat{\boldsymbol{\Lambda}}_2 &= \begin{pmatrix} 7.55 & 2.60 \\ 2.60 & 18.59 \end{pmatrix} \end{aligned}$$

The variance of the additional random effect in model 2 is 3.01.

The Akaike information criterion (AIC) for both of models are  $AIC_1 = 2304.226$  and  $AIC_2 = 2306.226$ . Having a difference of AIC values of 2 is equivalent to both models being equally as good as the other. The AIC values for the Carstensen 'unlinked' and 'linked' models are 1994.66 and 1955.48 respectively.

The  $\hat{\mathbf{A}}$  matrices are informative as to the difference between Carstensen's unlinked and linked models. For the oximetry data, the covariance terms (given above as 11.67 and 2.6 respectively) are of similar magnitudes to the variance terms. Conversely for the 'fat' data the covariance term (-0.00032) is negligible. When the interaction term

is added to the model, the covariance term remains negligible. (For the ‘fat’ data, the difference in AIC values is also 2).

To conclude, Carstensen’s models provided a rigorous way to determine limits of agreement, but don’t provide for the computation of  $\hat{\mathbf{D}}$  and  $\hat{\mathbf{\Lambda}}$ . Therefore the test’s proposed by Roy (2009a) can not be implemented. Conversely, accurate limits of agreement can be found using Roy’s method.

Addition of the interaction term erodes the capability of Roy’s methodology to compare candidate models, and therefore shall not be adopted.

(N.B. To complement the blood pressure ‘J vs S’ analysis, the limits of agreement are  $15.62 \pm 1.96 \times 20.33 = (-24.22, 55.46)$ .)



## 1.12 Conclusion

Carstensen et al. (2008) and Roy (2009a) highlight the need for method comparison methodologies suitable for use in the presence of replicate measurements. Roy (2009a) presents a comprehensive methodology for assessing the agreement of two methods, for replicate measurements. This methodology has the added benefit of overcoming the problems of unbalanced data and unequal numbers of replicates. Implementation of the methodology, and interpretation of the results, is relatively easy for practitioners who have only basic statistical training. Furthermore, it can be shown that widely used existing methodologies, such as the limits of agreement, can be incorporated into Roy's methodology.

# Chapter 2

## Influence Diagnostics

Cook (1986) introduces powerful tools for local-influence assessment and examining perturbations in the assumptions of a model. In particular the effect of local perturbations of parameters or observations are examined.

### 2.1 Influence Diagnostics: Basic Idea and Statistics

The general idea of quantifying the influence of one or more observations relies on computing parameter estimates based on all data points, removing the cases in question from the data, refitting the model, and computing statistics based on the change between full-data and reduced-data estimation.

Influence statistics can be coarsely grouped by the aspect of estimation that is their primary target:

- overall measures compare changes in objective functions: (restricted) likelihood distance (Cook and Weisberg 1982, Ch. 5.2)
- influence on parameter estimates: Cooks (Cook 1977, 1979), MDFFITS (Belsley, Kuh, and Welsch 1980, p. 32)
- influence on precision of estimates: CovRatio and CovTrace

- influence on fitted and predicted values: PRESS residual, PRESS statistic (Allen 1974), DFFITS (Belsley, Kuh, and Welsch 1980, p. 15)
- outlier properties: internally and externally studentized residuals, leverage

Furthermore, closed-form expressions for computing the change in important model quantities might not be available. This section provides background material for the various influence diagnostics available with the MIXED procedure. See the section Mixed Models Theory for relevant expressions and definitions. The parameter vector denotes all unknown parameters in the and matrix. The observations whose influence is being ascertained are represented by the set and referred to simply as "the observations in ." The estimate of a parameter vector, such as , obtained from all observations except those in the set is denoted . In case of a matrix , the notation represents the matrix with the rows in removed; these rows are collected in . If is symmetric, then notation implies removal of rows and columns. The vector comprises the responses of the data points being removed, and is the variance-covariance matrix of the remaining observations. When , lowercase notation emphasizes that single points are removed, such as .

## Residual Analysis for LME, Applications to MCS Data

This short section will look at residual analysis for LME models. The underlying assumptions for LME models are similar to those of classical linear models. There are two key techniques: a residual plot and the normal probability plot. Using the nlme package it is possible to create plots specific to each method. This is useful in determining which methods ‘disagree’ with the rest. Analysis of the residuals would determine if the methods of measurement disagree systematically, or whether or not erroneous measurements associated with a subset of the cases are the cause of disagreement. Erroneous measurements are incorrect measurements that indicate disagreement between methods that would otherwise be in agreement.

### 2.2 Deletion Diagnostics

Since the pioneering work of Cook in 1977, deletion measures have been applied to many statistical models for identifying influential observations.

Deletion diagnostics provide a means of assessing the influence of an observation (or groups of observations) on inference on the estimated parameters of LME models.

Data from single individuals, or a small group of subjects may influence non-linear mixed effects model selection. Diagnostics routinely applied in model building may identify such individuals, but these methods are not specifically designed for that purpose and are, therefore, not optimal. We describe two likelihood-based diagnostics for identifying individuals that can influence the choice between two competing models.

Case-deletion diagnostics provide a useful tool for identifying influential observations and outliers.

The computation of case deletion diagnostics in the classical model is made simple by the fact that estimates of  $\beta$  and  $\sigma^2$ , which exclude the  $i$ th observation, can be computed without re-fitting the model. Such update formulas are available in the mixed model only if you assume that the covariance parameters are not affected by the removal of the observation in question. This is rarely a reasonable assumption.

