

Overview of the package LMMstar

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This vignette describes the main functionalities of the **LMMstar** package. This package implements specific types of linear mixed models, mainly useful when having repeated observations over a discrete variable: $\mathbf{Y} = (Y_1, \dots, Y_T)$ where T can be for example be time (chronological ordering of the repetitions) or brain region (arbitrary ordering of the repetitions). Denoting by \mathbf{X} the associated covariates and $\boldsymbol{\varepsilon} = (\varepsilon_1, \dots, \varepsilon_T)$, the model can be written:

$$\mathbf{Y} = \mathbf{X}\beta + \boldsymbol{\varepsilon} \text{ where } \varepsilon \sim \mathcal{N}(0, \Omega)$$

where β are the mean parameters and the residual variance-covariance matrix, Ω , depends on a set of variance-covariance parameters (say γ) distinct of β . Key assumptions are:

- we observe n independent replicates of $\mathcal{O} = (\mathbf{Y}, \mathbf{X})$, i.e. at the cluster level, observations $(\mathcal{O}_1, \dots, \mathcal{O}_n)$ are independent. The replicates should also be identically distributed up to a categorical variable (called strata variable in the following).
- the residual variance is independent of the mean value.

Additional assumptions are necessary in presence of missing values, typically correct specification of the conditional mean to have consistent estimates of the mean parameters. This case will sometimes be exemplified by considering that only last outcome may be missing: the conditional mean $\mathbb{E}[Y_T | Y_1, Y_2, \dots, Y_{T-1}]$ is then abbreviated as $\mathbb{E}[Y_T | Y_{T-1}]$. Note that we do not require the residuals to be normally distributed to have valid estimates or statistical inference in large samples.

To get start, one should load the **LMMstar** package in the  session:

```
library(LMMstar)
```

This package is under active development. Newer package versions may include additional functionalities and fix previous bugs. The version of the package that is being used for this overview is:

```
utils::packageVersion("LMMstar")
```

```
[1] '1.1.0'
```

It is recommended to also load the following packages, as some of the methods implemented in the package are relative to a generic method implemented in other packages:

```
library(ggplot2) ## autoplot method
library(nlme)    ## ranef method
library(lava)    ## iid, information, manifest methods
```

The user interface of the **LMMstar** package is made of the following functions:

- functions to describe or visualize the dataset:
 - `scatterplot` to visualize the marginal and bivariate distribution of continuous variables.
 - `summarize` to compute summary statistics, possibly stratified on a categorical variable.
 - `summarizeNA` to identify missing data patterns.
 - `partialCor` to compute partial correlation between two variables.
- the function `mt.test` to perform multiple Student's t-Tests and adjust the results for multiple testing.
- the function `lmm` is the main function of the package which fits linear mixed models. The user can interact with `lmm` objects using:
 - `anova` to perform Wald tests, i.e. test linear combinations of coefficients ($\hat{\beta}_1 + \hat{\beta}_2 = 0$ or $\hat{\beta}_1 = \hat{\beta}_2 = 0$). The output obtained with different `lmm` can be combined using `rbind`.
 - `coef` to extract the estimated model parameters ($\hat{\beta}$ and possibly $\hat{\gamma}$).
 - `confint` to extract the estimates with their confidence intervals.
 - `effects` to evaluate marginal effects, e.g. $\mathbb{E}[\mathbb{E}[Y|X_1 = 1] - \mathbb{E}[Y|X_1 = 0]]$ when $\mathbf{X} = (X_1, X_2)$.
 - `estimate` to test non-linear combinations of coefficients (Wald test via a first order delta method, e.g. $\hat{\beta}_1/\hat{\beta}_2 = 1$).
 - `fitted` to output the fitted mean ($X\hat{\beta}$) or the conditional mean for observations with missing outcome (e.g. $X\hat{\beta} + \hat{\mathbb{E}}[\varepsilon_T|\varepsilon_{-T}]$).
 - `iid` to extract the influence function of the estimated parameters (φ), which satisfies $\sqrt{n}(\hat{\beta} - \beta) = \frac{1}{\sqrt{n}} \sum_{i=1}^n \varphi(\mathcal{O}_i) + o_p(1)$
 - `levels` to extract the reference level for the mean structure. (i.e. what (**Intercept**) refers to in presence of categorical covariates).
 - `logLik` to output the log-likelihood of the estimated model.
 - `model.tables` to extract the estimates, standard errors, p-value, and confidence intervals.
 - `plot` to obtain a diagnostic plots, partial residual plots, or a graphical display of the fitted values.
 - `predict` to compute the mean conditional on covariates and possible outcome values.
 - `profile` to display the likelihood or profile likelihood of the model.
 - `resample` to use non-parametric bootstrap or permutation test for statistical inference.
 - `residuals` to extract the observed residuals of the fitted model, possibly normalized ($\hat{\Omega}^{-\frac{1}{2}}\hat{\varepsilon}$).
 - `sigma` to extract the modeled residual variance covariance matrix ($\hat{\Omega}$).
 - `summary` to obtain a summary of the input, model fit, and estimated values.
 - `vcov` to extract the variance-covariance matrix of the mean parameters ($\hat{\Sigma}_{\hat{\beta}}$).
- the `mlmm` function to fit group-specific linear mixed models and gather the estimated coefficients.
- the `sampleRem` function to simulate longitudinal data.
- the `LMMstar.options` function enables the user to display the default values used in the **LMMstar** package. The function can also change the default values to better match the user needs.

1 Illustrative dataset

To illustrate the functionalities of the package, we will use the `gastricbypass` dataset. The long format can be imported using:

```
data(gastricbypassL, package = "LMMstar")
head(gastricbypassL)
```

	id	visit	time	weight	glucagonAUC
1	1	1	-13	127.2	20.69014
2	2	1	-13	165.2	49.92151
3	3	1	-13	109.7	42.43420
4	4	1	-13	146.2	27.51696
5	5	1	-13	113.1	29.15120
6	6	1	-13	158.8	42.70000

See `?gastricbypassL` for a presentation of the dataset. It is convenient to encode the time variable in two formats:

- numeric, e.g. here with the time in week since surgery (`time` variable taking values -13,-1,1,13 - negative times referring to before an intervention and positive times after the intervention).
- factor, e.g. here with the visit index (`visit` variable taking value 1,2,3,4)

To illustrate certain functionalities we will use an (artificial) group variable:

```
gastricbypassL$group <- as.numeric(gastricbypassL$id)%2
```

and dichotomize time as before and after the intervention:

```
gastricbypassL$baseline <- gastricbypassL$time<0
```

The corresponding wide format is

```
data(gastricbypassW, package = "LMMstar")
head(gastricbypassW)
```

	id	weight1	weight2	weight3	weight4	glucagonAUC1	glucagonAUC2	glucagonAUC3	glucagonAUC4
1	1	127.2	120.7	115.5	108.1	20.69014	20.53486	92.60017	43.43377
2	2	165.2	153.4	149.2	132.0	49.92151	58.51346	49.63298	35.74695
3	3	109.7	101.6	97.7	87.1	42.43420	25.76985	91.23980	83.13719
4	4	146.2	142.4	136.7	123.0	27.51696	27.55224	59.35960	21.37069
5	5	113.1	105.6	99.9	87.7	29.15120	NA	86.85942	57.96993
6	6	158.8	143.6	134.6	108.7	42.70000	31.61559	53.40800	37.63608

for which we can also add the group variable:

```
gastricbypassW$group <- as.numeric(gastricbypassW$id)%2
```

In some cases we will, for comparison, perform complete case analyses with the following dataset:

```
gastricbypassL.NNA <- gastricbypassL[!is.na(gastricbypassL$glucagonAUC),]
```

2 Visualization & descriptive statistics

2.1 Graphical display

A scatterplot of the data can be obtained by specifying which columns to display when using the wide format:

