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Statistical Models for Assessing Agreement in Method Comparison Studies with Replicate Measurements

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Statistical Models for Assessing Agreement in Method Comparison Studies with Replicate Measurements*

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Abstract

Method comparison studies are usually analyzed by computing limits of agreement. It is recommended that replicate measurements be taken by each method, but the resulting data are more cumbersome to analyze. We discuss the statistical model underlying the classical limits of agreement and extend it to the case with replicate measurements. As the required code to fit the models is non-trivial, we provide example computer code to fit the models, and show how to use the output to derive measures of repeatability and limits of agreement.

KEYWORDS: method comparison, Bland-Altman plot, mixed models

*We are grateful to Peter Dalgaard for (much needed) advice on the lme syntax.

1 Introduction

The problem of comparing two methods of measurement is still occasionally approached by computing correlation coefficients, despite the fact that this has been discouraged as irrelevant and misleading for more than 20 years [1, 2]. The preferred approach is to consider the differences between measurements by the two methods, and produce prediction limits for the difference between pairs of future measurements, known as the *limits of agreement*.

When replicate measurements are taken with each method on each item (i.e. person or sample) measuring agreement becomes slightly more complicated. Bland and Altman [3] presented details of various approaches to adopt in this case, mainly based on calculations that can be performed “by hand”. Such tedious computations are unnecessary since the underlying concept of limits of agreement is merely a prediction from a statistical model that can be fitted with modern software for random effects models. The estimates of the variance components are given directly in the program output and can be used directly to generate limits of agreement and measures of repeatability of the methods.

This has the advantage of bypassing a lot of hand-calculations and makes it irrelevant whether the design is perfectly balanced or not.

Moreover, setting up a model focuses on the implications of the exchangeability properties of the replicate measurements, e.g. whether replicates are exchangeable *within* each method by item stratum or only within items (paired or linked replicates).

2 Notation

In this paper we set up models for method comparison data with replicate measurements. The models that are needed are models where the residual variances differ by method, and this type of model is not very clearly presented in the manuals of any of the major software packages, so therefore we provide the code needed in R, Stata and SAS.

We assume the data are formatted as a dataset with four columns named:

`meth`, method of measurement, the number of methods being M ,

`item`, items (persons, samples) measured by each method, of which there are I ,

`repl`, replicate indicating repeated measurement of the same item by the same method, and

`y`, the measurement.

We denote the measurement by method m on item i , replicate r by y_{mir} .

When specifying mixed models we use Greek letters for fixed effects and Latin letters for random effects.

3 The classical approach

The classical setup for comparison of two measurement methods is one where one measurement by each method is taken on each item, that is, without replicates. In that case the recommendation is to compute the *limits of agreement*, a prediction interval for the difference between future measurements with the two methods on a new individual.

Underlying this approach is the two-way analysis of variance model:

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

The differences $y_{1i} - y_{2i}$ have variance $\sigma_1^2 + \sigma_2^2$, and the prediction interval for a difference between two new measurements is therefore:

$$\alpha_1 - \alpha_2 \pm 1.96 \times \sqrt{\sigma_1^2 + \sigma_2^2}$$

In practice, the term $\alpha_1 - \alpha_2$ is estimated by the mean difference, the last term is computed as the empirical standard deviation of the differences, and the 1.96 is replaced by 2 for convenience:

$$\bar{d} \pm 2 \text{ s.d.}(d_i)$$

— this is what is commonly termed the limits of agreement.

This is formally incorrect as a prediction interval, since the errors in estimation of the parameters are not taken into account; formally the 95% prediction interval for the difference should be computed as:

$$\bar{d} \pm t_{0.975}(I-1) \sqrt{1 + 1/I} \times \text{s.d.}(d_i)$$

where I is the number of items. The term $t_{0.975}(I-1) \sqrt{1 + 1/I}$ is 2.05 for $I = 30$ and less than 2 if $I > 61$, so the pragmatic method gives slight underestimates of the width of the limits of agreement for small studies. This is however based on a heavy exploitation of the normality assumption of the error terms (e_{mi}).

There are two rather more interesting assumptions in the model:

1. The variation of the differences is constant over the range of measurements.
2. The difference between the methods is constant over the range of measurements.

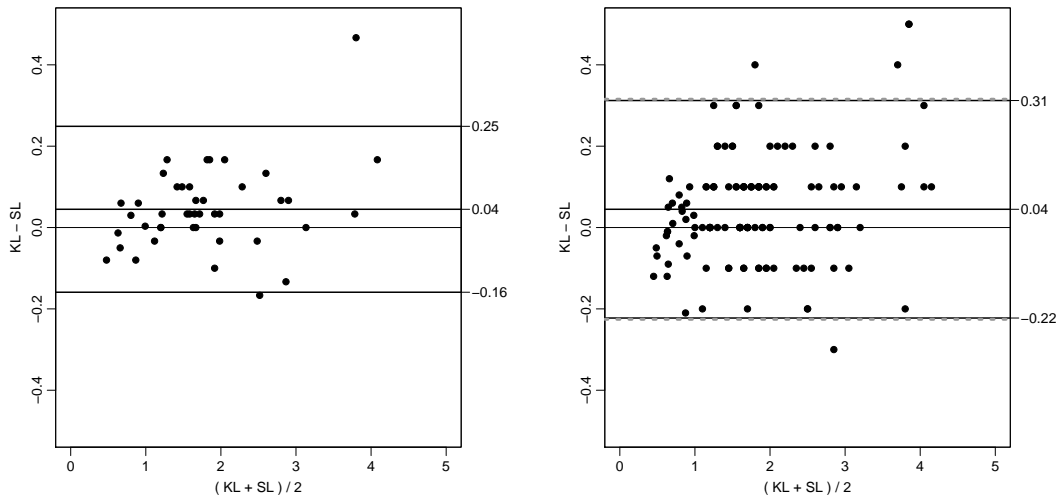


Figure 1: *Measurements of subcutaneous fat (in mm) by two different observers. Data from the Steno Diabetes Center, 2006. The left panel is a Bland-Altman plot based on the means over replicates with limits of agreement based on these. The right panel is a Bland-Altman plot where the replicates are randomly matched, and $(item \times repl)$ are used as independent items ignoring the exchangeability. The thick broken (gray) lines almost on top of the limits of agreement represent the correct limits of agreement computed from the variance component model in section 4.*

These assumptions are checked by making a so-called Bland-Altman plot [2], where differences are plotted against averages of methods.

Figure 1 presents data from a comparison of measurements of subcutaneous fat by two observers at the Steno Diabetes Center. Measurements are in millimeters (mm). Each person is measured three times by each observer. The sequence of measurements is not considered to be of importance, so the replicate measurements are exchangeable within person (item) and observer (method).

The graph indicates that the underlying assumptions are reasonably well fulfilled. The limits of agreement in the first graph are based on the means of repeats within item and method. These limits of agreement can only be interpreted as prediction limits for the difference between means of three measurements by both methods, which is normally not relevant. Hence we must set up a framework that allows us to address the relevant prediction question based on *single* measurements.