### 3. Case Deletion Diagnostics for LME Data: Cooks Distance, DFBetas

In this section we introduce influence analysis and case deletion diagnostics. A full overview of the topic will be provided although there are specific tools that are particularly useful in the case of MCS problems: specifically the Cook's Distance and the DFBeta.

A discussion of how leave-k-out diagnostics would work in the context of MCS problems is required. There are several scenarios. Suppose we have two methods of measurement X and Y, each with three measurements for a specific case:  $(x_1, x_2, x_3, y_1, y_2, y_3)$

- Leave One Out - one observation is omitted (e.g.  $x_1$ )
- Leave Pair Out - one pair of observation is omitted (e.g.  $x_1$  and  $y_1$ )
- Leave Case (or Subject) Out - All observations associated with a particular case or subject are omitted. (e.g.  $\{x_1, x_2, x_3, y_1, y_2, y_3\}$ )

Other metrics, such as the likelihood distance, will also be introduced, and revisited in a later section.

## 2.3 Case Deletion Diagnostics for Mixed Models

? notes the case deletion diagnostics techniques have not been applied to linear mixed effects models and seeks to develop methodologies in that respect.

? develops these techniques in the context of REML

## 2.4 Terminology for Case Deletion diagnostics

Preisser (1996) describes two type of diagnostics. When the set consists of only one observation, the type is called

## 2.5 Case Deletion Diagnostics

Since the pioneering work of Cook in 1977, deletion measures have been applied to many statistical models for identifying influential observations.

Deletion diagnostics provide a means of assessing the influence of an observation (or groups of observations) on inference on the estimated parameters of LME models.

Data from single individuals, or a small group of subjects may influence non-linear mixed effects model selection. Diagnostics routinely applied in model building may identify such individuals, but these methods are not specifically designed for that purpose and are, therefore, not optimal. We describe two likelihood-based diagnostics for identifying individuals that can influence the choice between two competing models. Case-deletion diagnostics provide a useful tool for identifying influential observations and outliers.

The computation of case deletion diagnostics in the classical model is made simple by the fact that estimates of  $\beta$  and  $\sigma^2$ , which exclude the  $i$ th observation, can be computed without re-fitting the model. Such update formulas are available in the mixed model only if you assume that the covariance parameters are not affected by the removal of the observation in question. This is rarely a reasonable assumption.

## 2.6 Case Deletion Diagnostics

? develops case deletion diagnostics, in particular the equivalent of Cook's distance, for diagnosing influential observations when estimating the fixed effect parameters and variance components.

## 2.7 Terminology for Case Deletion diagnostics

Preisser (1996) describes two type of diagnostics. When the set consists of only one observation, the type is called '*observation-diagnostics*'. For multiple observations, Preisser describes the diagnostics as '*cluster-deletion*' diagnostics. When applied to LME models, such update formulas are available only if one assumes that the covariance parameters are not affected by the removal of the observation in question. However, this is rarely a reasonable assumption.

## 2.8 Case Deletion Diagnostics

Case-deletion diagnostics provide a useful tool for identifying influential observations and outliers.

The computation of case deletion diagnostics in the classical model is made simple by the fact that estimates of  $\beta$  and  $\sigma^2$ , which exclude the  $i$ th observation, can be computed without re-fitting the model. Such update formulas are available in the mixed model only if you assume that the covariance parameters are not affected by the removal of the observation in question. This is rarely a reasonable assumption.

Linear models for uncorrelated data have well established measures to gauge the influence of one or more observations on the analysis. For such models, closed-form update expressions allow efficient computations without refitting the model.

Since the pioneering work of Cook in 1977, deletion measures have been applied to many statistical models for identifying influential observations. Case-deletion diagnos-

tics provide a useful tool for identifying influential observations and outliers.

The key to making deletion diagnostics useable is the development of efficient computational formulas, allowing one to obtain the case deletion diagnostics by making use of basic building blocks, computed only once for the full model.

The computation of case deletion diagnostics in the classical model is made simple by the fact that estimates of  $\beta$  and  $\sigma^2$ , which exclude the  $i$ -th observation, can be computed without re-fitting the model.

## 2.9 Measures of Influence

The impact of an observation on a regression fitting can be determined by the difference between the estimated regression coefficient of a model with all observations and the estimated coefficient when the particular observation is deleted. DFBETA and DFFITS are well known measures of influence. The measure DFBETA is the studentized value of this difference. DFFITS is a statistical measure designed to show how influential an observation is in a statistical model. DFFITS is closely related to the studentized residual.

$$DFBETA_a = \hat{\beta} - \hat{\beta}_{(a)} \quad (2.1)$$

$$= B(Y - Y_{\bar{a}}) \quad (2.2)$$

$$DFFITS = \frac{\hat{y}_i - \widehat{y}_{i(k)}}{s_{(k)}\sqrt{h_{ii}}} \quad (2.3)$$

The prediction residual sum of squares (PRESS) is a value associated with this calculation. When fitting linear models, PRESS can be used as a criterion for model selection, with smaller values indicating better model fits.

$$PRESS = \sum (y - y^{(k)})^2$$



## 2.10 DFBETAs

The measure that measures how much impact each observation has on a particular predictor is DFBETAs. The DFBETA for a predictor and for a particular observation is the difference between the regression coefficient calculated for all of the data and the regression coefficient calculated with the observation deleted, scaled by the standard error calculated with the observation deleted.