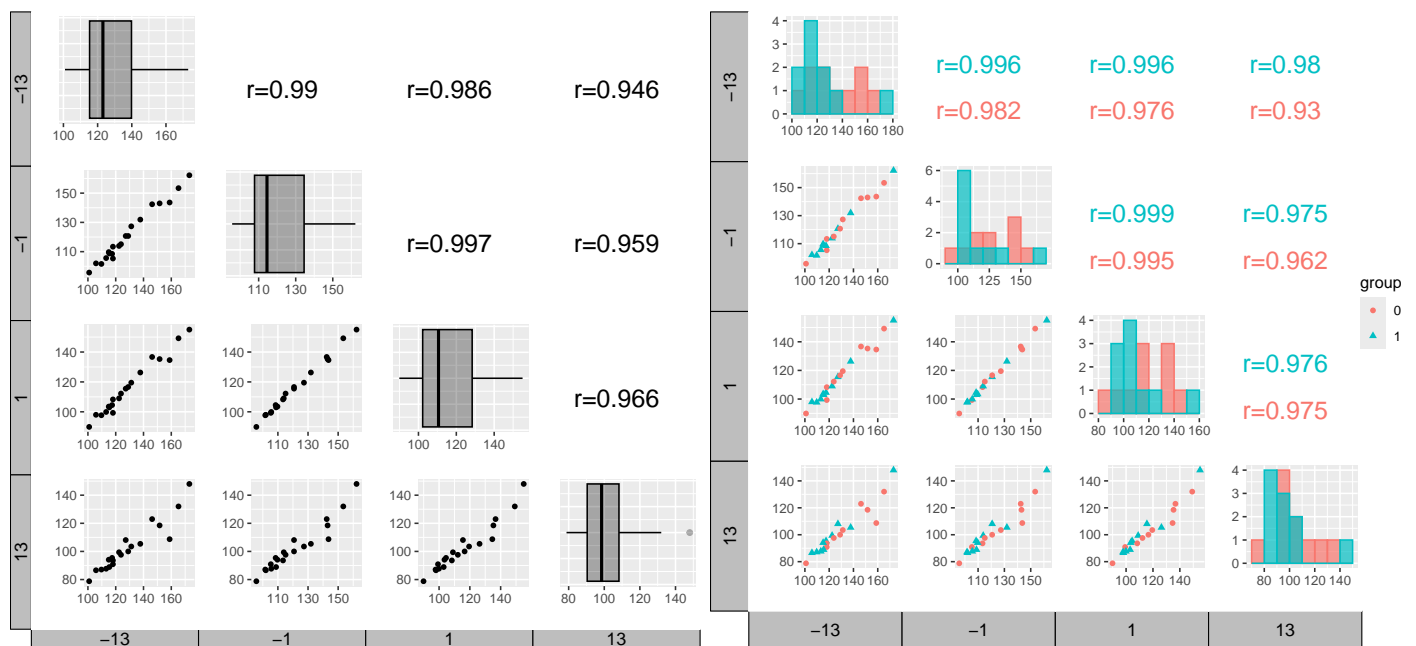
```
scatterplot(gastricbypassW, ## left panel
            columns = c("weight1", "weight2", "weight3", "weight4"))
```

When using the long format, a formula should describe the structure of the data: `outcome ~ order|cluster`

- the left hand side indicates the values to be displayed (here weight)
- the right hand side indicates the ordering of the repetitions (here over time) and how the repetitions are grouped within clusters (here within subject).

When calling `scatterplot`, the argument `group` leads to different color per group and the argument `type.diag` enables to use histograms (or density plots) instead of boxplots:

```
scatterplot(weight~time|id, data = gastricbypassL, ## right panel
            type.diag = "hist", group = "group")
```



By default the resulting object will be of class `list`. A `ggplot2` object can be obtained by setting the argument `facet` to `"grid2"`. This requires to have installed the package `ggh4x` and will produce a slightly different graphical display.

There is (currently) not dedicated function to obtain spaghetti plots. Instead one can use the `ggplot2` package with the long format, e.g.:

```
gg.spa <- ggplot(gastricbypassL, aes(x=time,y=weight,group=id,color=id))
gg.spa <- gg.spa + geom_point() + geom_line()
gg.spa
```

2.2 Missing data patterns

The `summarizeNA` function identifies the possible combinations of observed/missing data:

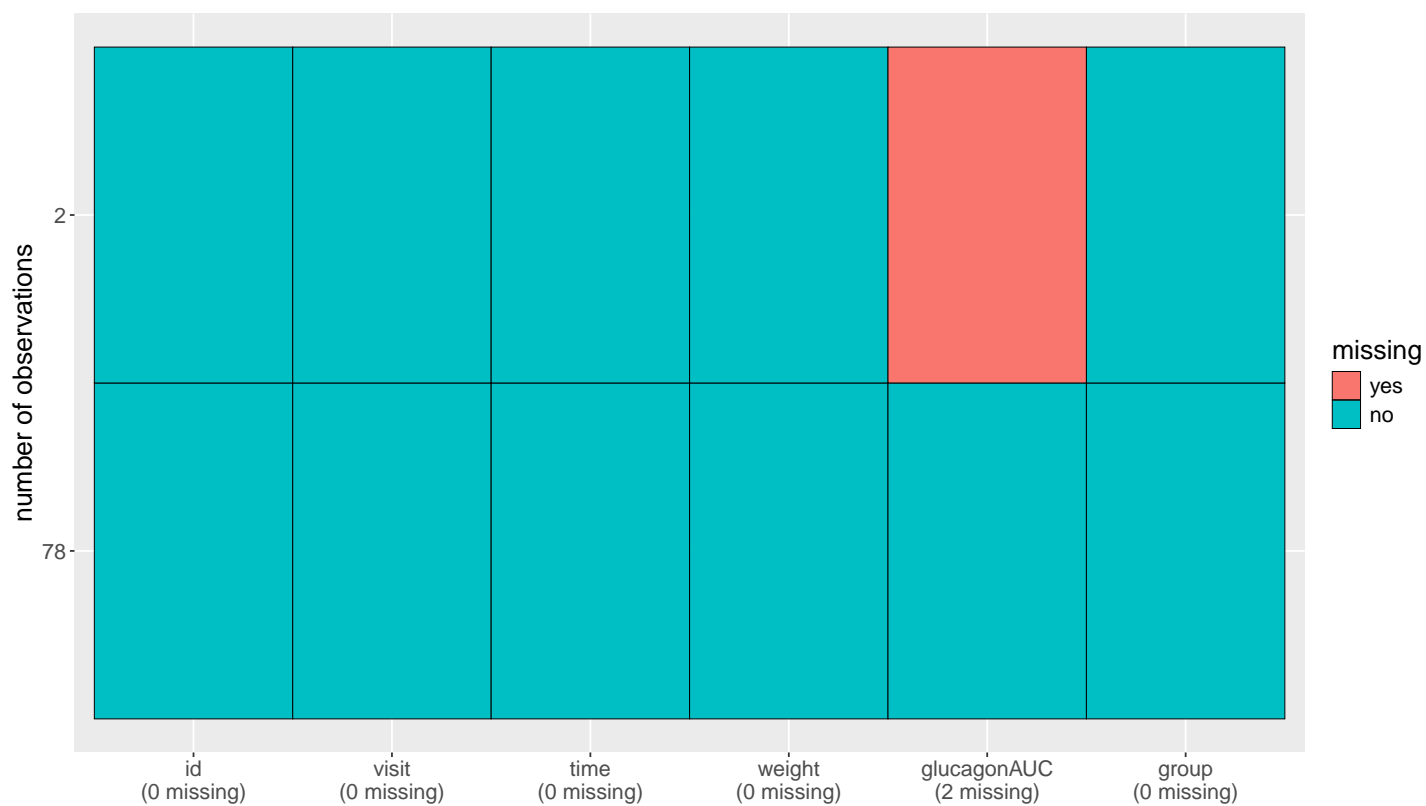
```
mp <- summarizeNA(gastricbypassL)
mp
```

frequency	missing.pattern	n.missing	id	visit	time	weight	glucagonAUC	group	baseline
78	0000000	0	0	0	0	0	0	0	0
2	0000100	1	0	0	0	0	1	0	0

A graphical representation can be obtained using `plot`:

```
plot(mp)
```

See `help(plot.summarizeNA)` for options to customize the graphical display.



