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## Chapter 1

## Method Comparison Studies

#### 1.1 What is a method comparison study?

The problem of compare the results of two different measurement techniques, or in other words, assessing the *agreement* between two or more methods of measurement is ubiquitous in scientific research, and is commonly referred to as a 'method comparison study'. Ludbrook (1997) states that the purpose of comparing two measurements "of a continuous biological variable" is to uncover systematic differences, not to point to similarities". The need to compare the results of two different measurement techniques is common in medical statistics.

In particular, in medicine, new methods or devices that are cheaper, easier to use, or less invasive, are routinely developed. Agreement between a new method and either a traditional reference or gold standard must be evaluated before the new one is put into practice. Various methodologies have been proposed for this purpose in recent years.

Published examples of method comparison studies can be found in disciplines as diverse as pharmacology (Ludbrook, 1997), anaesthesia (Myles, 2007), and cardiac imaging methods (Krummenauer et al., 2000).

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.

To illustrate the characteristics of a typical method comparison study consider the data in Table I (Grubbs, 1973). In each of twelve experimental trials, a single round of ammunition was fired from a 155mm gun and its velocity was measured simultaneously (and independently) by three chronographs devices, identified here by the labels 'Fotobalk', 'Counter' and 'Terma'.

Round	Fotobalk [F]	Counter [C]	Terma [T]
1	793.8	794.6	793.2
2	793.1	793.9	793.3
3	792.4	793.2	792.6
4	794.0	794.0	793.8
5	791.4	792.2	791.6
6	792.4	793.1	791.6
7	791.7	792.4	791.6
8	792.3	792.8	792.4
9	789.6	790.2	788.5
10	794.4	795.0	794.7
11	790.9	791.6	791.3
12	793.5	793.8	793.5

Table 1.1.1: Velocity measurement from the three chronographs (Grubbs 1973).

An important aspect of the these data is that all three methods of measurement are assumed to have an attended measurement error, and the velocities reported in Table 1.1 can not be assumed to be 'true values' in any absolute sense.

A method of measurement should ideally be both accurate and precise. Barnhart et al. (2007) describes agreement as being a broader term that contains both of those

qualities. An accurate measurement method will give results close to the unknown 'true value'. The precision of a method is indicated by how tightly measurements obtained under identical conditions are distributed around their mean measurement value. A precise and accurate method will yield results consistently close to the true value. Of course a method may be accurate, but not precise, if the average of its measurements is close to the true value, but those measurements are highly dispersed. Conversely a method that is not accurate may be quite precise, as it consistently indicates the same level of inaccuracy. The tendency of a method of measurement to consistently give results above or below the true value is a source of systematic bias. The smaller the systematic bias, the greater the accuracy of the method.

In the context of the agreement of two methods, there is also a tendency of one measurement method to consistently give results above or below the other method. Lack of agreement is a consequence of the existence of 'inter-method bias'. For two methods to be considered in good agreement, the inter-method bias should be in the region of zero. A simple estimation of the inter-method bias can be calculated using the differences of the paired measurements. The data in Table 1.2 are a good example of possible inter-method bias; the 'Fotobalk' consistently recording smaller velocities than the 'Counter' method. Consequently one would conclude that there is lack of agreement between the two methods.

The absence of inter-method bias by itself is not sufficient to establish whether two measurement methods agree. The two methods must also have equivalent levels of precision. Should one method yield results considerably more variable than those of the other, they can not be considered to be in agreement. With this in mind a methodology is required that allows an analyst to estimate the inter-method bias, and to compare the precision of both methods of measurement.

Round	Fotobalk (F)	Counter (C)	F-C
1	793.8	794.6	-0.8
2	793.1	793.9	-0.8
3	792.4	793.2	-0.8
4	794.0	794.0	0.0
5	791.4	792.2	-0.8
6	792.4	793.1	-0.7
7	791.7	792.4	-0.7
8	792.3	792.8	-0.5
9	789.6	790.2	-0.6
10	794.4	795.0	-0.6
11	790.9	791.6	-0.7
12	793.5	793.8	-0.3

Table 1.1.2: Difference between Fotobalk and Counter measurements.

#### 1.2 Agreement

Bland and Altman (1986) defined perfect agreement as the case where all of the pairs of measurement data, when plotted on a conventional scatter-plot, lie along the line of equality, where the line of equality is defined as the 45 degree line passing through the origin ,(i.e. The line X = Y on the Cartesian plane).

To carry their idea a step further, we define a specific numerical measure of agreement as twice the expected squared perpendicular distance of the pair of random variables  $(X_1, X_2)$  to the line of equality or agreement in the  $(X_1, X_2)$ -plane, that is,  $E(X_1 - X_2)/2$ , where  $X_1$  and  $X_2$  denote the continuous measurements of rater 1 and rater 2, respectively.

Obviously, other  $L_p$  norms may be considered for the purpose of numerically measuring agreement and warrant future consideration.