DFBETA is a measure found for each observation in a dataset. The DFBETA for a particular observation is the difference between the regression coefficient for an included variable calculated for all of the data and the regression coefficient calculated with the observation deleted, scaled by the standard error calculated with the observation deleted.

The cut-off value for DFBETAs is  $\frac{2}{\sqrt{n}}$ , where  $n$  is the number of observations. However, another cut-off is to look for observations with a value greater than 1.00. Here cutoff means, "this observation could be overly influential on the estimated coefficient."

## 2.11 DFBETAS

DFBETAS (standardized difference of the beta) is a measure that standardizes the absolute difference in parameter estimates between a (mixed effects) regression model based on a full set of data, and a model from which a (potentially influential) subset of data is removed. A value for DFBETAS is calculated for each parameter in the model separately.

This function computes the DFBETAS based on the information returned by the `estex()` function.

The `dfbeta` refers to how much a parameter estimate changes if the observation or case in question is dropped from the data set.

Cook's distance is presumably more important to you if you are doing predictive modeling, whereas `dfbeta` is more important in explanatory modeling.

The DFBETAS statistics are the scaled measures of the change in each parameter estimate and are calculated by deleting the  $i$ th observation:

where  $\hat{\beta}_i$  is the  $i$ th element of  $\hat{\beta}$ . In general, large values of DFBETAS indicate observations that are influential in estimating a given parameter. **Belsley, Kuh, and Welsch (1980)** recommend 2 as a general cutoff value to indicate influential observations and  $2/\sqrt{n}$  as a size-adjusted cutoff.

$$DFBETA_{A_a} = \hat{\beta} - \hat{\beta}_{(a)} \quad (2.4)$$

$$= B(Y - Y_{\hat{a}}) \quad (2.5)$$

In the case of method comparison studies, there are two covariates, and one can construct scatterplots of the pairs of dfbeta values accordingly, both for LOO and LSO calculations. Furthermore 95% confidence ellipse can be constructed around these scatterplots. Note that with  $k$  covariates, there will be  $k + 1$  dfbetas (the intercept,  $\beta_0$ , and 1  $\beta$  for each covariate). For example there would be 2 sets of dfbeta, 510 values for each in the case of LOO, and 85 for LSO diagnostics.

### 2.11.1 DFFITS

DFFITS is a statistical measure designed to show how influential an observation is in a statistical model.

$$DFFITS = \frac{\hat{y}_i - \widehat{y_{i(k)}}}{s_{(k)}\sqrt{h_{ii}}}$$

It is closely related to the studentized residual. For the sake of brevity, we will concentrate on the Studentized Residuals.

DFFITS is a diagnostic meant to show how influential a point is in a statistical regression. It was proposed in 1980. It is defined as the change ("DFFIT"), in the predicted value for a point, obtained when that point is left out of the regression, "Studentized" by dividing by the estimated standard deviation of the fit at that point:

$$DFFITS = \frac{\hat{y}_i - \widehat{y_{i(i)}}}{s_{(i)}\sqrt{h_{ii}}}$$

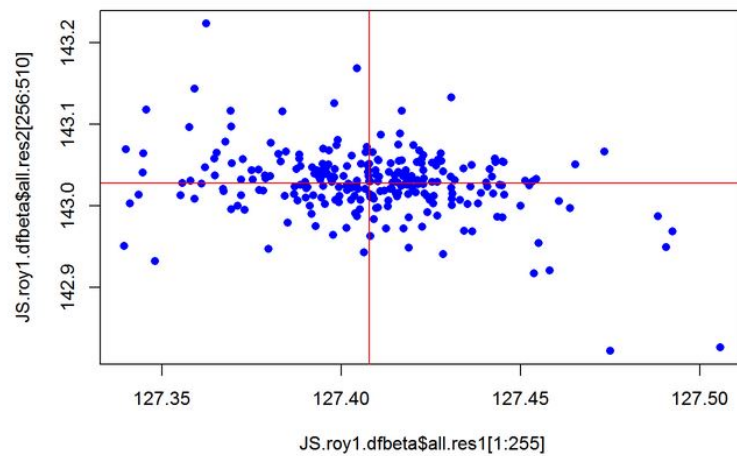


Figure 2.11.1:

### DFbetas for Blood Data

```
plot(JS.roy1.dfbeta$all.res1[1:255], JS.roy1.dfbeta$all.res2[256:510],
     pch=16, col="blue")
abline(v=JS.roy1.dfbeta$all.res1[256], col="red")
abline(h=JS.roy1.dfbeta$all.res2[1], col="red")
```

## 2.12 Likelihood Distance

The likelihood distance is a global, summary measure, expressing the joint influence of the observations in the set  $U$  on all parameters in  $\phi$  that were subject to updating.

The likelihood distance gives the amount by which the log-likelihood of the full data changes if one were to evaluate it at the reduced-data estimates. The important point is that  $l(\psi_U)$  is not the log-likelihood obtained by fitting the model to the reduced data set.