2.3 Summary statistics

Mean, standard deviation, and other summary statistic can be computed with respect to a categorical variable (typically time) using the `summarize` function:

(⚠ this function has the same name as a function from the `dplyr` package. If you have loaded `dplyr`, you should use `LMMstar::summarize`)

```
sss <- summarize(weight+glucagonAUC ~ time, data = gastricbypassL, na.rm = TRUE)
print(sss, digits = 3)
```

	outcome	time	observed	missing	mean	sd	min	q1	median	q3	max
1	weight	-13	20	0	129.0	20.3	100.90	115.3	123.1	139.8	173.0
2		-1	20	0	121.2	18.9	95.70	107.8	114.5	134.5	162.2
3		1	20	0	115.7	18.3	89.90	102.2	110.6	128.4	155.0
4		13	20	0	102.4	17.1	78.80	90.4	98.5	108.2	148.0
5	glucagonAUC	-13	20	0	32.3	15.5	10.28	21.3	27.9	42.5	69.1
6		-1	19	1	29.7	13.7	9.87	21.2	25.8	33.6	67.7
7		1	19	1	76.9	27.9	35.85	56.5	73.8	91.9	135.9
8		13	20	0	52.0	21.0	21.37	37.2	51.2	57.9	109.2

Specifying a cluster (`id`) and ordering variable (`time`) enable to output correlation matrices:

(⚠ there should be no duplicated value of the ordering variable within cluster)

```
sss2 <- summarize(weight ~ time|id, data = gastricbypassL, na.rm = TRUE)
print(sss2, digits = 3)
```

	time	observed	missing	mean	sd	min	q1	median	q3	max
1	-13	20	0	129	20.3	100.9	115.3	123.1	140	173
2	-1	20	0	121	18.9	95.7	107.8	114.5	135	162
3	1	20	0	116	18.3	89.9	102.2	110.6	128	155
4	13	20	0	102	17.1	78.8	90.4	98.5	108	148

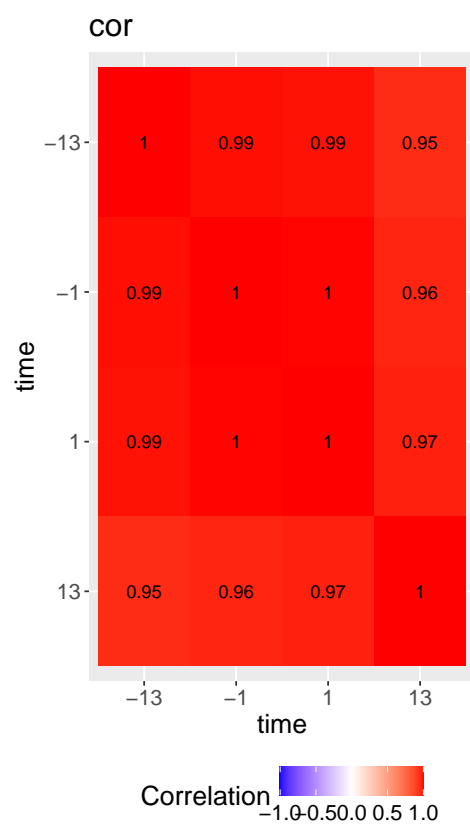
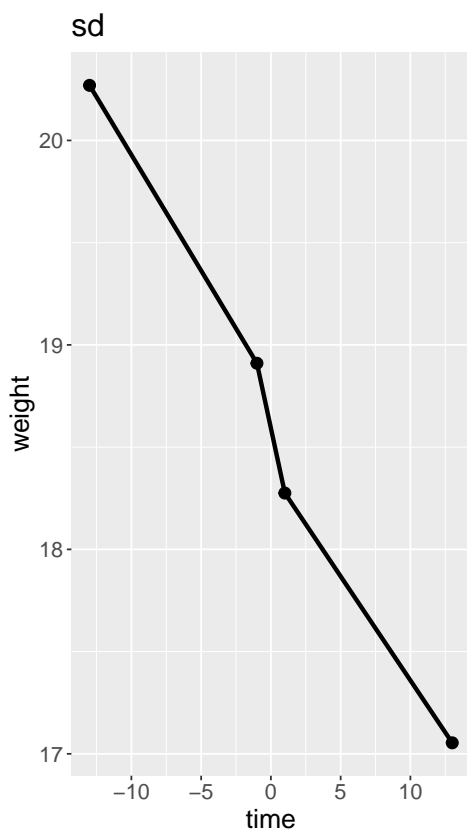
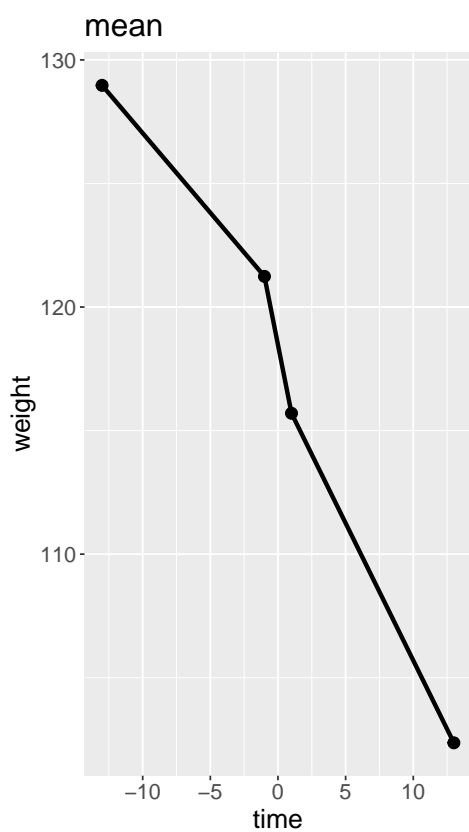
Pearson's correlation:

	-13	-1	1	13
-13	1.000	0.990	0.986	0.946
-1	0.990	1.000	0.997	0.959
1	0.986	0.997	1.000	0.966
13	0.946	0.959	0.966	1.000

Graphical displays of the summary statistics can be obtained via the `plot` method, where the argument `type` specifies the summary statistic to be displayed:

```
plot(sss2, type = "mean") ## left panel
plot(sss2, type = "sd") ## middle panel
plot(sss2, type = "cor") ## right panel
```

See `help(plot.summarize)` for options to customize the graphical display.