4 Models for replicate measurements

To determine prediction limits for differences between single measurements we must resort to a more elaborate model for our data, where replicate measurements are explicitly modeled:

$$y_{mir} = \alpha_m + \mu_i + c_{mi} + e_{mir}, \quad c_{mi} \sim \mathcal{N}(0, \tau_m^2), \quad e_{mir} \sim \mathcal{N}(0, \sigma_m^2) \quad (1)$$

This is a model where the variation between items for method m is captured by τ_m and the within item variation by σ_m . The formulation of this model is general and refers to comparison of any number of methods — however, if only two methods are compared, separate values of τ_1^2 and τ_2^2 cannot be estimated, only their average, so in the case of only two methods we are forced to assume that $\tau_1 = \tau_2 = \tau$.

Under this model the limits of agreement should be computed based on the standard deviation of the difference between a pair of measurements by the two methods on a new individual, j , say:

$$\text{var}(y_{1j} - y_{2j}) = 2\tau^2 + \sigma_1^2 + \sigma_2^2$$

Therefore the limits of agreement are estimated by:

$$\hat{\alpha}_1 - \hat{\alpha}_2 \pm 2 \times \sqrt{2\hat{\tau}^2 + \hat{\sigma}_1^2 + \hat{\sigma}_2^2}$$

It therefore only remains to estimate the variance components in this linear mixed model, which can be done using standard software. Using the subcutaneous fat example, we present below the code and output for the statistical packages R, Stata and SAS.

4.1 Practical estimation of the variance components

4.1.1 Data

For generality the dataset was set up with the variable names `meth`, `item`, `repl` and `y`. All three examples below are using this data set-up:

	meth	item	repl	y
1	KL	1	1	1.6
2	KL	1	2	1.7
3	KL	1	3	1.7
4	KL	3	1	2.8
5	KL	3	2	2.9
6	KL	3	3	2.8
7	KL	5	1	2.7
8	KL	5	2	2.8

```

9    KL    5    3    2.9
...
10   SL    1    1    1.7
11   SL    1    2    1.6
12   SL    1    3    1.7
13   SL    3    1    2.8
14   SL    3    2    2.7
15   SL    3    3    2.8
16   SL    5    1    3.0
17   SL    5    2    2.9
18   SL    5    3    2.9
...

```

4.1.2 R

The function to use in R is `lme`, but the syntax is somewhat arcane, see e.g. [6]. If the `random` argument in `lme` is a list, and the name of the first element is the name of a variable in the dataset, all terms are nested in this variable. The example here requires that the variables `meth`, `item` and `repl` are factors.

```

> lme( y ~ meth + item,
+      random = list( item = pdIdent( ~ meth-1 ) ),
+      weights = varIdent( form = ~1 | meth ),
+      data=fat
+      )
Linear mixed-effects model fit by REML
Data: fat
Log-restricted-likelihood: 188.3488
Fixed: y ~ meth + item
      (Intercept)      methSL      item2      item3
1.6896001995 -0.0448837209 -0.8653286307  1.1326030428
...

Random effects:
Formula: ~meth - 1 | item
Structure: Multiple of an Identity
           methKL  methSL  Residual
StdDev:  0.059556  0.059556  0.07717392

Variance function:
Structure: Different standard deviations per stratum
Formula: ~1 | meth
Parameter estimates:
           KL           SL
1.0000000  0.9383578
Number of Observations: 258
Number of Groups: 43

```

R gives the interaction s.d. and *one* of the residual s.d.s in the section named `Random effects:`, whereas the ratio of the residual standard deviations is found

under the section Variance function. In this case the interaction s.d. is 0.059556, the residual s.d. for method KL is 0.077174 and for method SL it is $0.077174 \times 0.938358 = 0.072417$. The estimated difference in means between method 1 and 2 is 0.044837, so the limits of agreement are then given by:

$$0.044883 \pm 2 \times \sqrt{2 \times 0.059556^2 + 0.077174^2 + 0.072417^2} = (-0.23, 0.32)$$

4.1.3 Stata

The function to use in Stata is `xtmixed`, which is only available as of Stata version 9, [5, 7]. To calculate separate residual variances for each of the methods, `xtmixed` requires generation of new variables that has a unique code for each (method×item) and each (method×item×replicate) combination. Additionally, `xtmixed` parametrizes the residual variances, as the variance for the method with the smallest residual variance and the difference in residual variances between the two methods. Therefore we must take care to use the method with the smallest residual variance as the reference. Doing it the wrong way around produces some warning messages and estimates without standard errors.

Using the `var` option produces estimates of the variance parameters and not the s.d.s. The `nocons` option is required to exclude the usual residual variation term which is no longer required (output truncated to the right):

```
gen meth1 = ( meth == 1 )
gen MI = item + 100 * meth1
gen MIR = _n

xi: xtmixed y i.meth1 i.item || MI: || MIR:meth1, nocons var
...
-----
          y |          Coef.   Std. Err.      z    P>|z|    [95% Co
-----+-----
   _Imeth1_1 |   .0448837     .015868      2.83   0.005   .013782
   _Iitem_2  |  -.8653287     .0735594    -11.76  0.000  -1.00950
   _Iitem_3  |   1.132603     .0735594     15.40  0.000   .988429
...
-----
Random-effects Parameters | Estimate   Std. Err.   [95% Co
-----+-----
MI: Identity      var(_cons) | .0035469   .0011984   .001829
MIR: Identity    var(meth1) | .0007116   .0012102   .000025
-----+-----
                  var(Residual) | .0052442   .0007997   .003889
-----
```


The residual variance for method 2 is 0.0052442 and for method 1 $0.0052442 + 0.0007116 = 0.0059558$, and the method by item interaction variance is 0.0035469. The estimated difference in means between method 1 and method 2 is 0.0448837, so the limits of agreement for the difference between method 1 and method 2 are:

$$0.0448837 \pm 2 \times \sqrt{2 \times 0.0035469 + 0.0052442 + 0.0059558} = (-0.23, 0.32)$$

4.1.4 SAS

The procedure to use is `proc mixed`[4], and with the generic names of the variables we use the following code to fit the model (output truncated to the right):

```
proc mixed data = rdata ;
  class meth item ;
  model y = meth item / s;
  random meth * item ;
  repeated item / group = meth ;
run ;
```

...

Covariance Parameter Estimates

Cov Parm	Group	Estimate
meth*item		0.003547
item	meth 1	0.005956
item	meth 2	0.005244

...

Solution for Fixed Effects

Effect	meth item	Estimate	Error	Standard DF	t Value
Intercept		1.6277	0.05259	42	30.95
meth	1	0.04488	0.01587	42	2.83
meth	2	0	.	.	.
item	1	0.01703	0.07356	42	0.23
item	2	-0.8483	0.07356	42	-11.53
item	3	1.1496	0.07356	42	15.63

...