Agreement is the extent to which the measure of the variable of interest, under a constant set of experimental conditions, yields the same result on repeated trials (Sanchez et al). The more consistent the results, the more reliable the measuring procedure.

#### 1.3 Purpose of Method Comparison Studies

Carstensen (2010) provides a review of many descriptions of the purpose of Method Comparison studies, several of which are reproduced here.

"The question being answered is not always clear, but is usually epxressed as an attempt to quantify the agreement between two methods" (Bland and Altman, 1995).

"Some lack of agreement between different methods of measurement is inevitable. What matters is the amount by which they disagree. We want to know by how much the new method is likely to differ from the old, so that it is not enough to cause problems in the mathematical interpretation we can preplace the old method by the new, or even use the two interchangeably" (Bland and Altman, 1999).

"It often happens that the same physical and chemical property can be measured in different ways. For example, one can determine For example, one can determine sodium in serum by flame atomic emission spectroscopy or by isotope dilution mass spectroscopy. The question arises as to which method is better" (Mandel, 1991).

"In areas of inter-laboratory quality control, method comparisons, assay validations and individual bio-equivalence, etc, the agreement between observations and target (reference) values is of interest" (Lin et al., 2002).

"The purpose of comparing two methods of measurement of a continuous

biological variable is to uncover systematic differences, not to point to similarities'" (Ludbrook, 1997).

"In the pharmaceutical industry, measurement methods that measure the quantity of products are regulated. The FDA (U.S. Food and Drug Administration) requires that the manufacturer show equivalency prior to approving the new or alternatice method in quality control" (Tan & Inglewicz, 1999).

While several major commonalities are present in each definitions, there is a different emphasis for each, which will inevitably give rise to confusion. Carstensen (2010) seems to endorse a simple phrasing of the research question that is proposed by Altman and Bland (1983), i.e. "do the two methods of measurement agree sufficiently closely?" with Carstensen (2010) expressing the view that other considerations (for example, the "equivalence" of two methods) to be treated as separate research questions. As such, we will revert to other research questions, such as "equivalence of methods" later, focusing on agreement and repeatability of methods.

#### 1.4 Bias as a source of Lack Of Agreement

Bland and Altman define bias (referred to hereafter as inter-method bias) as a *a consistent tendency for one method to exceed the other* and propose estimating its value by determining the mean of the case-wise differences. The variation about this mean shall be estimated by the standard deviation of the case-wise differences. Bland and Altman remark that these estimates are based on the assumption that bias and variability are constant throughout the range of measures.

#### 1.5 Statement of a Model

Carstensen (2010) presents a useful formulation for comparing two methods X and Y, in their measurement of item i, where the unknown 'true value' is  $\tau_i$ . Other authors,

such as Kinsella (1986), present similar formulations of the same model, as well as modified models to account for multiple measurements by each methods on each item, known as replicate measurements.

In some types of analysis, such as the conversion problems described by Lewis et al. (1991), an estimate for the scaling factor  $\beta$  may also be sought. For the time being, we will restrict ourselves to problems where  $\beta$  is assumed to be 1.

$$X_i = \tau_i + \delta_i, \qquad \delta_i \sim \mathcal{N}(0, \sigma_\delta^2)$$
 (1.1)

$$Y_i = \alpha + \beta \tau_i + \epsilon_i, \qquad \epsilon_i \sim \mathcal{N}(0, \sigma_{\epsilon}^2)$$
 (1.2)

In this formulation,  $\alpha$  represents the inter-method bias, and can be estimated as E(X - Y). That is to say, a simple estimate of the inter-method bias is given by the differences between pairs of measurements.

Table 1.1 is a good example of possible inter-method bias; the 'Fotobalk' consistently recording smaller velocities than the 'Counter' method. A cursory inspection of the table will indicate a systematic tendency for the Counter method to result in higher measurements than the Fotobalk method.

## Chapter 2

## Improper MCS Techniques

#### 2.1 Methods of assessing agreement

Historically comparison of two methods of measurement was carried out by use of paired sample t—test, correlation coefficients or simple linear regression. Simple linear regression is unsuitable for method comparison studies because of the required assumption that one variable is measured without error. In comparing two methods, both methods are assume to have attendant random error.

#### 2.2 Paired sample t test

This method can be applied to test for statistically significant deviations in bias. This method can be potentially misused for method comparison studies. Paired t tests test only whether the mean responses are the same. Certainly, we want the means to be the same, but this is only a small part of the story. The means can be equal while the (random) differences between measurements can be huge.

Bartko (1994) discusses the use of the well known paired sample t test to test for inter-method bias;  $H: \mu_d = 0$ . The test statistic is distributed a t random variable with n-1 degrees of freedom and is calculated as follows,

$$t^* = \frac{\bar{d}}{\frac{s_d}{\sqrt{n}}} \tag{2.1}$$

where  $\bar{d}$  and  $s_d$  is the average of the differences of the n observations. Only if the two methods show comparable precision then the paired sample student t-test is appropriate for assessing the magnitude of the bias.