2.4 Correlation and partial correlations

The `partialCor` function can be used to evaluate group-specific correlations, e.g.:

```
partialCor(weight + glucagonAUC ~ 1, by = "group", data = gastricbypassL)
```

		estimate	se	df	lower	upper	p.value
0:	<code>rho(weight,glucagonAUC)</code>	-0.328	0.143	21.8	-0.587	-0.00886	0.0447
1:	<code>rho(weight,glucagonAUC)</code>	-0.354	0.141	22.5	-0.607	-0.03631	0.0313

This will lead to the same estimate as the `cor.test` function (Pearson correlation):

```
gastricbypassL.0 <- gastricbypassL[gastricbypassL$group==0,]  
rho <- cor.test(gastricbypassL.0$weight, gastricbypassL.0$glucagonAUC)  
c(rho$estimate, p.value = rho$p.value)
```

	cor	p.value
	-0.32848119	0.03850536

However the p-value may differ, especially in small samples, as `partialCor` uses a different (and probably more crude) small sample approximation for the estimator's distribution. Nevertheless `partialCor` enables to compare correlation coefficients across groups, by specifying the argument `effects`:

```
partialCor(weight + glucagonAUC ~ 1, by = "group", effects = "Dunnett",  
           data = gastricbypassL)
```

		estimate	se	df	lower	upper	p.value
1:	<code>rho(weight,glucagonAUC)</code>	-0.0255	NA	NA	NA	NA	0.899
0:	<code>rho(weight,glucagonAUC)</code>						

Partial correlations can be also computed by specifying covariate to adjust for on the right-hand side:

```
partialCor(weight4 + glucagonAUC4 ~ weight1,  
           data = gastricbypassW)
```

		estimate	se	df	lower	upper	p.value
	<code>rho(weight4,glucagonAUC4)</code>	0.112	0.233	9.12	-0.397	0.568	0.645

When the set of covariates is outcome-dependent, a list of formulas can be used instead:

```
partialCor(list(weight1 ~ glucagonAUC1, weight4 ~ glucagonAUC4),  
           data = gastricbypassW)
```

		estimate	se	df	lower	upper	p.value
	<code>rho(weight1,weight4)</code>	0.946	0.0252	26.4	0.861	0.979	5.51e-08

These partial correlations are defined as the residual correlation between the outcomes, i.e. the correlation once the covariate effects have been subtracted from the outcome, and a linear mixed model is used to estimate them.

3 Multiple Student's t-tests

When working with multiple outcomes and having no missing data, mean comparisons between exposure groups can be carried out using Student's t-tests at each timepoint, e.g.:

```
restt <- t.test(weight1 ~ group, data = gastricbypassW)
c(estimate = unname(diff(restt$estimate)), p.value = restt$p.value)
```

```
estimate    p.value
-10.6000000  0.2528174
```

And so on for the three other timepoints. Moreover results would typically need to be adjusted for multiple comparisons, e.g. when looking for any mean difference. This can be facilitated by

```
## single step max-test adjustment (see help(confint.Wald_lmm) for details)
mt.test(weight1+weight2+weight3+weight4~group, data = gastricbypassW)
```

	by	parameter	estimate	se	df	lower	upper	p.value
1	weight1	group	-10.60	8.971747	17.96464	-30.93243	9.732430	0.3160768
2	weight2	group	-9.50	8.395143	17.98540	-28.52569	9.525687	0.3399966
3	weight3	group	-8.92	8.129458	17.95876	-27.34357	9.503573	0.3569464
4	weight4	group	-4.59	7.760674	17.68244	-22.17781	12.997807	0.6640834

The method used to adjust confidence intervals and p-values for multiple comparisons can be specified via the `method` argument, e.g.:

```
## no adjustment
mt.test(weight1+weight2+weight3+weight4~group, data = gastricbypassW, method = "none")
```

	by	parameter	estimate	se	df	lower	upper	p.value
1	weight1	group	-10.60	8.971747	17.96464	-29.45160	8.251601	0.2528144
2	weight2	group	-9.50	8.395143	17.98540	-27.13857	8.138567	0.2726630
3	weight3	group	-8.92	8.129458	17.95876	-26.00217	8.162169	0.2870297
4	weight4	group	-4.59	7.760674	17.68244	-20.91558	11.735584	0.5617090

```
## bonferroni adjustment
mt.test(weight1+weight2+weight3+weight4~group, data = gastricbypassW, method = "bonferroni")
```

	by	parameter	estimate	se	df	lower	upper	p.value
1	weight1	group	-10.60	8.971747	17.96464	-35.49775	14.29775	1
2	weight2	group	-9.50	8.395143	17.98540	-32.79464	13.79464	1
3	weight3	group	-8.92	8.129458	17.95876	-31.48110	13.64110	1
4	weight4	group	-4.59	7.760674	17.68244	-26.16472	16.98472	1

4 Linear mixed model (LMM)

4.1 Classical covariance patterns

Several build-in covariance patterns can be used when specifying the linear model. The most basic ones are the **identity** structure:

```
eId.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,  
              structure = "ID", data = gastricbypassL)  
eId.lmm  
cat(" modeled residual variance-covariance: \n");sigma(eId.lmm)
```

Linear regression

```
outcome/cluster/time: glucagonAUC/id/time  
data                  : 78 observations from 20 clusters  
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis  
                      1 variance (sigma)  
log-restr.likelihood: -316.461119970244  
convergence           : TRUE (0 iterations)  
modeled residual variance-covariance:  
      -13      -1       1       13  
-13 381.3536  0.0000  0.0000  0.0000  
-1   0.0000 381.3536  0.0000  0.0000  
1    0.0000  0.0000 381.3536  0.0000  
13   0.0000  0.0000  0.0000 381.3536
```

and the **independence** structure:

```
eInd.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,  
              structure = "IND", data = gastricbypassL)  
eInd.lmm  
cat(" modeled residual variance-covariance: \n");sigma(eInd.lmm)
```

Linear regression with heterogeneous residual variance

```
outcome/cluster/time: glucagonAUC/id/time  
data                  : 78 observations from 20 clusters  
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis  
                      4 variance (sigma k.-1 k.1 k.13)  
log-restr.likelihood: -310.428096419287  
convergence           : TRUE (0 iterations)  
modeled residual variance-covariance:  
      -13      -1       1       13  
-13 209.4422  0.0000  0.0000  0.0000  
-1   0.0000 174.8103  0.0000  0.0000  
1    0.0000  0.0000 768.2278  0.0000  
13   0.0000  0.0000  0.0000 382.9524
```

The most common linear mixed model uses a **compound symmetry** structure:

```
eCS.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,
               structure = "CS", data = gastricbypassL)
eCS.lmm
cat(" modeled residual variance-covariance: \n");sigma(eCS.lmm)
```

Linear Mixed Model with a compound symmetry covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time
data                  : 78 observations from 20 clusters
parameter            : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis
                      1 variance (sigma)
                      1 correlation (rho(id))
log-restr.likelihood: -314.394203759159
convergence           : TRUE (6 iterations)
modeled residual variance-covariance:
      -13      -1       1      13
-13 380.57991  82.74055  82.74055  82.74055
-1   82.74055 380.57991  82.74055  82.74055
1    82.74055 82.74055 380.57991  82.74055
13   82.74055 82.74055 82.74055 380.57991
```

A more flexible model can be obtained with a **toeplitz** covariance matrix:

```
eTOE.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,
               structure = "TOEPLITZ", data = gastricbypassL)
eTOE.lmm
cat(" modeled residual correlation: \n");cov2cor(sigma(eTOE.lmm))
```

Linear Mixed Model with a block Toeplitz covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time
data                  : 78 observations from 20 clusters
parameter            : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis
                      4 variance (sigma k.-1 k.1 k.13)
                      4 correlation (rho(12) rho(14) rho(26) rho(2))
log-restr.likelihood: -297.525485582536
convergence           : TRUE (15 iterations)
modeled residual correlation:
      -13      -1       1      13
-13 1.00000000 0.70001988 0.09361507 -0.08296276
-1   0.70001988 1.00000000 0.01679455  0.09361507
1    0.09361507 0.01679455 1.00000000  0.70001988
13   -0.08296276 0.09361507 0.70001988  1.00000000
```

And an even more flexible model can be obtained with an **unstructured** covariance matrix:

```
eUN.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,  
               structure = "UN", data = gastricbypassL)  
eUN.lmm  
cat(" modeled residual variance-covariance: \n");sigma(eUN.lmm)
```