SAS gives the desired variance components directly as in the model formulation and also the difference between means, so the limits of agreement are:

$$0.04488 \pm 2 \times \sqrt{2 \times 0.003547 + 0.005956 + 0.005244} = (-0.23, 0.32)$$

Note that SAS requires considerably less fidgeting with variables than do Stata, it has a syntax that is more in line with the way models are usually specified than that of R, and it gives estimates of the parameters used in the specification of the model. No wonder that `proc mixed` has become a *de facto* standard for fitting variance components models!

4.2 Limits of agreement

The limits of agreement based on the mixed model are shown in the right hand panel of figure 1. These correct limits are virtually indistinguishable from those based on a random pairing of replicates within item and using these item by replicate pairings as observations. We shall return to this point below.

5 Linked replicates

In the example above, we have assumed that the replicates were exchangeable *within* each method by item stratum. Sometimes, however, replicates are taken in parallel by each of the methods, which means that the values are linked by a common environment; typically time or sampling occasion.

5.1 The oximetry example

An example of this is the oximetry study, done at the Royal Children's Hospital in Melbourne to examine the agreement between pulse oximetry and co-oximetry in small babies. Many were very sick and therefore had very low oxygen saturation levels — the normal range is between 95 and 100%. Each baby was measured three times by each method; performed at three different times for each infant.

There were 61 babies in the study, of these, four had only measurements on two occasions, and one on only one occasion.

Since replicates are linked across methods we need to incorporate this in the model by including an extra random effect common within each item by replicate stratum:

$$\begin{aligned} y_{mir} &= \alpha_m + \mu_i + a_{ir} + c_{mi} + e_{mir}, \\ a_{ir} &\sim \mathcal{N}(0, \omega^2), \quad c_{mi} \sim \mathcal{N}(0, \tau_m^2), \quad e_{mir} \sim \mathcal{N}(0, \sigma_m^2) \end{aligned} \quad (2)$$

Recall that with only two methods we cannot estimate two separate, method-specific values of τ .

Note that the variance of the extra random effect (a_{ir}) cannot depend on method, but in principle it could depend on item-specific features, or some of it might be taken as a fixed effect, the latter could for example include an effect of time if replicates were taken at specific times.

When subtracting measurements by the two methods the effects a_{ir} cancel, so under this extended model we have the same expression for the variance of the differences as before:

$$\text{var}(y_{1j} - y_{2j}) = 2\tau^2 + \sigma_1^2 + \sigma_2^2,$$

so the limits of agreement are again:

$$\hat{\alpha}_1 - \hat{\alpha}_2 \pm 2 \times \sqrt{2\hat{\tau}^2 + \hat{\sigma}_1^2 + \hat{\sigma}_2^2}$$

Model (2) differs from the previous model (1) in the *estimation* of the variance components. The model where the replicates are non-exchangeable within method has some of the variation allocated to the `item`×`replicate` method.

It should be noted that the model with random effects of both `method`×`item` and `item`×`replicate` is a so-called “crossed” model and therefore usually will take longer time to fit.

5.2 Fitting the model

In the following we briefly indicate the code to fit the model with the crossed effects of `meth`×`item` and `item`×`repl`. The full code and the output generated is shown in the appendix.

5.2.1 R

The convention in the `lme` syntax is that when the `random` option is a list and the first element has the name of a variable from the dataset all the effects are nested in this. In the example below, both `meth` and `repl` are nested in `item`, i.e. we have `meth`×`item` and `item`×`repl` as random effects.

The R-code for fitting the model is:

```
lme( y ~ meth + item,
     random=list( item = pdIdent( ~ meth-1 ),
                 repl = ~1 ),
     weights = varIdent( form = ~1 | meth ),
     data=ox )
```

5.2.2 Stata

When using Stata we need to generate a few interaction variables prior to calling `xtmixed`:

```
. gen meth1 = (meth==1)
. gen meth2 = (meth==2)
. gen MI = item + 100*meth
. gen IR = item + 100*repl
. gen MIR = _n
. xi:xtmixed y i.meth i.item || _all:R.MI || _all:R.IR ///
              || MIR:meth2, nocons var
```

5.2.3 SAS

SAS has the absolutely simplest syntax — we just need to add the desired interaction:

```
proc mixed data = rdata ;  
  class meth item repl;  
  model y = meth item / s;  
  random meth * item item * repl ;  
  repeated item / group = meth ;  
run ;
```

5.3 Results

For the oximetry data we have the following results for the variance components, when fitting the correct model as well as the model where we (wrongly) assume exchangeable replicates:

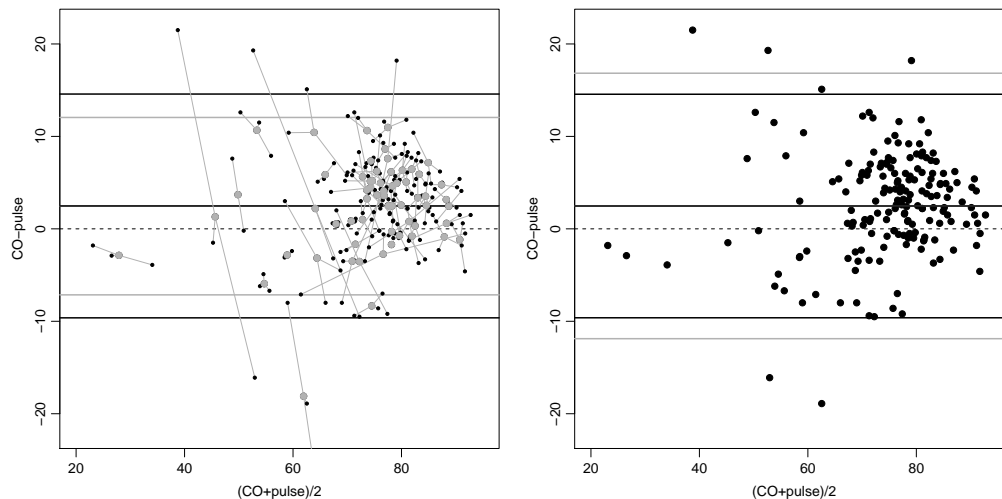


Figure 2: *The oximetry data. Left panel: Bland-Altman plot for means over replicates (gray), and paired replicates (black). The individual replicates are connected with a gray line to the mean. Right panel: Bland-Altman plot for the individual replicates. Gray limits of agreement are based on estimates from a model assuming exchangeability of replicates within methods, black limits on the correct model for the linked replicates.*

Model (random eff.)	$m \times i$	$i \times r$	Residual		Total		Limits of agreement
	τ	ω	σ_1	σ_2	Σ_1	Σ_2	
$m \times i, i \times r$	2.93	3.42	2.22	3.99	5.02	6.02	2.47 (-9.87;14.81)
$m \times i$	2.19		4.07	5.24	4.62	5.68	2.47 (-12.18;17.12)

We see that failure to account for the $i \times r$ interaction only slightly underestimates the total s.d.s, $\Sigma_1 = \sqrt{\tau^2 + \omega^2 + \sigma_1^2}$ and $\Sigma_2 = \sqrt{\tau^2 + \omega^2 + \sigma_2^2}$, but a substantial part of it is allocated to the wrong variance component, and so produces too wide limits of agreement.