#### 2.3 Inappropriate use of the Correlation Coefficient

It is well known that Pearson's correlation coefficient is a measure of the linear association between two variables, not the agreement between two variables (e.g., see Bland and Altman 1986).

This is a well known as a measure of linear association between two variables. Nonetheless this is not necessarily the same as Agreement. This method is considered wholly inadequate to assess agreement because it only evaluates only the association of two sets of observations.

Use of the Pearson Correlation Coefficient, although seemingly intuitive, is not appropriate approach to assessing agreement of two methods. Arguments against its usage have been made repeatedly in the relevant literature. It is possible for two analytical methods to be highly correlated, yet have a poor level of agreement.

It is intuitive when dealing with two sets of related data, i.e the results of the two raters, to calculate the correlation coefficient (r). Bland and Altman attend to this in their 1999 paper.

They present a data set from two sets of meters, and an accompanying scatterplot. An hypothesis test on the data set leads us to conclude that there is a relationship between both sets of meter measurements. The correlation coefficient is determined to be r = 0.94. However, this high correlation does not mean that the two methods agree. It is possible to determine from the scatterplot that the intercept is not zero, a requirement for stating both methods have high agreement. Essentially, should two methods have highly correlated results, it does not follow that they have high agreement.

It is a poor measure of agreement when the rater's measurements are perpendicular to the line of equality[Hutson et al]. In this context, an average difference of zero between the two raters, yet the scatter plot displays strong negative correlation.

The correlation coefficient measures linear agreement—whether the measurements go up-and-down together. Certainly, we want the measures to go up-and-down together, but the correlation coefficient itself is deficient in at least three ways as a measure of agreement.

The correlation coefficient can be close to 1 (or equal to 1!) even when there is considerable bias between the two methods. For example, if one method gives measurements that are always 10 units higher than the other method, the correlation will be 1 exactly, but the measurements will always be 10 units apart.

The magnitude of the correlation coefficient is affected by the range of subjects/units studied.

The correlation coefficient can be made smaller by measuring samples that are similar to each other and larger by measuring samples that are very different from each other. The magnitude of the correlation says nothing about the magnitude of the differences between the paired measurements which, when you get right down to it, is all that really matters.

The usual significance test involving a correlation coefficient—whether the population value is 0—is irrelevant to the comparability problem. What is important is not merely that the correlation coefficient be different from 0. Rather, it should be close to (ideally, equal to) 1!

#### Regression Methods

Regression analysis is typically misused by regressing one measurement on the other and declare them equivalent if and only if the confidence interval for the regression coefficient includes 1. Some simple mathematics shows that if the measurements are comparable, the population value of the regression coefficient will be equal to the correlation coefficient between the two methods.

The population correlation coefficient may be close to 1, but is never 1 in practice. Thus, the only things that can be indicated by the presence of 1 in the confidence interval for the regression coefficient is (1) that the measurements are comparable but there weren't enough observations to distinguish between 1 and the population regression coefficient, or (2) the population regression coefficient is 1 and therefore, the measurements aren't comparable.

There is a line whose slope will be 1 if the measurements are comparable. It is known as a structural equation and is the method advanced by Kelly (1985). Altman and Bland (1987) criticize it for a reason that should come as no surprise: Knowing the data are consistent with a structural equation with a slope of 1 says something about the absence of bias but \*nothing\* about the variability about Y = X (the difference between the measurements), which, as has already been stated, is all that really matters.

#### Intra-class correlation coefficient

The ICC, which takes on values between 0 and 1, is based on analysis of variance techniques. It is close to 1 when the differences between paired measurements is very small compared to the differences between subjects. Of these three procedures—t test, correlation coefficient, intra-class correlation coefficient—the ICC is best because it can be large only if there is no bias and the paired measurements are in good agreement, but it suffers from the same faults ii and iii as ordinary correlation coefficients. The magnitude of the ICC can be manipulated by the choice of samples to split and says nothing about the magnitude of the paired differences.

## Chapter 3

## Repeated Measurements and Repeatability

#### 3.1 Replicate Measurements

Thus far, the formulation for comparison of two measurement methods is one where one measurement by each method is taken on each subject. Should there be two or more measurements by each methods, these measurement are known as 'replicate measurements'. Carstensen et al. (2008) recommends the use of replicate measurements, but acknowledges the additional computational complexity.

Bland and Altman (1986) address this problem by offering two different approaches. The premise of the first approach is that replicate measurements can be treated as independent measurements. The second approach is based upon using the mean of the each group of replicates as a representative value of that group. Using either of these approaches will allow an analyst to estimate the inter method bias.

However, because of the removal of the effects of the replicate measurements error, this would cause the estimation of the standard deviation of the differences to be unduly small. Bland and Altman (1986) propose a correction for this.