Linear Mixed Model with an unstructured covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time  
data                  : 78 observations from 20 clusters  
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group visit4:group)  
                      4 variance (sigma k.-1 k.1 k.13)  
                      6 correlation (rho(-13,-1) rho(-13,1) rho(-13,13) rho(-1,1) rho(-1,13) rho(-1,13))  
log-restr.likelihood: -295.314056198772  
convergence           : TRUE (8 iterations)  
modeled residual variance-covariance:  
      -13      -1       1       13  
-13 209.44221 150.250246 106.399969 -24.20227  
-1  150.25025 168.113758  1.306397 -23.88443  
1   106.39997  1.306397 748.076864 288.18395  
13  -24.20227 -23.884434 288.183947 382.95234
```

Stratification of the covariance structure on a categorical variable is also possible:

- e.g. to get a **stratified compound symmetry**

```
eSCS.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,  
               structure = CS(group~1), data = gastricbypassL)  
eSCS.lmm
```

Linear Mixed Model with a stratified compound symmetry covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time  
data                  : 78 observations from 20 clusters  
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group visit4:group)  
                      2 variance (sigma:0 sigma:1)  
                      2 correlation (rho(id):0 rho(id):1)  
log-restr.likelihood: -314.123797063042  
convergence           : TRUE (7 iterations)
```

- e.g. **stratified unstructured** covariance matrix:

```
eSUN.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,
               structure = UN(group~1), data = gastricbypassL)
eSUN.lmm
```

Linear Mixed Model with a stratified unstructured covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time
data                  : 78 observations from 20 clusters
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group visit4:group)
                       8 variance (sigma:0 sigma:1 k.-1:0 k.1:0 k.13:0 k.-1:1 k.1:1 k.13:1)
                       12 correlation (rho(-13,-1):0 rho(-13,1):0 rho(-13,13):0 rho(-1,1):0 rho(-1,13):0 rho(1,-1):0 rho(1,1):0 rho(1,13):0)
log-restr.likelihood: -286.536815485471
convergence            : TRUE (10 iterations)
```

with modeled residual variance-covariance:

sigma(eSCS.lmm)					sigma(eSUN.lmm)				
\$'0'					\$'0'				
	-13	-1	1	13		-13	-1	1	13
-13	334.28864	50.78164	50.78164	50.78164	-13	309.8534	251.51195	102.18885	-42.24970
-1	50.78164	334.28864	50.78164	50.78164	-1	251.5120	274.75220	-79.81124	-90.71752
1	50.78164	50.78164	334.28864	50.78164	1	102.1888	-79.81124	579.10966	163.76749
13	50.78164	50.78164	50.78164	334.28864	13	-42.2497	-90.71752	163.76749	173.43923
\$'1'					\$'1'				
	-13	-1	1	13		-13	-1	1	13
-13	428.4618	115.0917	115.0917	115.0917	-13	109.030904	48.66651	104.90774	-6.154856
-1	115.0917	428.4618	115.0917	115.0917	-1	48.666509	59.39530	93.97619	43.214357
1	115.0917	115.0917	428.4618	115.0917	1	104.907735	93.97619	967.58280	450.889944
13	115.0917	115.0917	115.0917	428.4618	13	-6.154856	43.21436	450.88994	592.465459

Finally the some covariance patterns like the compound symmetry structure may depend on covariates:

- e.g. to obtain a **block compound symmetry** structure¹:

```
eBCS.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,
               structure = CS(~baseline, type = "homogeneous"), data = gastricbypassL)
eBCS.lmm
cat(" modeled residual variance-covariance: \n");sigma(eBCS.lmm)
```

Linear Mixed Model with a block compound symmetry covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time
data                  : 78 observations from 20 clusters
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis
                      1 variance (sigma)
                      2 correlation (rho(id/baseline) rho(id))
log-restr.likelihood: -308.994835006264
convergence           : TRUE (6 iterations)
modeled residual variance-covariance:
      -13      -1       1       13
-13 380.95687 226.40271 15.46503 15.46503
-1 226.40271 380.95687 15.46503 15.46503
1 15.46503 15.46503 380.95687 226.40271
13 15.46503 15.46503 226.40271 380.95687
```

- e.g. to obtain a **block unstructured** covariance matrix:

```
eBUN.lmm <- lmm(glucagonAUC ~ visit*group, repetition = ~time|id,
               structure = CS(~baseline, type = "heterogeneous"), data = gastricbypassL)
eBUN.lmm
cat(" modeled residual variance-covariance: \n");sigma(eBUN.lmm)
```

Linear Mixed Model with a block unstructured covariance matrix

```
outcome/cluster/time: glucagonAUC/id/time
data                  : 78 observations from 20 clusters
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group vis
                      2 variance (sigma k.TRUE)
                      3 correlation (rho(FALSE) rho(FALSE,TRUE) rho(TRUE))
log-restr.likelihood: -300.047474124556
convergence           : TRUE (7 iterations)
modeled residual variance-covariance:
      -13      -1       1       13
-13 189.41998 150.35555 15.35285 15.35285
-1 150.35555 189.41998 15.35285 15.35285
1 15.35285 15.35285 570.90792 300.07084
13 15.35285 15.35285 300.07084 570.90792
```

¹similar to nested random effects

4.2 User-specific covariance patterns

It is possible input user-specific covariance patterns under the following model for the residuals:

$$\Omega = \sigma^\top R \sigma$$

- $\sigma = f(\theta_\sigma, Z_\sigma)$ is a vector of residual standard errors depending on a vector of parameters θ_σ and possible covariates via the design matrix Z_σ .
- $R = g(\theta_R, Z_R)$ is a matrix of residual correlations depending on a vector of parameters θ_R and possible covariates via the design matrix Z_R .

To be more concrete, consider the following correlation matrix

```
rho.2block <- function(p,n.time,X){
  rho <- matrix(1, nrow = n.time, ncol = n.time)
  rho[1,2] <- rho[2,1] <- rho[4,5] <- rho[5,4] <- p["rho1"]
  rho[1,3] <- rho[3,1] <- rho[4,6] <- rho[6,4] <- p["rho2"]
  rho[2,3] <- rho[3,2] <- rho[5,6] <- rho[6,5] <- p["rho3"]
  rho[4:6,1:3] <- rho[1:3,4:6] <- p["rho4"]
  return(rho)
}
Rho <- rho.2block(p = c(rho1=0.25,rho2=0.5,rho3=0.4,rho4=0.1),
  n.time = 6)
Rho
```

```
      [,1] [,2] [,3] [,4] [,5] [,6]
[1,] 1.00 0.25  0.5 0.10 0.10  0.1
[2,] 0.25 1.00  0.4 0.10 0.10  0.1
[3,] 0.50 0.40  1.0 0.10 0.10  0.1
[4,] 0.10 0.10  0.1 1.00 0.25  0.5
[5,] 0.10 0.10  0.1 0.25 1.00  0.4
[6,] 0.10 0.10  0.1 0.50 0.40  1.0
```

and the corresponding dataset:

```
set.seed(11)
Y <- mvtnorm::rmvnorm(1000, mean = rep(0,6), sigma = Rho)
dfW <- cbind(id = 1:NROW(Y), as.data.frame(Y))
dfL <- reshape2::melt(dfW, id.vars = "id", variable.name = "time")
dfL[dfL$id %in% 1:2,]
```

	id	time	value		id	time	value	
	1	1	V1	-0.9842079	2	2	V1	1.2402726
	1001	1	V2	-0.3681245	1002	2	V2	0.6494215
	2001	1	V3	-1.6174652	2002	2	V3	0.3272105
	3001	1	V4	-1.4994103	3002	2	V4	-1.0626973
	4001	1	V5	0.7493107	4002	2	V5	-0.9013244
	5001	1	V6	-1.0719657	5002	2	V6	-0.6696714

To estimate the corresponding mixed model we first define a new covariance structure:

```
myStruct <- CUSTOM(~time,
  FCT.sigma = function(p,n.time,X){rep(p,n.time)}, ## function f
  init.sigma = c("sigma"=1),
  FCT.rho = rho.2block, ## function g
  init.rho = c("rho1"=0.25,"rho2"=0.25,"rho3"=0.25,"rho4"=0.25))
```

and then call `lmm` with this structure structure:

```
e.lmmCUSTOM <- lmm(value~time, repetition=~time|id,
  structure = myStruct, data=dfL,
  df = FALSE) ## df = FALSE to save computation time
logLik(e.lmmCUSTOM)
```

```
[1] -7962.243
```