Failure to take the replication structure into account results in over-estimation of the prediction interval for the difference between future measurements. This is illustrated in figure 2, where the left panel shows the limits obtained using classical methods, and the right panel shows the limits derived from mixed effects models. The difference between limits obtained by using the linked replicates as items, and fitting the correct model is very small in this case, whereas the effect of using means strongly underestimates the limits and failing to take account of the replication structure in the models strongly overestimates the limits.

6 Repeatability

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 σ 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 ω^2 may be considered irrelevant in the repeatability and one would therefore base the repeatability on the measurement errors alone, i.e. use $2.8\sigma_m$. In such cases one would presumably try to model the effects of differing replication circumstances by a systematic ef-

fect. Hence there is no subject-matter-free way of defining repeatability from the variance components in the models.

In the oximetry example the measurements were taken rather close in time and hence it would be natural to include the between replicate variation in the calculation of repeatability. For co-oximetry the repeatability is $2.8 \times \sqrt{3.42^2 + 2.22^2} = 2.8 \times 4.08 = 11.4\%$ and for pulse oximetry it is $2.8 \times \sqrt{3.42^2 + 3.99^2} = 2.8 \times 5.25 = 14.7\%$. Hence the upper 95% limits for the absolute difference between two repeat measurements by the two methods is 11.4 and 14.7% respectively, whereas the limits of agreement (CO–pulse) are (−9.9; 14.8)%. Thus the discrepancy between the two methods is largely attributable to the rather poor repeatability of both methods.

This conclusion would clearly not have been possible without taking replicate measurements by the two methods.

Had we deemed the between replicate variation to be irrelevant, the repeatabilities would have been only $2.8 \times 2.22 = 6.2\%$ for CO and $2.8 \times 3.99 = 11.2\%$ for pulse; substantially smaller, but still major contributors to the width of the limits of agreement.

7 Getting it wrong and getting it almost right

In a dataset with replicate measurements there are two ways to treat the data along the lines indicated by Bland & Altman [2] which covers the situation with only one measurement per method and item:

1. Take means over replicates within each method by item stratum.
2. Replicates within item are taken as items.

Suppose that we have the following model (model 2) for the measurements:

$$\begin{aligned} y_{mir} &= \alpha_m + \mu_i + a_{ir} + c_{mi} + e_{mir}, \\ a_{ir} &\sim \mathcal{N}(0, \omega^2), \quad c_{mi} \sim \mathcal{N}(0, \tau_m^2), \quad e_{mir} \sim \mathcal{N}(0, \sigma_m^2) \end{aligned} \quad (3)$$

Note that we are allowing the interaction between method and item to have separate variances for each method — with only two methods these cannot be estimated separately, but they can of course still be used in calculations. The random $i \times r$ interaction term is only relevant if the replicates are linked across methods (paired replicates).

In the model the correct limits of agreement would be:

$$\alpha_1 - \alpha_2 \pm 2\sqrt{\tau_1^2 + \tau_2^2 + \sigma_1^2 + \sigma_2^2}$$

7.1 Averaging over replicates

If we are using means of replicates to form the differences we have (R_{mi} is the number of replicates by method m on item i):

$$\bar{d}_i = \bar{y}_{1i} - \bar{y}_{2i} = \alpha_1 - \alpha_2 + \frac{\sum_r a_{ir}}{R_{1i}} - \frac{\sum_r a_{ir}}{R_{2i}} + c_{1i} - c_{2i} + \frac{\sum_r e_{1ir}}{R_{1i}} - \frac{\sum_r e_{2ir}}{R_{2i}}$$

The terms with a_{ir} are only relevant for linked replicates in which case $R_{1i} = R_{2i}$ and therefore the term vanishes. Thus:

$$\text{var}(\bar{d}_i) = \tau_1^2 + \tau_2^2 + \sigma_1^2/R_{1i} + \sigma_2^2/R_{2i} < \tau_1^2 + \tau_2^2 + \sigma_1^2 + \sigma_2^2$$

so the limits of agreement calculated based on the means are much too narrow as prediction limits for differences between future *single* measurements.

7.2 Replicates as items

If replicates are taken as items, then the calculated differences are:

$$d_{ir} = y_{1ir} - y_{2ir} = \alpha_1 - \alpha_2 + c_{1i} - c_{2i} + e_{1ir} - e_{2ir}$$

which has variance $\tau_1^2 + \tau_2^2 + \sigma_1^2 + \sigma_2^2$, and therefore using the empirical variance of the differences in principle gives the correct limits of agreement. However the differences are not independent:

$$\text{cov}(d_{ir}, d_{is}) = \tau_1^2 + \tau_2^2, \quad \text{cor}(d_{ir}, d_{is}) = \frac{\tau_1^2 + \tau_2^2}{\tau_1^2 + \tau_2^2 + \sigma_1^2 + \sigma_2^2}$$

This is negligible if the residual variances are very large compared to the interaction, so the estimate of the “correct” variance based on these differences is likely to be only slightly downwards biased.

If replicates are exchangeable within method by item strata it is not clear how to produce the differences — it can be done in a number of different ways since the replicates can be matched within item in several different ways. If replicates are paired at random, the variance will still be correct, assuming model (2) (without the $i \times r$ interaction term)

$$\text{var}(y_{1ir} - y_{2is}) = \tau_1^2 + \sigma_1^2 + \tau_2^2 + \sigma_2^2$$

but again the differences will be positively correlated within item:

$$\text{cov}(y_{1ir} - y_{2is}, y_{1it} - y_{2iu}) = \tau_1^2 + \tau_2^2$$

so the estimate of $\tau_1^2 + \sigma_1^2 + \tau_2^2 + \sigma_2^2$ as the empirical variance of $y_{1ir} - y_{2is}$ for a random matching of replicates between methods will be an underestimate, albeit not a large one. In the fat dataset (with exchangeable replicates) the correct upper limit of agreement based on the model is 0.315, the upper limit based on the numbering in the dataset is 0.312, but the median upper limit over 1000 random matchings of replicates within items is 0.309.

8 Conclusion

Based on this, we offer the following general advice in the analysis of method comparison studies with replicate measurements:

- Do not use hand calculations — they are overly complicated and outdated in the computer age — software for mixed models was constructed for a reason.
- Set up the correct model, taking the exchangeability structure of the data into account: If replicates are linked across methods, include the item by replicate random effect, otherwise not.
- Fit the model and use the estimated parameters (and your subject-matter knowledge) to draw conclusions based on:
 - the limits of agreement between methods
 - repeatability of methods
- If you absolutely refuse to use modern statistical software, use (item \times replicate) as items; if replicates are not linked, then make a random pairing. However, the correlations will bias the limits of agreement downward, and you will miss important information on the repeatability by not knowing the variance components. Your analysis will still be suboptimal, but not a totally wrong as it would be if you used averages over replicates.

Appendix: Programs

In this section we show the total results from fitting the models to the two datasets by the three packages.