Carstensen et al. (2008) takes issue with the limits of agreement based on mean

values of replicate measurements, in that they can only be interpreted as prediction limits for difference between means of repeated measurements by both methods, as opposed to the difference of all measurements. Incorrect conclusions would be caused by such a misinterpretation. Carstensen et al. (2008) demonstrates how the limits of agreement calculated using the mean of replicates are 'much too narrow as prediction limits for differences between future single measurements'. This paper also comments that, while treating the replicate measurements as independent will cause a downward bias on the limits of agreement calculation, this method is preferable to the 'mean of replicates' approach.

#### 3.2 Definition of Replicate measurements

Further to Bland and Altman (1999), a formal definition is required of what exactly replicate measurements are

By replicates we mean two or more measurements on the same individual taken in identical conditions. In general this requirement means that the measurements are taken in quick succession.

Roy accords with Bland and Altman's definition of a replicate, as being two or more measurements on the same individual under identical conditions. Roy allows the assumption that replicated measurements are equi-correlated. Roy allows unequal numbers of replicates.

Replicate measurements are linked over time. However the method can be easily extended to cover situations where they are not linked over time.

#### 3.3 Exchangeable and Linked measurements

Repeated measurements are said to be exchangeable if no relationship exists between successive measurements across measurements. If the condition of exchangeability exists, a group of measurement of the same item determined by the same method can be re-arranged in any permutation without prejudice to proper analysis. There is no reason to believe that the true value of the underlying variable has changed over the course of the measurements.

Exchangeable repeated measurements can be treated as true replicates. For the purposes of method comparison studies the following remarks can be made. The r-th measurement made by method 1 has no special correspondence to the r-th measurement made by method 2, and consequently any pairing of repeated measurements are as good as each other.

Replicate measurements are linked over time. However the method can be easily extended to cover situations where they are not linked over time.

#### 3.4 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 Childrens 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 addition 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 Roy's model and the modified model (denoted 1 and 2 respectively);

$$\hat{\mathbf{\Lambda}}_1 = \begin{pmatrix} 16.61 & 11.67 \\ 11.67 & 27.65 \end{pmatrix} \qquad \hat{\mathbf{\Lambda}}_2 = \begin{pmatrix} 7.55 & 2.60 \\ 2.60 & 18.59 \end{pmatrix}. \tag{3.1}$$

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

Akaike (1974) introduces the Akaike information criterion (AIC), a model selection tool based on the likelihood function. Given a data set, candidate models are ranked according to their AIC values, with the model having the lowest AIC being considered the best fit. Two candidate models can said to be equally good if there is a difference of less than 2 in their AIC values.

The Akaike information criterion (AIC) for both models are  $AIC_1 = 2304.226$  and  $AIC_2 = 2306.226$ , indicating little difference in models. The AIC values for the Carstensen 'unlinked' and 'linked' models are 1994.66 and 1955.48 respectively, indicating an improvement by adding the interaction term.

The  $\hat{\Lambda}$  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 approximately 2).

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

Finally, to complement the blood pressure (i.e. 'J vs S') method comparison from the previous section (i.e. 'J vs S'), the limits of agreement are  $15.62 \pm 1.96 \times 20.33 = (-24.22, 55.46)$ .)

#### 3.5 Repeated Measurements in LME models

In many statistical analyzes, the need to determine parameter estimates where multiple measurements are available on each of a set of variables often arises. Further to Lam et al. (1999), Hamlett et al. (2004) performs an analysis of the correlation of replicate measurements, for two variables of interest, using LME models.

Let  $y_{Aij}$  and  $y_{Bij}$  be the jth repeated observations of the variables of interest A and B taken on the ith subject. The number of repeated measurements for each variable may differ for each individual. Both variables are measured on each time points. Let  $n_i$  be the number of observations for each variable, hence  $2 \times n_i$  observations in total.

It is assumed that the pair  $y_{Aij}$  and  $y_{Bij}$  follow a bivariate normal distribution.

$$\left(egin{array}{c} y_{Aij} \ y_{Bij} \end{array}
ight) \sim \mathcal{N}(oldsymbol{\mu},oldsymbol{\Sigma}) ext{ where } oldsymbol{\mu} = \left(egin{array}{c} \mu_A \ \mu_B \end{array}
ight)$$

The matrix  $\Sigma$  represents the variance component matrix between response variables at a given time point j.

$$oldsymbol{\Sigma} = \left(egin{array}{cc} \sigma_A^2 & \sigma_{AB} \ \sigma_{AB} & \sigma_B^2 \end{array}
ight)$$

 $\sigma_A^2$  is the variance of variable A,  $\sigma_B^2$  is the variance of variable B and  $\sigma_{AB}$  is the covariance of the two variable. It is assumed that  $\Sigma$  does not depend on a particular time point, and is the same over all time points.