The optimization procedure may be slow but should eventually reaches an optimum. We can then output the estimated correlation matrix:

```
cov2cor(sigma(e.lmmCUSTOM))
```

	V1	V2	V3	V4	V5	V6
V1	1.00000000	0.24898095	0.50058994	0.09053785	0.09053785	0.09053785
V2	0.24898095	1.00000000	0.36110943	0.09053785	0.09053785	0.09053785
V3	0.50058994	0.36110943	1.00000000	0.09053785	0.09053785	0.09053785
V4	0.09053785	0.09053785	0.09053785	1.00000000	0.24898095	0.50058994
V5	0.09053785	0.09053785	0.09053785	0.24898095	1.00000000	0.36110943
V6	0.09053785	0.09053785	0.09053785	0.50058994	0.36110943	1.00000000

Comparison to build-in structure: consider the following model using a build-in compound symmetry structure:

```
system.time(
  e.lmmDEFAULT.CS <- lmm(value~time, repetition = ~time|id,
    structure = "CS", data = dfL,
    df = FALSE)
)
```

```
user  system elapsed
0.097   0.000   0.097
```

Using instead `CUSTOM` to specifying this structure:

```
myCS <- CUSTOM(~1,
  FCT.sigma = function(p,n.time,X){rep(p,n.time)},
  init.sigma = c("sigma"=1),
  FCT.rho = function(p,n.time,X){p+diag(1-p,n.time,n.time)},
  init.rho = c("rho"=0.5))
```


is considerably slower than using the pre-specified structure:

```
system.time(  
  e.lmmCUSTOM.CS <- lmm(value~time, repetition = ~time|id,  
                        structure = myCS, data = dfL,  
                        df = FALSE)  
)
```

```
user system elapsed  
0.952  0.019  0.972
```

but will lead to the same estimates:

```
logLik(e.lmmDEFAULT.CS)  
logLik(e.lmmCUSTOM.CS)
```

```
[1] -8186.859  
[1] -8186.859
```

There are two reasons for the slower execution time: slower evaluation of the derivatives (since they are obtained by numerical differentiation) and worse starting point, as reflected by the larger number of iterations needed to reach convergence:

```
e.lmmDEFAULT.CS$opt$n.iter  
e.lmmCUSTOM.CS$opt$n.iter
```

```
[1] 1  
[1] 4
```

Faster execution time can be obtained by specifying the first and second derivative regarding each parameter:

```
myCS.wD <- CUSTOM(~1,  
  FCT.sigma = function(p,n.time,X){rep(p,n.time)},  
  dFCT.sigma = function(p,n.time,X){list(sigma = rep(1,n.time))},  
  d2FCT.sigma = function(p,n.time,X){list(sigma = rep(0,n.time))},  
  init.sigma = c("sigma"=1),  
  FCT.rho = function(p,n.time,X){p+diag(1-p,n.time,n.time)},  
  dFCT.rho = function(p,n.time,X){list(rho = 1-diag(1,n.time,n.time))},  
  d2FCT.rho = function(p,n.time,X){list(rho = matrix(0,n.time,n.time))},  
  init.rho = c("rho"=0.5))
```

```
system.time(  
  e.lmmCUSTOMwD.CS <- lmm(value~time,  
                          repetition = ~time|id,  
                          structure = myCS.wD,  
                          data = dfL, df = FALSE  
  )  
)
```

```
user system elapsed  
0.699  0.004  0.703
```

4.3 Estimation procedure

Initialiation: by default the mean parameters are initialized using Ordinary Least Squares (OLS) and the variance and correlation parameters are initialized by minimizing the difference between the observed and residuals variance-covariance matrix. These values can be visualized by specifying the argument **control**:

```
eCS.lmm.bis <- update(eCS.lmm, control = list(trace = 2))
```

Initialization:

(Intercept)	visit2	visit3	visit4	group	visit2:group	visit3:group
38.7289746	-4.7343331	31.4330293	4.5213806	-12.8246159	3.7594585	27.0014967
visit4:group	sigma	rho(id)				
30.2239055	19.5282760	0.2281903				

Loop:

(Intercept)	visit2	visit3	visit4	group	visit2:group	visit3:group
38.7289746	-4.7343331	31.4330293	4.5213806	-12.8246159	3.8033683	27.4810338
visit4:group	sigma	rho(id)				
30.2239055	19.5084575	0.2174065				

Convergence after 6 iterations: max score=1.241263e-05 | max change in coefficient=4.516656e-06

It is possible to input user-defined value:

- for all parameters (vector)

```
init.all <- coef(eCS.lmm, effects = "all")
eCS.lmm.bis <- update(eCS.lmm, control = list(init = init.all, trace = 1))
```

Convergence after 0 iteration: max score=1.241263e-05

- the mean parameters only (vector)

```
init.mean <- coef(eCS.lmm, effects = "mean")
eCS.lmm.bis <- update(eCS.lmm, control = list(init = init.mean, trace = 2))
```

Initialization:

(Intercept)	visit2	visit3	visit4	group	visit2:group	visit3:group
38.7289746	-4.7343331	31.4330293	4.5213806	-12.8246159	3.8033683	27.4810338
visit4:group	sigma	rho(id)				
30.2239055	19.5290394	0.2284891				

Loop:

(Intercept)	visit2	visit3	visit4	group	visit2:group	visit3:group
38.7289746	-4.7343331	31.4330293	4.5213806	-12.8246159	3.8033683	27.4810339
visit4:group	sigma	rho(id)				
30.2239055	19.5084576	0.2174065				

Convergence after 6 iterations: max score=1.489256e-05 | max change in coefficient=5.386569e-06

- a full data variance-covariance matrix (matrix).

```
init.vcov <- sigma(eCS.lmm)
eCS.lmm.bis <- update(eCS.lmm, control = list(init = init.vcov, trace = 1))
```

Convergence after 0 iteration: max score=1.241263e-05

Optimizer: by default the optimizer is a Newton Raphson algorithm with backtracking. At each iteration:

- it computes the first two moments (score, information) according to the current parameters values.
- it updates the variance-covariance parameters according to the gradient multiplied by the inverse of the information.
- it updates the mean parameters by generalized least squares (using the updated variance-covariance parameters).
- it checks whether the log-likelihood at the updated estimates is well defined and higher than at the previous estimates. If this is not the case, the step is re-run with half the update of the variance-covariance parameters (backtracking).

One can modify the maximum number of iterations (**n.iter**), maximum number of backtracking steps (**n.backtracking**), the maximum score (absolute) value over all parameters (**tol.score**) and (absolute) maximum difference in parameter value between two iterations (**tol.param**) used to declare convergence. It is also possible to use another optimizer (**optimizer**). All these elements should be passed to the argument **control** of **lmm** using a list.