R

The R-programs are completely self-contained since the two datasets used for illustration are part of the MethComp package. Currently (June 2008) the package is only available at www.biostat.ku.dk/~bxc/MethComp.

Exchangeable replicates

```
> library( MethComp )
Loading required package: R2WinBUGS
> library( nlme )
>
> data( fat )
> fat <- data.frame( item=factor(fat$Id),
+                   meth=fat$Obs,
+                   repl=factor(fat$Rep),
+                   y=fat$Sub )
> str( fat )
'data.frame': 258 obs. of 4 variables:
 $ item: Factor w/ 43 levels "1","2","3","4",...: 1 1 1 3 3 3 5 5 5 11 ...
 $ meth: Factor w/ 2 levels "KL","SL": 1 1 1 1 1 1 1 1 1 1 ...
 $ repl: Factor w/ 3 levels "1","2","3": 1 2 3 1 2 3 1 2 3 1 ...
 $ y : num 1.6 1.7 1.7 2.8 2.9 2.8 2.7 2.8 2.9 3.9 ...
>
> # The convention is that within a list in random, the terms subsequent to
> # item are nested within item
>
> lme( y ~ meth + item,
+      random = list( item = pdIdent( ~ meth-1 ) ),
+      weights = varIdent( form = ~1 | meth ),
+      data=fat
+      )
Linear mixed-effects model fit by REML
Data: fat
Log-restricted-likelihood: 188.3488
Fixed: y ~ meth + item
(Intercept)      methSL      item2      item3      item4
1.6896001995 -0.0448837209 -0.8653286307 1.1326030428 -1.0077856154
      item5      item6      item7      item8      item9
1.2014605811 -0.7673239282 -0.1844287691 -0.2510954358 0.6155712309
      item10     item11     item13     item14     item15
-0.5496348547 2.1282212996 -0.6750365145 1.2326030428 -0.9973239282
      item16     item17     item18     item19     item20
-0.3851590597 -0.0007302905 -0.0844287691 -0.0836984786 0.1815076070
      item21     item22     item24     item25     item27
-0.4347939144 0.2510954358 0.3170318119 0.0496348547 -0.4503651453
      item28     item29     item30     item31     item32
-1.0365206086 0.9318727523 0.3163015214 0.0992697095 -1.1891236514
      item33     item34     item35     item36     item37
-0.0333333333 2.1163015214 0.8170318119 0.3815076070 1.4666666667
      item38     item39     item40     item41     item42
-0.4666666667 -0.7991236514 0.8518257263 2.4148409403 -0.4666666667
      item43     item44     item45     item46
-0.1170318119 0.2496348547 0.1481742737 -0.0170318119

Random effects:
Formula: ~meth - 1 | item
Structure: Multiple of an Identity
      methKL  methSL  Residual
StdDev: 0.059556 0.059556 0.07717392

Variance function:
Structure: Different standard deviations per stratum
Formula: ~1 | meth
Parameter estimates:
      KL      SL
1.0000000 0.9383578
Number of Observations: 258
Number of Groups: 43
```

From the output (red entries) we get the following quantities:

$$\begin{aligned}\alpha_{SL} - \alpha_{KL} &= -0.0448837209 \\ \tau &= 0.059559 \\ \sigma_{KL} &= 0.07717392 \\ \sigma_{SL}/\sigma_{KL} &= 0.9383578\end{aligned}$$

Linked replicates

```
> library( MethComp )
Loading required package: R2WinBUGS
> library( nlme )
>
> data( ox )
> ox$item <- factor(ox$item)
> ox$repl <- factor(ox$repl)
> str( ox )
'data.frame': 354 obs. of 4 variables:
 $ meth: Factor w/ 2 levels "CO","pulse": 1 1 1 1 1 1 1 1 1 1 ...
 $ item: Factor w/ 61 levels "1","2","3","4",...: 1 1 1 2 2 2 3 3 4 ...
 $ repl: Factor w/ 3 levels "1","2","3": 1 2 3 1 2 3 1 2 3 1 ...
 $ y : num 78 76.4 77.2 68.7 67.6 68.3 82.9 80.1 80.7 62.3 ...
>
> # The convention is that within a list in random, the termes subsequent to
> # item are nested within item
>
> lme( y ~ meth + item,
+      random=list( item = pdIdent( ~ meth-1 ),
+                  repl = ~1 ),
+      weights = varIdent( form = ~1 | meth ),
+      data=ox
+      )
Linear mixed-effects model fit by REML
Data: ox
Log-restricted-likelihood: -911.7401
Fixed: y ~ meth + item
(Intercept) methpulse item2 item3 item4 item5
76.0428384 -2.4704462 -7.0216227 5.1497034 -10.7281860 -1.1137199
 item6 item7 item8 item9 item10 item11
3.1649924 9.7065633 3.5568599 -4.1821374 -14.4222445 12.7503731
 item12 item13 item14 item15 item16 item17
-47.3135668 3.3219575 -1.1293724 6.2565251 -0.5367298 13.9153464
 item18 item19 item20 item21 item22 item23
1.5322522 -2.0861271 -1.0351969 6.4653272 -0.4416475 4.5820322
 item24 item25 item26 item27 item28 item29
8.2772197 2.1049894 2.7779659 -10.3186089 -10.8197187 0.7833716
 item30 item31 item32 item33 item34 item35
2.6444795 -29.2466418 5.6528703 6.8769614 8.7365767 0.9285974
 item36 item37 item38 item39 item40 item41
3.0492155 3.6735649 7.5298316 2.7392939 -8.6159587 -0.1044011
 item42 item43 item44 item45 item46 item47
-4.3450727 -20.7468236 -16.2943647 2.0329985 4.5130501 3.4254305
 item48 item49 item50 item51 item52 item53
-3.0309414 10.4662553 -24.8350417 -20.8508611 -0.3525354 -3.6222924
 item54 item55 item56 item57 item58 item59
1.4299082 12.8385572 9.7971680 13.3501148 13.4953406 15.6657386
 item60 item61
7.3963452 -1.7503731

Random effects:
Formula: ~meth - 1 | item
Structure: Multiple of an Identity
methCO methpulse
StdDev: 2.928042 2.928042

Formula: ~1 | repl %in% item
(Intercept) Residual
StdDev: 3.415692 2.224868
```