#### 3.6 Replicate measurements

Bland and Altman attend to the issue of repeated measures in 1996.

Repeated measurements on several subjects can be used to quantify measurement error, the variation between measurements of the same quantity on the same individual. Bland and Altman discuss two metrics for measurement error; the within-subject standard deviation, and the correlation coefficient.

The above plot incorporates both the conventional limits of agreement ( the inner pair of dashed lines), the 't-' limits of agreement ( the outer pair of dashed lines) centred around the inter-method bias (indicated by the full line). This plot is intended for expository purposes only, as the sample size is small.

Roy (2009b) accords with Bland and Altman?s definition of a replicate, as being two or more measurements on the same individual under identical conditions. Roy allows the assumption that replicated measurements are equi-correlated. Roy allows unequal numbers of replicates.

Replicate measurements are linked over time. However the method can be easily extended to cover situations where they are not linked over time.

Repeated measurements are said to be linked if a direct correspondence exists between successive measurements across measurements, i.e. pairing. Such measurements are commonly made with a time interval between them, but simultaneously for both methods. Paired measurements are exchangeable, but individual measurements are not.

If the paired measurements are taken in a short period of time so that no real systemic changes can take place on each item, they can be considered true replicates. Should enough time elapse for systemic changes, linked repeated measurements can not be treated as true replicates.

In this model, the variances of the random effects must depend on m, since the different methods do not necessarily measure on the same scale, and different methods naturally must be assumed to have different variances. Carstensen (2004) attends to the issue of comparative variances.

Bland and Altman (1999) also remark that an important feature of replicate observations is that they should be independent of each other. This issue is addressed by Carstensen (2010), in terms of exchangeability and linkage. Carstenen advises that repeated measurements come in two *substantially different* forms, depending on the circumstances of their measurement: exchangable and linked.

#### 3.7 Model for replicate measurements

We generalize the single measurement model for the replicate measurement case, by additionally specifying replicate values. Let  $y_{mir}$  be the r-th replicate measurement for subject "i" made by method "m". Further to ? fixed effect can be expressed with a single term  $\alpha_{mi}$ , which incorporate the true value  $\mu_i$ .

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

Combining fixed effects (?), we write,

$$y_{mir} = \alpha_{mi} + e_{mir}$$
.

The following assumptions are required

- $e_{mir}$  is independent of the fixed effects with mean  $E(e_{mir}) = 0$ .
- Further to ? between-item and within-item variances  $Var(\alpha_{mi}) = \sigma_{Bm}^2$  and  $Var(e_{mir}) = \sigma_{Wm}^2$

#### 3.8 Statistical Model For Replicate Measurements

Let  $y_{Aij}$  and  $y_{Bij}$  be the jth repeated observations of the variables of interest A and B taken on the ith subject. The number of repeated measurements for each variable may differ for each individual. Both variables are measured on each time points. Let  $n_i$  be the number of observations for each variable, hence  $2 \times n_i$  observations in total.

It is assumed that the pair  $y_{Aij}$  and  $y_{Bij}$  follow a bivariate normal distribution.

$$\begin{pmatrix} y_{Aij} \\ y_{Bij} \end{pmatrix} \sim \mathcal{N}(\boldsymbol{\mu}, \boldsymbol{\Sigma}) \text{ where } \boldsymbol{\mu} = \begin{pmatrix} \mu_A \\ \mu_B \end{pmatrix}$$
 (3.2)

The matrix  $\Sigma$  represents the variance component matrix between response variables

at a given time point j.

$$\Sigma = \begin{pmatrix} \sigma_A^2 & \sigma_{AB} \\ \sigma_{AB} & \sigma_B^2 \end{pmatrix} \tag{3.3}$$

 $\sigma_A^2$  is the variance of variable A,  $\sigma_B^2$  is the variance of variable B and  $\sigma_{AB}$  is the covariance of the two variable. It is assumed that  $\Sigma$  does not depend on a particular time point, and is the same over all time points.

#### 3.9 What is Repeatability

The quality of repeatability is the ability of a measurement method to give consistent results for a particular subject. That is to say that a measurement will agree with prior and subsequent measurements of the same subject.

Repeatability describes the variation in measurements taken by a single method of measurement on the same item and under the same conditions. A less-than-perfect test-retest reliability causes test-retest variability. Such variability can be caused by, for example, intra-individual variability and intra-observer variability. A measurement may be said to be repeatable when this variation is smaller than some agreed limit.

Test-retest variability is practically used, for example, in medical monitoring of conditions. In these situations, there is often a predetermined "critical difference", and for differences in monitored values that are smaller than this critical difference, the possibility of pre-test variability as a sole cause of the difference may be considered in addition to, for examples, changes in diseases or treatments.

Repeatability is defined by the **IUPAC** as 'the closeness of agreement between independent results obtained with the same method on identical test material, under the same conditions (same operator, same apparatus, same laboratory and after short intervals of time)' and is determined by taking multiple measurements on a series of subjects.