4.4 Model output

The `summary` method can be used to display the main information relative to the model fit:

```
summary(eUN.lmm)
```

Linear Mixed Model

Dataset: `gastricbypassL`

- 20 clusters
- 78 observations were analyzed, 2 were excluded because of missing values
- between 3 and 4 observations per cluster

Summary of the outcome and covariates:

```
$ glucagonAUC: num  20.7 49.9 42.4 27.5 29.2 ...
$ visit       : Factor w/ 4 levels "1","2","3","4": 1 1 1 1 1 1 1 1 1 1 ...
$ group       : num   1 0 1 0 1 0 1 0 1 0 ...
reference level: visit=1
```

Estimation procedure

- Restricted Maximum Likelihood (REML)
- log-likelihood : -295.3141
- parameters: mean = 8, variance = 4, correlation = 6
- convergence: TRUE (8 iterations)
- largest |score| = 4.677128e-05 for $\rho(-1,1)$
|change| = 1.68033723859651e-05 for visit3:group

Residual variance-covariance: unstructured

- correlation structure: ~0 + time
 -13 -1 1 13
-13 1.0000 0.80072 0.26880 -0.0855
-1 0.8007 1.00000 0.00368 -0.0941
1 0.2688 0.00368 1.00000 0.5384
13 -0.0855 -0.09413 0.53842 1.0000
- variance structure: ~time
 standard.deviation ratio
sigma.-13 14.5 1.000
sigma.-1 13.0 0.896
sigma.1 27.4 1.890
sigma.13 19.6 1.352

Fixed effects: glucagonAUC ~ visit * group

	estimate	se	df	lower	upper	p.value	
(Intercept)	38.729	4.576	18	29.114	48.344	< 1e-04	***
visit2	-4.734	2.776	17.5	-10.577	1.109	0.10574	
visit3	31.433	8.63	17.6	13.272	49.594	0.00192	**
visit4	4.521	8.005	18	-12.297	21.34	0.57917	
group	-12.825	6.472	18	-26.422	0.773	0.06302	.
visit2:group	3.987	3.996	17.9	-4.41	12.383	0.33169	
visit3:group	27.571	12.42	17.8	1.461	53.682	0.03963	*
visit4:group	30.224	11.321	18	6.439	54.008	0.01562	*

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Columns lower and upper contain 95% pointwise confidence intervals for each coefficient.

Model-based standard errors are derived from the observed information (column se).

Degrees of freedom were computed using a Satterthwaite approximation (column df).

Note: the calculation of the degrees of freedom, especially when using the observed information can be quite slow. Setting the arguments `df` to `FALSE` and `type.information` to `"expected"` when calling `lmm` should lead to a more reasonable computation time.

4.5 Extract estimated coefficients

The value of the estimated coefficients can be output using `coef`:

```
coef(eUN.lmm)
```

(Intercept)	visit2	visit3	visit4	group	visit2:group	visit3:group
38.728975	-4.734333	31.433029	4.521381	-12.824616	3.986559	27.571424
visit4:group						
30.223906						

Variance coefficients can be output by specifying the `effects` argument:

```
coef(eUN.lmm, effects = "variance")
```

sigma	k.-1	k.1	k.13
14.4721183	0.8959206	1.8899095	1.3521979

The first coefficient is the residual standard deviation at the reference timepoint (here -13 week) and the remaining coefficient the residual standard deviation at later timepoints relative to the reference timepoint. It is possible to apply specific transformation on the variance coefficients, for instance to obtain the residual variance at each timepoint:

```
coef(eUN.lmm, effects = "variance", transform.k = "sd")
```

sigma.-13	sigma.-1	sigma.1	sigma.13
14.47212	12.96587	27.35099	19.56917

4.6 Extract estimated coefficient and associated uncertainty

The uncertainty about the mean coefficients can be obtained using the `model.tables` method ²:

```
model.tables(eUN.lmm)
```

	estimate	se	df	lower	upper	p.value
(Intercept)	38.728975	4.576486	18.00314	29.114255	48.3436939	1.089106e-07
visit2	-4.734333	2.775887	17.54267	-10.577172	1.1085054	1.057398e-01
visit3	31.433029	8.629711	17.58484	13.271949	49.5941098	1.922898e-03
visit4	4.521381	8.004993	17.99506	-12.296816	21.3395773	5.791699e-01
group	-12.824616	6.472128	18.00316	-26.421881	0.7726496	6.301532e-02
visit2:group	3.986559	3.995676	17.93653	-4.410175	12.3832925	3.316866e-01
visit3:group	27.571424	12.419868	17.83119	1.460532	53.6823152	3.963353e-02
visit4:group	30.223906	11.320769	17.99537	6.439413	54.0083977	1.562388e-02

Values for the all correlation parameters can be displayed too, by specifying `effect=c("variance","correlation")`

```
model.tables(eUN.lmm, effect = c("variance","correlation"))
```

	estimate	se	df	lower	upper	p.value
sigma	14.472118325	2.41202002	15.315761	10.15147605	20.6317000	NA
k.-1	0.895920602	0.12703190	20.267083	0.66669608	1.2039575	0.4472196321
k.1	1.889909495	0.43109758	25.915694	1.18243799	3.0206725	0.0097415223
k.13	1.352197903	0.31755015	29.807376	0.83693748	2.1846783	0.2087440682
rho(-13,-1)	0.800721445	0.08517658	13.414224	0.52948759	0.9234265	0.0004292266
rho(-13,1)	0.268804346	0.21919999	7.928565	-0.26374140	0.6757630	0.2773574833
rho(-13,13)	-0.085457774	0.23398096	8.588214	-0.55306416	0.4230915	0.7250514470
rho(-1,1)	0.003683836	0.23723671	8.148741	-0.49424281	0.4997905	0.9879844454
rho(-1,13)	-0.094132798	0.23364886	8.919109	-0.55696763	0.4133142	0.6982138069
rho(1,13)	0.538423876	0.17622066	10.223258	0.05058039	0.8188256	0.0352264245

Because these parameters are constrained (e.g. strictly positive), they uncertainty is by default computed after transformation (e.g. `log`) and then backtransformed. The column argument can be used to extract more or less information, e.g.:

```
model.tables(eUN.lmm, columns = c("estimate","p.value"))
```

	estimate	p.value
(Intercept)	38.728975	1.089106e-07
visit2	-4.734333	1.057398e-01
visit3	31.433029	1.922898e-03
visit4	4.521381	5.791699e-01
group	-12.824616	6.301532e-02
visit2:group	3.986559	3.316866e-01
visit3:group	27.571424	3.963353e-02
visit4:group	30.223906	1.562388e-02

²it is equivalent to `confint` method except that by default it also outputs `se` and `p.value`

All parameters can be displayed by specifying `effect="all"`. The functions `add` (resp. `remove`) can be used to add (resp. remove) one or several columns from the default display, e.g.:

```
model.tables(eUN.lmm, columns = add("statistic"))
```

	estimate	se	statistic	df	lower	upper	p.value
(Intercept)	38.728975	4.576486	8.4626016	18.00314	29.114255	48.3436939	1.089106e-07
visit2	-4.734333	2.775887	-1.7055210	17.54267	-10.577172	1.1085054	1.057398e-01
visit3	31.433029	8.629711	3.6424197	17.58484	13.271949	49.5941098	1.922898e-03
visit4	4.521381	8.004993	0.5648201	17.99506	-12.296816	21.3395773	5.791699e-01
group	-12.824616	6.472128	-1.9815145	18.00316	-26.421881	0.7726496	6.301532e-02
visit2:group	3.986559	3.995676	0.9977182	17.93653	-4.410175	12.3832925	3.316866e-01
visit3:group	27.571424	12.419868	2.2199451	17.83119	1.460532	53.6823152	3.963353e-02
visit4:group	30.223906	11.320769	2.6697749	17.99537	6.439413	54.0083977	1.562388e-02

4.7 Extract estimated residual variance-covariance structure

The method `sigma` can be used to output the modeled residual covariance structure:

```
Sigma <- sigma(eUN.lmm)
Sigma
```

	-13	-1	1	13
-13	209.44221	150.250246	106.399969	-24.20227
-1	150.25025	168.113758	1.306397	-23.88443
1	106.39997	1.306397	748.076864	288.18395
13	-24.20227	-23.884434	288.183947	382.95234

and then converted to a correlation matrix using `cov2cor`:

```
cov2cor(Sigma)
```

	-13	-1	1	13
-13	1.00000000	0.800721445	0.268804346	-0.08545777
-1	0.80072145	1.000000000	0.003683836	-0.09413280
1	0.26880435	0.003683836	1.000000000	0.53842388
13	-0.08545777	-0.094132798	0.538423876	1.00000000

The method can also be used to extract the residual covariance relative to a "known" individual:

```
sigma(eUN.lmm, cluster = 5)
```

	-13	1	13
-13	209.44221	106.4000	-24.20227
1	106.39997	748.0769	288.18395
13	-24.20227	288.1839	382.95234

or for a new individual:

```
newdata <- data.frame(id = "X", time = c("-13", "-1", "1", "13"))
sigma(eUN.lmm, cluster = newdata)
```

```
      -13      -1      1      13
-13 209.44221 150.250246 106.399969 -24.20227
-1  150.25025 168.113758  1.306397 -23.88443
1   106.39997  1.306397 748.076864 288.18395
13  -24.20227 -23.884434 288.183947 382.95234
```

4.8 Random effects

Mixed model having a compound symmetry structure with positive correlation parameters may be equivalent to random intercept models, possibly with nested random effects. Indeed in some case the residual variance-covariance matrix can then be decomposed as:

$$\Omega = Z\Psi Z^T + \Delta$$

where:

- Z is the design matrix associated to the random effect (e.g. patient id)
- Ψ is the variance-covariance of the random effects
- Δ the residual variance covariance conditional to the random effects.