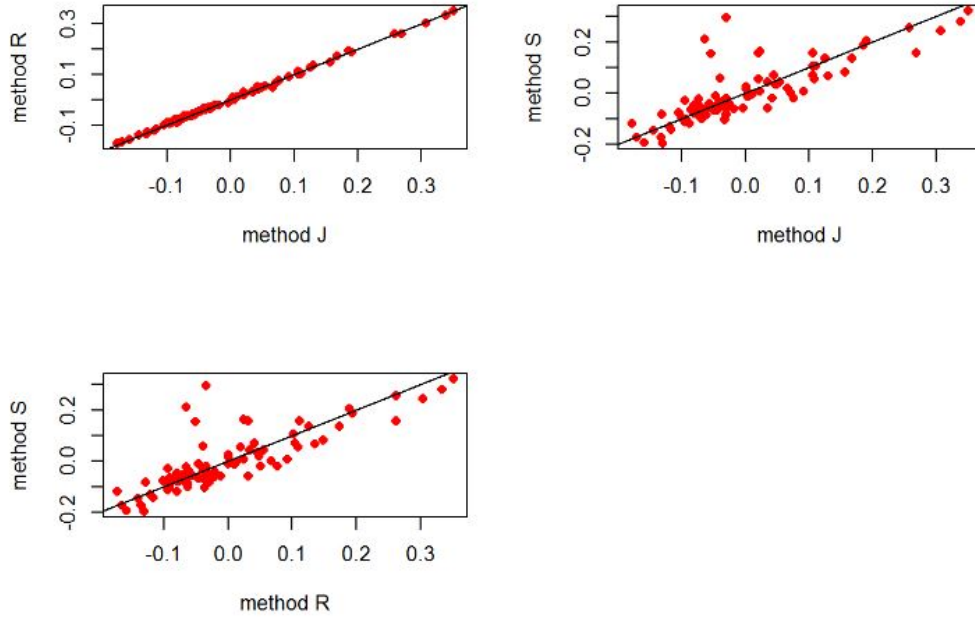
It is obtained by evaluating the likelihood function based on the full data set (containing all  $n$  observations) at the reduced-data estimates.

#### 4. Using DFBETAs to Assess Agreement

Suppose an LME model was formulated to model agreement for various (i.e. 2 or more) methods of measurement, with replicate measurements. If the methods are to be agreement, the DFBetas for each case would be the same for both methods. **As such, agreement between any two methods can be determined by a simple scatterplot of the DFBetas. If the points align along the line of equality, then both methods can be said to be in agreement.**

For the model fitted to the blood data with the lme4 R package, the results tabulated below can be produced. All 85 subjects are ranked by Cook's Distance (with only the top 6 being presented here). The remaining columns are the DFBeta for each of the fixed effects, for each of the 85 subject.

Subject	Cook's D	methodJ	methodR	methodS
78	0.61557407	-0.02934556	-0.03387780	0.2954937
80	0.41590973	-0.06305026	-0.06515241	0.2123881
68	0.22536651	-0.05334867	-0.05062375	0.1555187
72	0.09348500	0.02388626	0.02419887	0.1617474
48	0.08706988	0.02147541	0.03145273	0.1581591
30	0.07118415	0.26925807	0.26215970	0.1581569



In the first of the three plots (*Top Right*), strong agreement between method J and method R is indicated. The other plots indicate lack of agreement of methods J and R with method S.

If lack of agreement is indicated, a subsequent analysis using a technique proposed by Roy(2009) can be used to identify the specific cause for this lack of agreement (see next section).

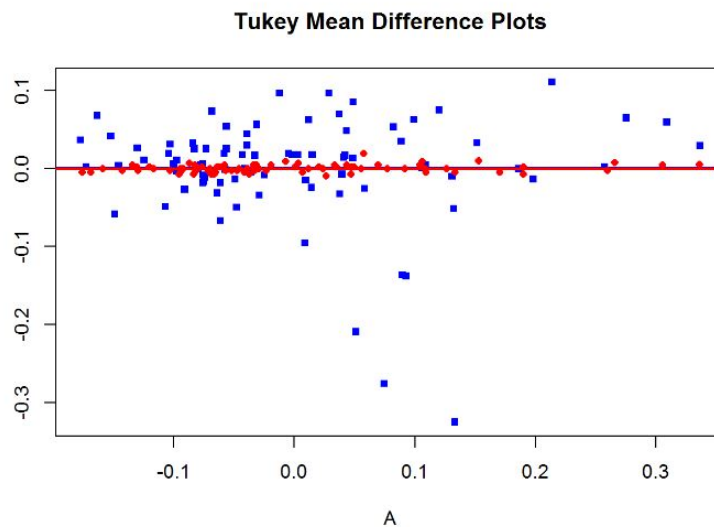
The Pearson Correlation coefficient of the DFBetas can be used in conjunction with this analysis. A high correlation confirms good agreement. No threshold value for agreement is suggested, and analysts are advised to perform model diagnostics regardless of the correlation coefficient.

The Bonferroni Outlier Test and Cook's Distance values can be used to identify unusual cases, when the relationship between sets of dfbeta is modelled as a (classical) linear model. In this model, the covariates should be homoskedastic. A test for non-constant variance may be used to verify this. These diagnostic procedures are implementable using the *car* R package.

Deming Regression can be used to verify the line of equality. Significance test for Deming regression estimates are not available, but 95% bootstrap confidence intervals for the slope estimate and intercept estimates can be computed.

Additionally a mean difference plot can be used to identify outliers. This mean-difference plot differs from the Bland-Altman plot in that the plot is denominated in terms of dfbeta values, and not in measurement units.

If lack of agreement is indicated between methods of measurement, use of Roy's Testing is advised (This is the subject of the next section).



## 5. Using Roy's Test to Identify cause of Lack of agreement

Barnhart specifies three conditions for method of measurement that are required for two methods of measurement to be considered in agreement.