y	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
_Imeth1_1	.0448837	.015868	2.83	0.005	.0137829	.0759845
_Iitem_2	-.8653287	.0735594	-11.76	0.000	-1.009502	-.7211549
_Iitem_3	1.132603	.0735594	15.40	0.000	.9884293	1.276777
_Iitem_4	-1.007786	.0735594	-13.70	0.000	-1.151959	-.8636119
_Iitem_5	1.201461	.0735594	16.33	0.000	1.057287	1.345634
_Iitem_6	-.7673239	.0735594	-10.43	0.000	-.9114977	-.6231502
_Iitem_7	-.1844288	.0735594	-2.51	0.012	-.3286025	-.040255
_Iitem_8	-.2510955	.0735594	-3.41	0.001	-.3952692	-.1069217
_Iitem_9	.6155712	.0735594	8.37	0.000	.4713975	.759745
_Iitem_10	-.5496349	.0735594	-7.47	0.000	-.6938086	-.4054611
_Iitem_11	2.128221	.0735594	28.93	0.000	1.984048	2.272395
_Iitem_13	-.6750365	.0735594	-9.18	0.000	-.8192103	-.5308628
_Iitem_14	1.232603	.0735594	16.76	0.000	1.088429	1.376777
_Iitem_15	-.9973239	.0735594	-13.56	0.000	-1.141498	-.8531502
_Iitem_16	-.3851591	.0735594	-5.24	0.000	-.5293328	-.2409853
_Iitem_17	-.0007303	.0735594	-0.01	0.992	-.144904	.1434435
_Iitem_18	-.0844288	.0735594	-1.15	0.251	-.2286025	.059745
_Iitem_19	-.0836985	.0735594	-1.14	0.255	-.2278723	.0604753
_Iitem_20	.1815076	.0735594	2.47	0.014	.0373338	.3256814
_Iitem_21	-.4347939	.0735594	-5.91	0.000	-.5789677	-.2906201
_Iitem_22	.2510953	.0735594	3.41	0.001	.1069216	.3952691
_Iitem_24	.3170317	.0735594	4.31	0.000	.172858	.4612055
_Iitem_25	.0496349	.0735594	0.67	0.500	-.0945389	.1938086
_Iitem_27	-.4503651	.0735594	-6.12	0.000	-.5945389	-.3061914
_Iitem_28	-1.036521	.0735594	-14.09	0.000	-1.180694	-.8923469
_Iitem_29	.9318727	.0735594	12.67	0.000	.787699	1.076047
_Iitem_30	.3163015	.0735594	4.30	0.000	.1721277	.4604752
_Iitem_31	.0992697	.0735594	1.35	0.177	-.0449041	.2434435
_Iitem_32	-1.189124	.0735594	-16.17	0.000	-1.333297	-1.04495
_Iitem_33	-.0333334	.0735594	-0.45	0.650	-.1775071	.1108404
_Iitem_34	2.116302	.0735594	28.77	0.000	1.972128	2.260475
_Iitem_35	.8170318	.0735594	11.11	0.000	.672858	.9612055
_Iitem_36	.3815076	.0735594	5.19	0.000	.2373338	.5256814
_Iitem_37	1.466667	.0735594	19.94	0.000	1.322493	1.61084
_Iitem_38	-.4666667	.0735594	-6.34	0.000	-.6108404	-.3224929
_Iitem_39	-.7991237	.0735594	-10.86	0.000	-.9432975	-.6549499
_Iitem_40	.8518256	.0735594	11.58	0.000	.7076519	.9959994
_Iitem_41	2.414841	.0735594	32.83	0.000	2.270667	2.559015
_Iitem_42	-.4666667	.0735594	-6.34	0.000	-.6108404	-.3224929
_Iitem_43	-.1170318	.0735594	-1.59	0.112	-.2612056	.0271419
_Iitem_44	.2496348	.0735594	3.39	0.001	.105461	.3938086
_Iitem_45	.1481743	.0735594	2.01	0.044	.0040005	.292348
_Iitem_46	-.0170318	.0735594	-0.23	0.817	-.1612056	.1271419
_cons	1.644717	.05259	31.27	0.000	1.541642	1.747791

Random-effects Parameters	Estimate	Std. Err.	[95% Conf. Interval]	
MI: Identity				
var(_cons)	.0035469	.0011984	.0018291	.0068779
MIR: Identity				
var(meth1)	.0007116	.0012102	.0000254	.0199439
var(Residual)	.0052442	.0007997	.0038893	.0070711

LR test vs. linear regression: chi2(2) = 23.45 Prob > chi2 = 0.0000

Note: LR test is conservative and provided only for reference

_Iitem_12		-47.31357	4.422289	-10.70	0.000	-55.98109	-38.64604
_Iitem_13		3.321958	4.422289	0.75	0.453	-5.345569	11.98948
_Iitem_14		-1.129371	4.422289	-0.26	0.798	-9.796898	7.538155
_Iitem_15		6.256526	4.422289	1.41	0.157	-2.411	14.92405
_Iitem_16		-.5367311	4.422289	-0.12	0.903	-9.204257	8.130795
_Iitem_17		13.91535	4.715682	2.95	0.003	4.672781	23.15791
_Iitem_18		1.532253	4.422289	0.35	0.729	-7.135274	10.19978
_Iitem_19		-2.086126	4.422289	-0.47	0.637	-10.75365	6.5814
_Iitem_20		-1.035196	4.715682	-0.22	0.826	-10.27776	8.20737
_Iitem_21		6.465328	4.422289	1.46	0.144	-2.202198	15.13285
_Iitem_22		-.4416481	4.422289	-0.10	0.920	-9.109174	8.225878
_Iitem_23		4.582033	4.422289	1.04	0.300	-4.085493	13.24956
_Iitem_24		8.27722	4.422289	1.87	0.061	-.3903066	16.94475
_Iitem_25		2.104989	4.715682	0.45	0.655	-7.137578	11.34756
_Iitem_26		2.777965	4.422289	0.63	0.530	-5.889561	11.44549
_Iitem_27		-10.31861	4.422289	-2.33	0.020	-18.98613	-1.651082
_Iitem_28		-10.81972	4.422289	-2.45	0.014	-19.48724	-2.152191
_Iitem_29		.7833705	4.422289	0.18	0.859	-7.884156	9.450897
_Iitem_30		2.64448	4.422289	0.60	0.550	-6.023047	11.31201
_Iitem_31		-29.24664	4.422289	-6.61	0.000	-37.91417	-20.57911
_Iitem_32		5.652869	4.422289	1.28	0.201	-3.014657	14.3204
_Iitem_33		6.876962	4.422289	1.56	0.120	-1.790565	15.54449
_Iitem_34		8.736578	4.422289	1.98	0.048	.0690514	17.4041
_Iitem_35		.928597	4.422289	0.21	0.834	-7.738929	9.596123
_Iitem_36		3.049215	4.422289	0.69	0.491	-5.618312	11.71674
_Iitem_37		3.673565	4.422289	0.83	0.406	-4.993961	12.34109
_Iitem_38		7.529832	4.422289	1.70	0.089	-1.137694	16.19736
_Iitem_39		2.739297	5.492224	0.50	0.618	-8.025264	13.50386
_Iitem_40		-8.615959	4.422289	-1.95	0.051	-17.28349	.0515677
_Iitem_41		-.1044024	4.422289	-0.02	0.981	-8.771929	8.563124
_Iitem_42		-4.345072	4.422289	-0.98	0.326	-13.0126	4.322455
_Iitem_43		-20.74682	4.422289	-4.69	0.000	-29.41435	-12.0793
_Iitem_44		-16.29436	4.422289	-3.68	0.000	-24.96189	-7.626837
_Iitem_45		2.032999	4.422289	0.46	0.646	-6.634527	10.70053
_Iitem_46		4.513051	4.422289	1.02	0.307	-4.154475	13.18058
_Iitem_47		3.425431	4.422289	0.77	0.439	-5.242095	12.09296
_Iitem_48		-3.03094	4.422289	-0.69	0.493	-11.69847	5.636586
_Iitem_49		10.46626	4.422289	2.37	0.018	1.798729	19.13378
_Iitem_50		-24.83504	4.715682	-5.27	0.000	-34.07761	-15.59248
_Iitem_51		-20.85086	4.422289	-4.71	0.000	-29.51839	-12.18333
_Iitem_52		-.3525351	4.422289	-0.08	0.936	-9.020062	8.314991
_Iitem_53		-3.622292	4.422289	-0.82	0.413	-12.28982	5.045235
_Iitem_54		1.42991	4.422289	0.32	0.746	-7.237617	10.09744
_Iitem_55		12.83856	4.422289	2.90	0.004	4.17103	21.50608
_Iitem_56		9.797168	4.422289	2.22	0.027	1.129642	18.46469
_Iitem_57		13.35012	4.422289	3.02	0.003	4.68259	22.01764
_Iitem_58		13.49534	4.422289	3.05	0.002	4.827816	22.16287
_Iitem_59		15.66574	4.422289	3.54	0.000	6.998213	24.33327
_Iitem_60		7.396344	4.422289	1.67	0.094	-1.271182	16.06387
_Iitem_61		-1.750373	4.422289	-0.40	0.692	-10.4179	6.917154
_cons		76.04284	3.138534	24.23	0.000	69.89142	82.19425