#### 3.10 Repeatability

As mentioned previously, Barnhart et al. (2007) emphasizes the importance of repeatability as part of an overall method comparison study. The coefficient of repeatability was proposed by Bland and Altman (1999), and is referenced in subsequent papers, such as Carstensen et al. (2008). The coefficient of repeatability is a measure of how well a measurement method agrees with itself over replicate measurements (Bland and Altman, 1999).

Repeatability is the ability of a measurement method to give consistent results for a particular subject, i.e. a measurement will agree with prior and subsequent measurements of the same subject. Barnhart et al. (2007) emphasizes the importance of repeatability as part of an overall method comparison study, a view endorsed by Carstensen et al. (2008). Before there can be good agreement between two methods, a method must have good agreement with itself. If one method has poor repeatability in the sense of considerable variability, then agreement between two methods is bound to be poor (Roy, 2009b).

The coefficient of repeatability is a measure of how well a measurement method agrees with itself over replicate measurements (Bland and Altman, 1999). Once the the standard deviations of the differences between the two measurements (in some texts called the residual standard deviation or within-item variability)  $sigma_m$  is determined, the computation of the coefficients of repeatability for both methods is straightforward. The coefficient is calculated from the (in some texts called the residual standard deviation) as  $1.96 \times \sqrt{2} \times \sigma_m = 2.83\sigma_m$ ).

Barnhart et al. (2007) remarks that it is important to report repeatability when assessing measurement, because it measures the purest form of random error not influenced by other factors, while further remarking 'curiously replicate measurements are rarely made in method comparison studies, so that an important aspect of comparability is often overlooked. Bland and Altman (1999) strongly recommends the simultaneous estimation of repeatability and agreement be collecting replicated data. However Roy

(2009b) notes the lack of convenience in such calculations. Repeatability is defined by the IUPAC (2009) as 'the closeness of agreement between independent results obtained with the same method on identical test material, under the same conditions (same operator, same apparatus, same laboratory and after short intervals of time)' and is determined by taking multiple measurements on a series of subjects.

According to the Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results, the following conditions need to be fulfilled in the establishment of repeatability:

- the same measurement procedure
- the same observer
- the same measuring instrument, used under the same conditions
- the same location
- repetition over a short period of time.
- same objectives

A measurement is said to be repeatable when this variation is smaller than some prespecified limit. In these situations, there is often a predetermined "critical difference", and for differences in monitored values that are smaller than this critical difference, the possibility of pre-test variability as a sole cause of the difference may be considered in addition to, for examples, changes in diseases or treatments. A measurement method can be said to have a good level of repeatability if there is consistency in repeated measurements on the same subject using that method. Conversely, a method has poor repeatability if there is considerable variation in repeated measurements.

The British Standards Institute (1979) defines a coefficient of repeatability as the value below which the difference between two single test results may be expected to lie within a specified probability. In the absence of other indications, the probability is 95%.

#### 3.11 Coefficient of Repeatability - Good

The coefficient of repeatability is a measure of how well a measurement method agrees with itself over replicate measurements (Bland and Altman, 1999). Once the withinitem variability is known, the computation of the coefficients of repeatability for both methods is straightforward. Bland and Altman (1999) introduces the repeatability coefficient for a method, which is defined as the upper limits of a prediction interval for the absolute difference between two measurements by the same method on the same item under identical circumstances (Carstensen et al., 2008). The British standards Insitute [1979] define a coefficient of repeatability as the value below which the difference between two single test results....may be expected to lie within a specified probability. Unless otherwise instructed, the probability is assumed to be 95%.

 $\sigma_x^2$  is the within-subject variance of method x. The repeatability coefficient is  $2.77\sigma_x$  (i.e.  $1.96 \times \sqrt{2}\sigma_x$ ). For 95% of subjects, two replicated measurement by the same method will be within this repeatability coefficient.

The Bland Altman method offers a measurement on the repeatability of the methods. The Coefficient of Repeatability (CR) can be calculated as 1.96 (or 2) times the standard deviations of the differences between the two measurements ( $d_2$  and  $d_1$ ).

Bland and Altman (1999) introduces the repeatability coefficient for a method, which is defined as the upper limits of a prediction interval for the absolute difference between two measurements by the same method on the same item under identical circumstances (Carstensen et al., 2008).

 $\sigma_x^2$  is the within-subject variance of method x. The repeatability coefficient is  $2.77\sigma_x$  (i.e.  $1.96 \times \sqrt{2}\sigma_x$ ). For 95% of subjects, two replicated measurement by the same method will be within this repeatability coefficient.

#### 3.12 Gold Standard

This is considered to be the most accurate measurement of a particular parameter.

#### 3.13 Repeatability and Gold Standards

Currently the phrase 'gold standard' describes the most accurate method of measurement available. No other criteria are set out. Further to Dunn (2002), various gold standards have a varying levels of repeatability. Dunn cites the example of the sphygmomanometer, which is prone to measurement error. Consequently it can be said that a measurement method can be the 'gold standard', yet have poor repeatability.