- (i) No Significant Inter-method bias
- (ii) No significant Difference in Within-Subject Variance
- (iii) No significant Difference in Within-Subject Variance

Roy(2009) demonstrates a LME model specification, and a series of tests that look at each of these agreement criteria individually. If two methods of measurement lack agreement, the specific reason or reasons for this lack of agreement can be identified.

Roy proposes an LME model with Kronecker product covariance structure in a doubly multivariate setup. Response for  $i$ th subject can be written as

$$y_i = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + b_{1i} z_{i1} + b_{2i} z_{i2} + \epsilon_i$$

- $\beta_1$  and  $\beta_2$  are fixed effects corresponding to both methods. ( $\beta_0$  is the intercept.)
- $b_{1i}$  and  $b_{2i}$  are random effects corresponding to both methods.

Overall variability between the two methods ( $\Omega$ ) is sum of between-subject ( $D$ ) and within-subject variability ( $\Sigma$ ),

$$\text{Block } \Omega_i = \begin{bmatrix} d_1^2 & d_{12} \\ d_{12} & d_2^2 \end{bmatrix} + \begin{bmatrix} \sigma_1^2 & \sigma_{12} \\ \sigma_{12} & \sigma_2^2 \end{bmatrix}.$$

## 6. Using Roy's Model to Compute LoAs and CR

In this short section, a demonstration of how Roy's technique can be used to compute two common MCS metrics: Limits of Agreement and the Coefficient of Repeatability. While Limits of Agreement are not used in the analysis proposed here, they are ubiquitous in literature, and a demonstration on how to compute them with the Roy Model would assist the adoption of this proposed method.



The coefficient of repeatability is encountered in Gage R & R analysis. (*A future exploration of how LME models can be used in that field would be of interest. This is something to include in the Conclusions Section*).

## 7. Model Diagnostics for Roy's Models

Further to previous work, this section revisits case-deletion and residual diagnostics, and explores how approaches devised by Galecki & Burzykowski (2013) can be used to appraise Roy's model. These authors specifically look at Cook's Distances and Likelihood Distances. For the Roy Model, Cook's Distances may also be generated using the *predictmeans*

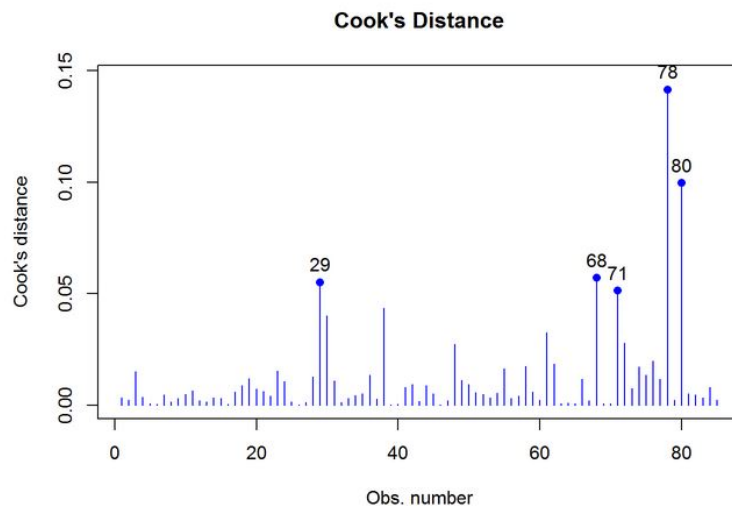


Figure 2.12.2:

As the model is structurally different from the models discussed in the earlier sections, Residual analysis will be briefly revisited.

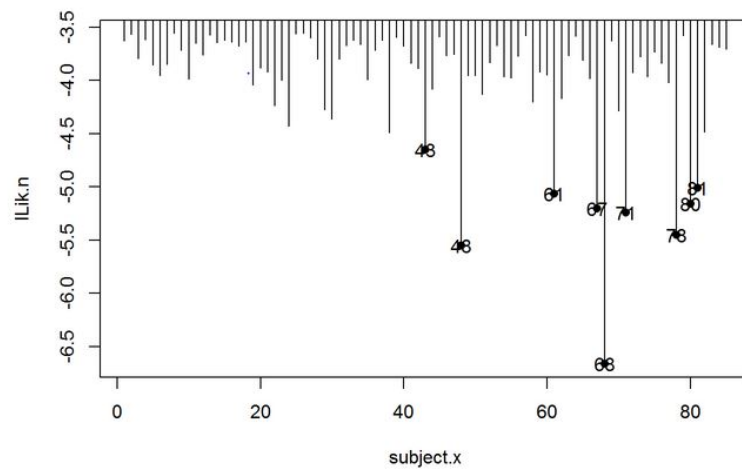


Figure 2.12.3:

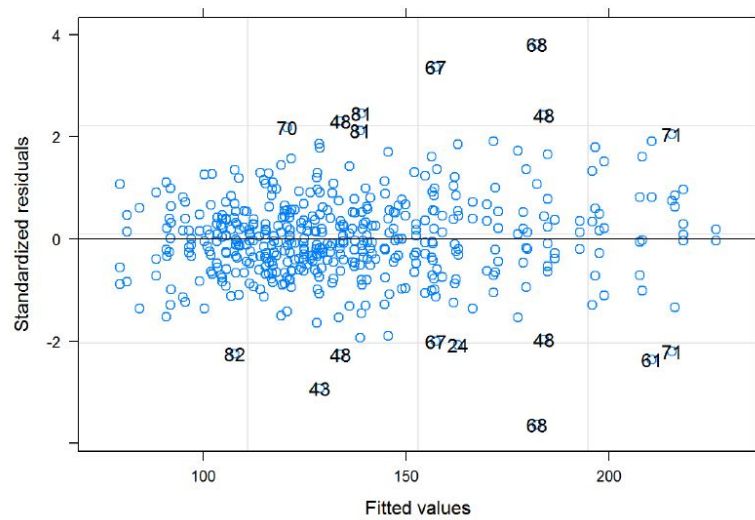


Figure 2.12.4:

## 8. Case Deletion Diagnostics for the Variance Ratios

Schabenberger advises on the use of deletion diagnostics for variance components of an LME model. Taking the core principals of his methods, and applying them to the Method Comparison problem, case deletion diagnostics are used on the variance components of the Roy model., specifically the ratio of between subject variances and the within subject covariances respectively.

$$\text{BSVR} = \frac{\sigma_2^2}{\sigma_2^2} \qquad \text{WSVR} = \frac{d_2^2}{d_2^2}$$

These variance ratios are re-computed for each case removed, and may be analysed separately or jointly for outliers.

The Grubbs' Test for Outliers is a commonly used technique for assessing outlier in a univariate data set. As there may be several outliers (i.e. influential cases) present, the Grubbs test is not practical. However outlier detection using to Tukey's specification for boxplots (i.e. greater than  $Q_3 + 1.5IQR$  or less than  $Q_1 - 1.5IQR$ ), will suffice. Ranking the absolute values of the standardized scores can also be used to identify influential cases, even if the data is not normally distributed.

Bivariate Analyses may be applied jointly to the both sets of data sets, e.g Mahalanobis distances. The Mahalanobis distance, while not an intuitive measure in the context of the data, can be used to rank highly influential cases.

## 9. Permutation Test, Power Tests and Missing Data

This section explores topics such as dependent variable simulation and power analysis, introduced by Galecki & Burzykowski (2013), and implementable with their ***nlmeU*** R package. Using the ***predictmeans*** R package, it is possible to perform permutation t-tests for coefficients of (fixed) effects and permutation F-tests.

The matter of missing data has not been commonly encountered in either Method Comparison Studies or Linear Mixed Effects Modelling. However Roy (2009) deals with the relevant assumptions regarding missing data. Galecki & Burzykowski (2013) approaches the subject of missing data in LME Modelling. The ***nlmeU*** package includes the **patMiss** function, which “*allows to compactly present pattern of missing data in a given vector/matrix/data frame or combination of thereof*”.

### 2.12.1 Influential Observations : DFBeta and DFBetas

## 2.13 Measures of Influence

The impact of an observation, or a case with multiple observations, on a regression fitting can be determined by the difference between the estimated regression coefficient of a model with all observations and the estimated coefficient when the particular observation is deleted. The measure DFBETA is the studentized value of this difference.

## 2.14 Overall Influence

An overall influence statistic measures the change in the objective function being minimized. For example, in OLS regression, the residual sums of squares serves that purpose. In linear mixed models fit by maximum likelihood (ML) or restricted maximum likelihood (REML), an overall influence measure is the likelihood distance [Cook and Weisberg ].

## 2.15 Effects on fitted and predicted values

$$\hat{e}_{i(U)} = y_i - x\hat{\beta}_{(U)} \quad (2.6)$$

## 2.16 Cook's Distance

Cook's Distance is a measure indicating to what extent model parameters are influenced by (a set of) influential data on which the model is based. This function computes the Cook's distance based on the information returned by the `estex()` function.

### 2.16.1 Cook's Distance

- For variance components  $\gamma$ :  $CD(\gamma)_i$ ,

- For fixed effect parameters  $\beta$ :  $CD(\beta)_i$ ,
- For random effect parameters  $\mathbf{u}$ :  $CD(u)_i$ ,
- For linear functions of  $\hat{\beta}$ :  $CD(\psi)_i$

## Random Effects

A large value for  $CD(u)_i$  indicates that the  $i$ -th observation is influential in predicting random effects.

## linear functions

$CD(\psi)_i$  does not have to be calculated unless  $CD(\beta)_i$  is large.

*Cook (1977)* greatly expanded the study of residuals and influence measures. Cook's key observation was the effects of deleting each observation in turn could be computed without undue additional computational expense. Consequently deletion diagnostics have become an integral part of assessing linear models.

Cook (1986) gave a completely general method for assessing influence of local departures from assumptions in statistical models.

### 2.16.2 Cook's Distance

In classical linear regression, a commonly used measure of influence is Cook's distance. It is used as a measure of influence on the regression coefficients.

For linear mixed effects models, Cook's distance can be extended to model influence diagnostics by defining.

$$C_{\beta i} = \frac{(\hat{\beta} - \hat{\beta}_{[i]})^T (\mathbf{X}' \mathbf{V}^{-1} \mathbf{X}) (\hat{\beta} - \hat{\beta}_{[i]})}{p}$$

It is also desirable to measure the influence of the case deletions on the covariance matrix of  $\hat{\beta}$ .

### 2.16.3 Cook's Distance

Cook's  $D$  statistics (i.e. colloquially Cook's Distance) is a measure of the influence of observations in subset  $U$  on a vector of parameter estimates (Cook, 1977).

$$\delta_{(U)} = \hat{\beta} - \hat{\beta}_{(U)}$$

If  $V$  is known, Cook's  $D$  can be calibrated according to a chi-square distribution with degrees of freedom equal to the rank of  $\mathbf{X}$  (?).