Random-effects Parameters	Estimate	Std. Err.	[95% Conf. Interval]	
_all: Identity				
var(R.MI)	8.573426	2.25398	5.121191	14.35284
_all: Identity				
var(R.IR)	11.66695	2.263471	7.976607	17.06462
MIR: Identity				
var(meth2)	11.00559	3.624397	5.771552	20.9862
var(Residual)	4.950042	1.784803	2.441726	10.03508

 LR test vs. linear regression: chi2(3) = 55.24 Prob > chi2 = 0.0000

Note: LR test is conservative and provided only for reference

From the output (red entries) we get the following quantities:

$$\begin{aligned} \alpha_{\text{pulse}} - \alpha_{\text{CO}} &= -2.470446 \\ \tau^2 &= 8.573426 \\ \omega^2 &= 11.66695 \\ \sigma_{\text{CO}}^2 &= 4.950042 \\ \sigma_{\text{pulse}}^2 - \sigma_{\text{CO}}^2 &= 11.00559 \end{aligned}$$

SAS

Exchangeable replicates

```
20      proc mixed data = rdata ;
21          class meth item ;
22          model y = meth item / s;
23          random meth * item ;
24          repeated item / group = meth ;
25      run ;
```

```
NOTE: Convergence criteria met.
NOTE: The PROCEDURE MIXED printed pages 1-2.
NOTE: PROCEDURE MIXED used (Total process time):
      real time      3.75 seconds
      cpu time       1.52 seconds
```

The Mixed Procedure

Model Information

Data Set	WORK.RDATA
Dependent Variable	y
Covariance Structure	Variance Components
Group Effect	meth
Estimation Method	REML
Residual Variance Method	None
Fixed Effects SE Method	Model-Based
Degrees of Freedom Method	Containment

Class Level Information

Class	Levels	Values
meth	2	KL SL
item	43	1 2 3 4 5 6 7 8 9 10 11 13 14 15 16 17 18 19 20 21 22 24 25 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46

Dimensions

Covariance Parameters	3
Columns in X	46
Columns in Z	86
Subjects	1
Max Obs Per Subject	258

Number of Observations

Number of Observations Read 258
 Number of Observations Used 258
 Number of Observations Not Used 0

Iteration History

Iteration	Evaluations	-2 Res Log Like	Criterion
0	1	-353.24387418	
1	1	-376.69765836	0.00000000

Convergence criteria met.

Covariance Parameter Estimates

Cov Parm	Group	Estimate
meth*item		0.003547
item	meth KL	0.005956
item	meth SL	0.005244

Fit Statistics

-2 Res Log Likelihood	-376.7
AIC (smaller is better)	-370.7
AICC (smaller is better)	-370.6
BIC (smaller is better)	-363.3

Solution for Fixed Effects

Effect	meth	item	Estimate	Standard Error	DF	t Value	Pr > t
Intercept			1.6277	0.05259	42	30.95	<.0001
meth	KL		0.04488	0.01587	42	2.83	0.0071
meth	SL		0
item		1	0.01703	0.07356	42	0.23	0.8180
item		2	-0.8483	0.07356	42	-11.53	<.0001
item		3	1.1496	0.07356	42	15.63	<.0001
item		4	-0.9908	0.07356	42	-13.47	<.0001
item		5	1.2185	0.07356	42	16.56	<.0001
item		6	-0.7503	0.07356	42	-10.20	<.0001
item		7	-0.1674	0.07356	42	-2.28	0.0280
item		8	-0.2341	0.07356	42	-3.18	0.0028
item		9	0.6326	0.07356	42	8.60	<.0001
item		10	-0.5326	0.07356	42	-7.24	<.0001
item		11	2.1453	0.07356	42	29.16	<.0001
item		13	-0.6580	0.07356	42	-8.95	<.0001
item		14	1.2496	0.07356	42	16.99	<.0001
item		15	-0.9803	0.07356	42	-13.33	<.0001
item		16	-0.3681	0.07356	42	-5.00	<.0001
item		17	0.01630	0.07356	42	0.22	0.8257
item		18	-0.06740	0.07356	42	-0.92	0.3648
item		19	-0.06667	0.07356	42	-0.91	0.3699
item		20	0.1985	0.07356	42	2.70	0.0100
item		21	-0.4178	0.07356	42	-5.68	<.0001
item		22	0.2681	0.07356	42	3.65	0.0007
item		24	0.3341	0.07356	42	4.54	<.0001
item		25	0.06667	0.07356	42	0.91	0.3699
item		27	-0.4333	0.07356	42	-5.89	<.0001
item		28	-1.0195	0.07356	42	-13.86	<.0001
item		29	0.9489	0.07356	42	12.90	<.0001
item		30	0.3333	0.07356	42	4.53	<.0001
item		31	0.1163	0.07356	42	1.58	0.1214
item		32	-1.1721	0.07356	42	-15.93	<.0001
item		33	-0.01630	0.07356	42	-0.22	0.8257
item		34	2.1333	0.07356	42	29.00	<.0001
item		35	0.8341	0.07356	42	11.34	<.0001
item		36	0.3985	0.07356	42	5.42	<.0001
item		37	1.4837	0.07356	42	20.17	<.0001
item		38	-0.4496	0.07356	42	-6.11	<.0001

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item	39	-0.7821	0.07356	42	-10.63	<.0001
item	40	0.8689	0.07356	42	11.81	<.0001
item	41	2.4319	0.07356	42	33.06	<.0001
item	42	-0.4496	0.07356	42	-6.11	<.0001
item	43	-0.10000	0.07356	42	-1.36	0.1813
item	44	0.2667	0.07356	42	3.63	0.0008
item	45	0.1652	0.07356	42	2.25	0.0300
item	46	0