Dunn (2002) recognizes this problem. Hence, if the most accurate method is considered to have poor repeatability, it is referred to as a "bronze standard". Again, no formal definition of a 'bronze standard' exists.

The coefficient of repeatability may provide the basis of formulation a formal definition of a 'gold standard'. For example, by determining the ratio of CR to the sample mean  $\bar{X}$ . Advisably the sample size should specified in advance. A gold standard may be defined as the method with the lowest value of  $\lambda = CR/\bar{X}$  with  $\lambda < 0.1\%$ . Similarly, a silver standard may be defined as the method with the lowest value of  $\lambda$ with  $0.1\% \le \lambda < 1\%$ . Such thresholds are solely for expository purposes.

#### 3.14 Importance of Repeatability in MCS

Repeatability is important in the context of method comparison because the repeatability of two methods influence the amount of agreement which is possible between those methods. If one method has poor repeatability, the agreement is bound to be poor. If both methods have poor repeatability, agreement is even worse. If one method has poor repeatability in the sense of considerable variability, then agreement between two methods is bound to be poor (Roy, 2009b).

Barnhart et al. (2007) and Roy (2009a) highlight the importance of reporting repeatability in method comparison, because it measures the purest random error not influenced by any external factors. Statistical procedures on within-subject variances of two methods are equivalent to tests on their respective repeatability coefficients. A formal test is introduced by Roy (2009a), which will discussed in due course.

Barnhart emphasizes the importance of repeatability as part of an overall method comparison study. Before there can be good agreement between two methods, a method must have good agreement with itself. The coefficient of repeatability, as proposed by Bland & Altman (1999) is an important feature of both Carstensen's and Roy's methodologies. The coefficient is calculated from the residual standard deviation (i.e.  $1.96 \times \sqrt{2} \times \sigma_m = 2.83\sigma_m$ ).

Bland and Altman (1999) strongly recommends the simultaneous estimation of repeatability and agreement be collecting replicated data. Roy (2009b) notes the lack of convenience in such calculations. It is important to report repeatability when assessing measurement, because it measures the purest form of random error not influenced by other factors (Barnhart et al., 2007).

importance of repeatability' curiously replicate measurements are rarely made in method comparison studies, so that an important aspect of comparability is often overlooked.

As noted by Bland and Altman 1999, the repeatability of two methods of measurement can potentially limit Repeatability (using Bland-Altman plot) The Bland-Altman plot may also be used to assess a method?s repeatability by comparing repeated measurements using one single measurement method on a sample of items. The plot can then also be used to check whether the variability or precision of a method is related to the size of the characteristic being measured. Since for the repeated measurements the same method is used, the mean difference should be zero. Therefore the Coefficient of Repeatability (CR) can be calculated as 1.96 (often rounded to 2) times the standard deviation of the case-wise differences.

# 3.15 Repeatability in Bland-Altman Blood Data Analysis

- Two readings by the same method will be within  $1.96\sqrt{2}\sigma_w$  or  $2.77\sigma_w$  for 95% of subjects. This value is called the repeatability coefficient.
- For observer J using the sphygmomanometer  $\sigma_w = \sqrt{37.408} = 6.116$  and so the repeatability coefficient is  $2:77 \times 6.116 = 16:95$  mmHg.
- For the machine  $S_{,\sigma_w} = \sqrt{83.141} = 9.118$  and the repeatability coefficient is  $2:77\times9.118 = 25.27$  mmHg.
- Thus, the repeatability of the machine is 50% greater than that of the observer.

#### 3.16 Carstensen Move to Chapter 2

- The limits of agreement are not always the only issue of interest the assessment of method specific repeatability and reproducibility are of interest in their own right.
- Repeatability can only be assessed when replicate measurements by each method are available.
- The repeatability coefficient for a method is defined as the upper limits of a prediction interval for the absolute difference between two measurements by the same method on the same item under identical circumstances.
- If the standard deviation of a measurement is  $\sigma$  the repeatability coefficient is  $2 \times \sqrt{2}\sigma = 2.83 \times \sigma \approx 2.8\sigma$ .
- The repeatability of measurement methods is calculated differently under the two models

- Under the model assuming exchangeable replicates (1), the repeatability is based only on the residual standard deviation, i.e.  $2.8\sigma_m$
- Under the model for linked replicates (2) there are two possibilities depending on the circumstances.
- If the variation between replicates within item can be considered a part of the repeatability it will be  $2.8\sqrt{\omega^2 + \sigma_m^2}$ .
- However, if replicates are taken under substantially different circumstances, the variance component  $\omega^2$  may be considered irrelevant in the repeatability and one would therefore base the repeatability on the measurement errors alone, i.e. use  $2.8\sigma_m$ .