For LME models, Cook's distance can be extended to model influence diagnostics by defining.

It is also desirable to measure the influence of the case deletions on the covariance matrix of  $\hat{\beta}$ .

## 2.17 Cook's Distance for LMEs

Diagnostic methods for fixed effects are generally analogues of methods used in classical linear models. Diagnostic methods for variance components are based on 'one-step' methods. Cook (1986) gives a completely general method for assessing the influence of local departures from assumptions in statistical models.

For fixed effects parameter estimates in LME models, the Cook's distance can be extended to measure influence on these fixed effects.

$$CD_i(\beta) = \frac{(c_{ii} - r_{ii}) \times t_i^2}{r_{ii} \times p}$$

For random effect estimates, the Cook's distance is

$$CD_i(b) = g'_{(i)}(I_r + \text{var}(\hat{b})D)^{-2}\text{var}(\hat{b})g_{(i)}.$$

Large values for Cook's distance indicate observations for special attention.

## Cook's Distance

Cooks Distance ( $D_i$ ) is an overall measure of the combined impact of the  $i$ th case of all estimated regression coefficients. It uses the same structure for measuring the

combined impact of the differences in the estimated regression coefficients when the  $i$ -th case is deleted.

Importantly,  $D_{(i)}$  can be calculated without fitting a new regression coefficient each time an observation is deleted.

Cook (1977) greatly expanded the study of residuals and influence measures. Cook's key observation was the effects of deleting each observation in turn could be computed without undue additional computational expense. Consequently deletion diagnostics have become an integral part of assessing linear models.

Cook's Distance is a well known diagnostic technique used in classical linear models, extended to LME models. For LME models, two formulations exist; a Cook's distance that examines the change in fixed fixed parameter estimates, and another that examines the change in random effects parameter estimates. The outcome of either Cook's distance is a scaled change in either  $\beta$  or  $\theta$ .

### 2.17.1 Cook's Distance

In statistics, Cook's Distance or Cook's D is a commonly used estimate of the influence of a data point when performing least squares regression analysis.[1] In a practical ordinary least squares analysis, Cook's distance can be used in several ways: to indicate data points that are particularly worth checking for validity; to indicate regions of the design space where it would be good to be able to obtain more data points. It is named after the American statistician R. Dennis Cook, who introduced the concept in 1977.

### 2.17.2 Change in the precision of estimates

The effect on the precision of estimates is separate from the effect on the point estimates. Data points that have a small Cook's distance, for example, can still greatly affect hypothesis tests and confidence intervals, if their influence on the precision of the estimates is large.



## Interpretation

Specifically  $D_i$  can be interpreted as the distance one's estimates move within the confidence ellipsoid that represents a region of plausible values for the parameters.[clarification needed] This is shown by an alternative but equivalent representation of Cook's distance in terms of changes to the estimates of the regression parameters between the cases where the particular observation is either included or excluded from the regression analysis.

### 2.17.3 Cook's Distance

Some texts tell you that points for which Cook's distance is higher than 1 are to be considered as influential. Other texts give you a threshold of  $4/N$  or  $4/(Nk+1)$ , where  $N$  is the number of observations and  $k$  the number of explanatory variables. In your case the latter formula should yield a threshold around 0.1 .

John Fox (1), in his booklet on regression diagnostics is rather cautious when it comes to giving numerical thresholds. He advises the use of graphics and to examine in closer details the points with "values of  $D$  that are substantially larger than the rest". According to Fox, thresholds should just be used to enhance graphical displays.

In your case the observations 7 and 16 could be considered as influential. Well, I would at least have a closer look at them. The observation 29 is not substantially different from a couple of other observations.

(1) Fox, John. (1991). Regression Diagnostics: An Introduction. Sage Publications.

# Chapter 3

## Appendices 1

### 3.1 Model Terms (Roy 2009)

- Let  $y_{mir}$  be the response of method  $m$  on the  $i$ th subject at the  $r$ —th replicate.
- Let  $\mathbf{y}_{ir}$  be the  $2 \times 1$  vector of measurements corresponding to the  $i$ —th subject at the  $r$ —th replicate.
- Let  $\mathbf{y}_i$  be the  $R_i \times 1$  vector of measurements corresponding to the  $i$ —th subject, where  $R_i$  is number of replicate measurements taken on item  $i$ .
- Let  $\alpha_{mi}$  be the fixed effect parameter for method for subject  $i$ .
- Formally Roy uses a separate fixed effect parameter to describe the true value  $\mu_i$ , but later combines it with the other fixed effects when implementing the model.
- Let  $u_{1i}$  and  $u_{2i}$  be the random effects corresponding to methods for item  $i$ .
- $\boldsymbol{\epsilon}_i$  is a  $n_i$ -dimensional vector comprised of residual components. For the blood pressure data  $n_i = 85$ .
- $\boldsymbol{\beta}$  is the solutions of the means of the two methods. In the LME output, the bias and corresponding t-value and p-values are presented. This is relevant to Roy's first test.

## 3.2 Algorithms : ML v REML

Maximum likelihood estimation is a method of obtaining estimates of unknown parameters by optimizing a likelihood function. The ML parameter estimates are the values of the argument that maximise the likelihood function, i.e. the estimates that make the observed values of the dependent variable most likely, given the distributional assumptions

The most common iterative algorithms used for the optimization problem in the context of LMEs are the EM algorithm, fisher scoring algorithm and NR algorithm, which [cite:West] commends as the preferred method.