From the output (red entries) we get the following quantities:

$$\begin{aligned} \alpha_{KL} - \alpha_{SL} &= 0.04488 \\ \tau^2 &= 0.003547 \\ \sigma_{KL}^2 &= 0.005956 \\ \sigma_{SL}^2 &= 0.005244 \end{aligned}$$

Linked replicates

```
20      proc mixed data = rdata ;
21          class meth item repl ;
22          model y = meth item / s;
23          random meth*item item*repl ;
24          repeated item / group = meth ;
25      run ;
```

NOTE: Convergence criteria met.
 NOTE: The PROCEDURE MIXED printed pages 1-2.
 NOTE: PROCEDURE MIXED used (Total process time):
 real time 3:22.36
 cpu time 2:51.92

The Mixed Procedure

Model Information

```
Data Set WORK.RDATA
Dependent Variable y
Covariance Structure Variance Components
Group Effect meth
Estimation Method REML
Residual Variance Method None
Fixed Effects SE Method Model-Based
Degrees of Freedom Method Containment
```

Class Level Information

```
Class Levels Values
meth 2 CO pu
item 61 1 2 3 4 5 6 7 8 9 10 11 12 13
14 15 16 17 18 19 20 21 22 23
24 25 26 27 28 29 30 31 32 33
34 35 36 37 38 39 40 41 42 43
44 45 46 47 48 49 50 51 52 53
54 55 56 57 58 59 60 61
repl 3 1 2 3
```

Dimensions

```
Covariance Parameters 4
Columns in X 64
Columns in Z 299
Subjects 1
```

Max Obs Per Subject 354

Number of Observations

Number of Observations Read 354
 Number of Observations Used 354
 Number of Observations Not Used 0

Iteration History

Iteration	Evaluations	-2 Res Log Like	Criterion
0	1	1878.72378376	
1	2	1823.48059503	0.00000054
2	1	1823.48033506	0.00000014
3	1	1823.48031459	0.00000010
4	1	1823.48024763	0.00000000

Convergence criteria met.

Covariance Parameter Estimates

Cov Parm	Group	Estimate
meth*item		8.5734
item*repl		11.6670
item	meth CO	4.9500
item	meth pu	15.9556

Fit Statistics

-2 Res Log Likelihood 1823.5
 AIC (smaller is better) 1831.5
 AICC (smaller is better) 1831.6
 BIC (smaller is better) 1842.7

Solution for Fixed Effects

Effect	meth	item	Estimate	Standard Error	DF	t Value	Pr > t
Intercept			71.8220	3.1482	60	22.81	<.0001
meth	CO		2.4704	0.6333	60	3.90	0.0002
meth	pu		0
item		1	1.7504	4.4223	60	0.40	0.6937
item		2	-5.2713	4.4223	60	-1.19	0.2380
item		3	6.9001	4.4223	60	1.56	0.1239
item		4	-8.9778	4.4223	60	-2.03	0.0468
item		5	0.6367	4.4223	60	0.14	0.8860
item		6	4.9154	4.4223	60	1.11	0.2708
item		7	11.4569	4.4223	60	2.59	0.0120
item		8	5.3072	4.4223	60	1.20	0.2348
item		9	-2.4318	4.4223	60	-0.55	0.5844
item		10	-12.6719	4.4223	60	-2.87	0.0057
item		11	14.5007	4.4223	60	3.28	0.0017
item		12	-45.5632	4.4223	60	-10.30	<.0001
item		13	5.0723	4.4223	60	1.15	0.2559
item		14	0.6210	4.4223	60	0.14	0.8888
item		15	8.0069	4.4223	60	1.81	0.0752
item		16	1.2136	4.4223	60	0.27	0.7847
item		17	15.6657	4.7157	60	3.32	0.0015
item		18	3.2826	4.4223	60	0.74	0.4608
item		19	-0.3358	4.4223	60	-0.08	0.9397
item		20	0.7152	4.7157	60	0.15	0.8800
item		21	8.2157	4.4223	60	1.86	0.0681
item		22	1.3087	4.4223	60	0.30	0.7683
item		23	6.3324	4.4223	60	1.43	0.1574
item		24	10.0276	4.4223	60	2.27	0.0270
item		25	3.8554	4.7157	60	0.82	0.4168
item		26	4.5283	4.4223	60	1.02	0.3100

item	27	-8.5682	4.4223	60	-1.94	0.0574
item	28	-9.0693	4.4223	60	-2.05	0.0447
item	29	2.5337	4.4223	60	0.57	0.5688
item	30	4.3949	4.4223	60	0.99	0.3243
item	31	-27.4963	4.4223	60	-6.22	<.0001
item	32	7.4032	4.4223	60	1.67	0.0993
item	33	8.6273	4.4223	60	1.95	0.0557
item	34	10.4869	4.4223	60	2.37	0.0209
item	35	2.6790	4.4223	60	0.61	0.5469
item	36	4.7996	4.4223	60	1.09	0.2821
item	37	5.4239	4.4223	60	1.23	0.2248
item	38	9.2802	4.4223	60	2.10	0.0401
item	39	4.4897	5.4922	60	0.82	0.4169
item	40	-6.8656	4.4223	60	-1.55	0.1258
item	41	1.6460	4.4223	60	0.37	0.7111
item	42	-2.5947	4.4223	60	-0.59	0.5596
item	43	-18.9965	4.4223	60	-4.30	<.0001
item	44	-14.5440	4.4223	60	-3.29	0.0017
item	45	3.7834	4.4223	60	0.86	0.3957
item	46	6.2634	4.4223	60	1.42	0.1618
item	47	5.1758	4.4223	60	1.17	0.2465
item	48	-1.2806	4.4223	60	-0.29	0.7731
item	49	12.2166	4.4223	60	2.76	0.0076
item	50	-23.0847	4.7157	60	-4.90	<.0001
item	51	-19.1005	4.4223	60	-4.32	<.0001
item	52	1.3978	4.4223	60	0.32	0.7530
item	53	-1.8719	4.4223	60	-0.42	0.6736
item	54	3.1803	4.4223	60	0.72	0.4748
item	55	14.5889	4.4223	60	3.30	0.0016
item	56	11.5475	4.4223	60	2.61	0.0114
item	57	15.1005	4.4223	60	3.41	0.0012
item	58	15.2457	4.4223	60	3.45	0.0010
item	59	17.4161	4.4223	60	3.94	0.0002
item	60	9.1467	4.4223	60	2.07	0.0429
item	61	0

From the output (red entries) we get the following quantities:

$$\begin{aligned}
 \alpha_{\text{CO}} - \alpha_{\text{pulse}} &= 2.4704 \\
 \tau^2 &= 8.5734 \\
 \omega^2 &= 11.6670 \\
 \sigma_{\text{CO}}^2 &= 4.9500 \\
 \sigma_{\text{pulse}}^2 &= 15.9556
 \end{aligned}$$

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