#### 3.16.1 Notes from BXC Book (chapter 9)

The assessment of method-specific repeatability and reproducibility is of interest in its own right. Repeatability and reproducibility can only be assessed when replicate measurements by each method are available. If replicate measurements by a method are available, it is simple to estimate the measurement error for a method, using a model with fixed effects for item, then taking the residual standard deviation as measurement error standard deviation. However, if replicates are linked, this may produce an estimate that biased upwards.

The repeatability coefficient (or simply repeatability) for a method is defined as the upper limit of a prediction interval for the absolute difference between two measurements by the same method on the same item under identical circumstances (see above conditions)

$$y_{mir} = \alpha_m + \beta_m(\mu_i + a_{ir} + c_{mi}) + e_{mir}$$

The variation between measurements under identical circumstances.

## 3.17 Sampling Scheme: Linked and Unlinked Replicates

Measurements taken in quick succession by the same observer using the same instrument on the same subject can be considered true replicates. Roy (2009b) notes that some measurements may not be 'true' replicates.

Roy's methodology assumes the use of 'true replicates'. However data may not be collected in this way. In such cases, the correlation matrix on the replicates may require a different structure, such as the autoregressive order one AR(1) structure. However determining MLEs with such a structure would be computational intense, if possible at all.

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)

## **Bibliography**

- Akaike, H. (1974). A new look at the statistical model identification. *IEEE Transactions* on Automatic Control 19(6), 716–723.
- Altman, D. and J. Bland (1983). Measurement in medicine: The analysis of method comparison studies. *Journal of the Royal Statistical Society. Series D (The Statistician)* 32(3), 307–317.
- Barnhart, H., M. Haber, and L. Lin (2007). An overview of assessing agreement with continuous measurements. *Journal of Biopharmaceutical Statistics* 17, 529–569.
- Bartko, J. (1994). Measures of agreement: A single procedure. Statistics in Medicine 13, 737–745.
- Bland, J. and D. Altman (1986). Statistical methods for assessing agreement between two methods of clinical measurement. *The Lancet i*, 307–310.
- Bland, J. and D. Altman (1995). Comparing methods of measurement why plotting difference against standard method is misleading. *The Lancet 346*, 1085–87.
- Bland, J. and D. Altman (1999). Measuring agreement in method comparison studies. Statistical Methods in Medical Research 8(2), 135–160.
- Carstensen, B. (2004). Comparing and predicting between several methods of measurement. *Biostatistics* 5(3), 399–413.
- Carstensen, B. (2010). Comparing Clinical Measurement Methods: A Practical Guide. Wiley.

- Carstensen, B., J. Simpson, and L. C. Gurrin (2008). Statistical models for assessing agreement in method comparison studies with replicate measurements. *The International Journal of Biostatistics* 4(1).
- Dunn, G. (2002). Statistical Evaluation of Measurement Error (Second ed.). Stanford: American Mathematical Society and Digital Press.
- Grubbs, F. (1973). Errors of measurement, precision, accuracy and the statistical comparison of measuring instruments. *Technometrics* 15(1), 53–66.
- Hamlett, A., L. Ryan, and R. Wolfinger (2004). On the use of PROC MIXED to estimate correlation in the presence of repeated measures. *Proceedings of the Statistics* and Data Analysis Section, SAS Users Group International 198-229, 1–7.
- IUPAC (2009). IUPAC Compendium of Chemical Terminology the Gold Book. http://goldbook.iupac.org/R05293.html.
- Kinsella, A. (1986). Estimating method precision. The Statistician 35, 421–427.
- Krummenauer, F., I. Genevriere, and U. Nixdorff (2000). The biometrical comparison of cardiac imaging methods. *Computer Methods and Programs in Biomedicine* 62, 21–34.
- Lam, M., K. Webb, and D. O'Donnell (1999). Correlation between two variables in repeated measurements. American Statistical Association, Proceedings of the Biometric Session, 213–218.
- Lewis, P., P. Jones, J. Polak, and H. Tillitson (1991). The problem of conversion in method comparison studies. *Applied Statistics* 40, 105–112.
- Lin, L., A. Hedayat, B. Sinha, and M. Yang (2002). Statistical methods in assessing agreement: Models issues and tools. *Journal of American Statistical Association 97*, 257–270.

- Ludbrook, J. (1997). Comparing methods of measurement. Clinical and Experimental Pharmacology and Physiology 24, 193–203.
- Myles, P. (2007). Using the Bland Altman method to measure agreement with repeated measures. *British Journal of Anaesthesia* 99(3), 309–311.
- Roy, A. (2009a). An application of linear mixed effects model to assess the agreement between two methods with replicated observations. *Journal of Biopharmaceutical Statistics* 19, 150–173.
- Roy, A. (2009b). An application of the linear mixed effects model to ass the agreement between two methods with replicated observations. *Journal of Biopharmaceutical Statistics* 19, 150–173.