A mixed model is an extension of the general linear models that can specify additional random effects terms.

Parameter of the mixed model can be estimated using either ML or REML, while the AIC and the BIC can be used as measures of "goodness of fit" for particular models, where smaller values are considered preferable.

(**Wikipedia**)The restricted (or residual, or reduced) maximum likelihood (REML) approach is a particular form of maximum likelihood estimation which does not base estimates on a maximum likelihood fit of all the information, but instead uses a likelihood function calculated from a transformed set of data, so that nuisance parameters have no effect.

In contrast to the earlier maximum likelihood estimation, REML can produce unbiased estimates of variance and covariance parameters.

### ML procedures for LME

The maximum likelihood procedure of Hartley and Rao yields simultaneous estimates for both the fixed effects and the random effect, by maximising the likelihood of  $\mathbf{y}$  with respect to each element of  $\boldsymbol{\beta}$  and  $\mathbf{b}$ .

### 3.3 Estimation of random effects

Estimation of random effects for LME models in the NLME package is accomplished through use of both EM (Expectation-Maximization) algorithms and Newton-Raphson algorithms.

- EM iterations bring estimates of the parameters into the region of the optimum very quickly, but convergence to the optimum is slow when near the optimum.
- Newton-Raphson iterations are computationally intensive and can be unstable when far from the optimum. However, close to the optimum they converge quickly.
- The LME function implements a hybrid approach, using 25 EM iterations to quickly get near the optimum, then switching to Newton-Raphson iterations to quickly converge to the optimum.
- If convergence problems occur, the “controlargument in LME can be used to change the way the model arrives at the optimum.

### 3.4 Covariance Parameters

The unknown variance elements are referred to as the covariance parameters and collected in the vector  $\theta$ .

#### 3.4.1 Methods and Measures

The key to making deletion diagnostics useable is the development of efficient computational formulas, allowing one to obtain the case deletion diagnostics by making use of basic building blocks, computed only once for the full model.

? lists several established methods of analyzing influence in LME models. These methods include

- Cook's distance for LME models,
- likelihood distance,
- the variance (information) ration,
- the Cook-Weisberg statistic,
- the Andrews-Prebigon statistic.

### 3.5 Computation and Notation

with  $\mathbf{V}$  unknown, a standard practice for estimating  $\mathbf{X}\boldsymbol{\beta}$  is to estimate the variance components  $\sigma_j^2$ , compute an estimate for  $\mathbf{V}$  and then compute the projector matrix  $\mathbf{A}$ ,  $\mathbf{X}\hat{\boldsymbol{\beta}} = \mathbf{A}\mathbf{Y}$ .

? remarks that  $\mathbf{D}$  is a block diagonal with the  $i$ -th block being  $u\mathbf{I}$

## 3.6 Measures 2

### 3.6.1 Cook's Distance

- For variance components  $\gamma$

Diagnostic tool for variance components

$$C_{\theta i} = (\hat{(\theta)}_{[i]} - \hat{(\theta)})^T \text{cov}(\hat{(\theta)})^{-1} (\hat{(\theta)}_{[i]} - \hat{(\theta)})$$

### 3.7 Haslett's Analysis

For fixed effect linear models with correlated error structure Haslett (1999) showed that the effects on the fixed effects estimate of deleting each observation in turn could be cheaply computed from the fixed effects model predicted residuals.



## Chapter 4

## Appendices

## 4.1 Measures 2

### 4.1.1 Cook's Distance

- For variance components  $\gamma$

Diagnostic tool for variance components

$$C_{\theta i} = ((\hat{\theta})_{[i]} - \hat{\theta})^T \text{cov}(\hat{\theta})^{-1} ((\hat{\theta})_{[i]} - \hat{\theta})$$

### 4.1.2 Variance Ratio

- For fixed effect parameters  $\beta$ .

### 4.1.3 Cook-Weisberg statistic

- For fixed effect parameters  $\beta$ .

### 4.1.4 Andrews-Pregibon statistic

- For fixed effect parameters  $\beta$ .

The Andrews-Pregibon statistic  $AP_i$  is a measure of influence based on the volume of the confidence ellipsoid. The larger this statistic is for observation  $i$ , the stronger the influence that observation will have on the model fit.

## 4.2 Computation and Notation

with  $\mathbf{V}$  unknown, a standard practice for estimating  $\mathbf{X}\beta$  is to estimate the variance components  $\sigma_j^2$ , compute an estimate for  $\mathbf{V}$  and then compute the projector matrix  $A$ ,  $\mathbf{X}\hat{\beta} = A\mathbf{Y}$ .

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## 4.3 Measures 2

### 4.3.1 Cook's Distance

- For variance components  $\gamma$

Diagnostic tool for variance components

$$C_{\theta i} = ((\hat{\theta})_{[i]} - \hat{\theta})^T \text{cov}(\hat{\theta})^{-1} ((\hat{\theta})_{[i]} - \hat{\theta})$$

### 4.3.2 Variance Ratio

- For fixed effect parameters  $\beta$ .

### 4.3.3 Cook-Weisberg statistic

- For fixed effect parameters  $\beta$ .

### 4.3.4 Andrews-Pregibon statistic

- For fixed effect parameters  $\beta$ .

The Andrews-Pregibon statistic  $AP_i$  is a measure of influence based on the volume of the confidence ellipsoid. The larger this statistic is for observation  $i$ , the stronger the influence that observation will have on the model fit.

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