One can use the `lme4` syntax to fit random intercept models with `lmm`:

```
eRI.lmm <- lmm(glucagonAUC ~ visit*group + (1|id), data = gastricbypassL)
eRI.lmm
```

Linear Mixed Model with a random intercept

```
outcome/cluster/time: glucagonAUC/id/XXtimeXX
data                  : 78 observations from 20 clusters
parameter            : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group visit4:group)
                      1 variance (sigma)
                      1 correlation (rho(id))
log-restr.likelihood: -314.394203759159
convergence           : TRUE (6 iterations)
```

It is also possible to specify cross or nested random effects, e.g.:

```
eNRI.lmm <- lmm(glucagonAUC ~ visit*group + (1|id/baseline), data = gastricbypassL)
eNRI.lmm
```


Linear Mixed Model with nested random intercepts

```
outcome/cluster/time: glucagonAUC/id/XXtimeXX
data                  : 78 observations from 20 clusters
parameter             : 8 mean ((Intercept) visit2 visit3 visit4 group visit2:group visit3:group visi
                        1 variance (sigma)
                        2 correlation (rho(id/baseline) rho(id))
log-restr.likelihood: -308.994835006264
convergence           : TRUE (6 iterations)
```

We obtain the same log-likelihood as, respectively, `eCS.lmm` and `eBCS.lmm`. Indeed, as previously mentioned, with positive residual correlation the random effect structure is equivalent to a compound symmetry structure.

⚠ random slopes are not currently supported in LMMstar.

⚠ the proposed implementation can be very inefficient compared to `lme4`.

The joint distribution between the outcome \mathbf{Y} and the random effects $\boldsymbol{\eta}$ can be explicit as:

$$\begin{bmatrix} \mathbf{Y} \\ \boldsymbol{\eta} \end{bmatrix} \sim \mathcal{N} \left(\begin{bmatrix} \boldsymbol{\mu} \\ \mathbf{0} \end{bmatrix}, \begin{bmatrix} \Omega & Z\Psi \\ \Psi Z^\top & \Psi \end{bmatrix} \right)$$

Denote by $\varepsilon_i = \mathbf{Y}_i - \boldsymbol{\mu}_i$ the vector of marginal residuals relative to individual i , Ω_i its variance-covariance matrix, and $\psi_j = (\Psi)_{jj}$ the variance of the j -th random effect. We can re-express the expected value of the j -th random effect for individual i as:

$$\eta_{ij} = \psi_j Z_{ij} \Omega_i^{-1} \varepsilon_i$$

This is what the `ranef` method returns:

```
head(ranef(eRI.lmm, format = "wide"))
```

```
head(ranef(eNRI.lmm, format = "wide"))
```

```
   id  estimate
1  1  0.9036038
2  2 32.5542378
3  3 -18.3099658
4  4 20.2561307
5  5 -15.4258816
6  6 19.3751847
```

```
   id  estimate estimate.0 estimate.1
1  1  4.931442  0.52901983 -0.4829138
2  2 28.390660 -0.09204109  0.3574766
3  3 -13.728389  0.18951039 -0.3178625
4  4 15.645550  0.82309894 -0.6768225
5  5 -11.246852 -0.30658155  0.2014303
6  6 15.002108 -2.64303027  2.7832909
```

It is also possible to extract the variance decomposition by setting the argument `effects` to `"variance"`:

```
ranef(eRI.lmm, effects = "variance", format = "wide", simplify = FALSE)
```

```
variable strata  variance  relative
1   total      1 355.30623 1.00000000
2     id       1 344.62363 0.96993408
3 residual     1  10.68261 0.03006592
```

```
ranef(eNRI.lmm, effects = "variance", format = "wide", simplify = FALSE)
```

	variable	strata	variance	relative
1	total	1	346.744085	1.000000000
2	id	1	336.182537	0.969540799
3	baseline	1	3.143104	0.009064622
4	residual	1	7.418444	0.021394579

4.9 Sum of squares

⚠ The definition of the sum of squares is not straightforward with mixed models. Intuitively summing residuals across several outcomes will be hard to interpret unless all outcomes have the same variance. This is why LMMstar does not provide them. Nevertheless for specific covariance structure, namely independence and compound symmetry (with positive correlation) structure, sum of squares can be deduced from the `lmm` object - see appendix C for the theoretical derivations. Importantly, with these structures the residuals can be reparametrised as random effects plus independent residuals, i.e. $\Omega = Z\Psi Z^\top + \delta I$ where I is the identity matrix and δ the variance of these independent residuals.

Appendix C illustrate how to extract the sum of squares for univariate linear regression (i.e. independence structure) and here we illustrate the case of a compound symmetry structure. A key step is to extract from the `lmm` object the conditional variance δ :

```
sigma2 <- coef(eCS.lmm, effect = "variance")^2
tau <- coef(eCS.lmm, effect = "correlation")*sigma2
delta <- unname(sigma2 - tau)
```

This step will typically depend on the covariance structure. The residual sum of squares (SSE) equals the residual degrees of freedom times the conditional variance:

```
df.res <- df.residual(eCS.lmm)
SSE <- df.res * delta
c(df.res = df.res, SSE = SSE)
```

```
df.res      SSE
73.0000 779.8304
```

For the regression sum of squares (SSR), we first extract the mean parameters and their variance-covariance based on the expected information:

```
eBeta.lmm <- coef(eCS.lmm)
eVcov.lmm <- vcov(eCS.lmm, type.information = "expected")
```

Parameters are grouped with respect to the original variable:

```
attr(model.matrix(eCS.lmm), "assign")
```

```
[1] 0 1 1 1 2
```

So we respect this grouping when computing the normalized SSR:

```
SSRstar.time <- eBeta.lmm[2:4] %*% solve(eVcov.lmm[2:4,2:4]) %*% eBeta.lmm[2:4]
SSRstar.glucagon <- eBeta.lmm[5] %*% solve(eVcov.lmm[5,5]) %*% eBeta.lmm[5]
```

The SSR is obtained by multiplying the normalized SSR by the conditional variance:

```
SSR.time <- as.double(SSRstar.time * delta)
SSR.glucagon <- as.double(SSRstar.glucagon * delta)
c(time = SSR.time, glucagon = SSR.glucagon)
```

```
      time    glucagon
6986.78351  18.83074
```

4.10 Proportion of explained variance and partial correlation

⚠ The definition of explained variance is not straightforward with mixed models. Intuitively considering the variance across several outcomes will be hard to interpret unless all outcomes have the same variance. Similar consideration holds for partial correlation. This is why LMMstar does not output these quantities by default. Nevertheless for specific covariance structure, namely independence and compound symmetry (with positive correlation) structure, explained variance and partial correlation can be deduced from the `lmm` object. Importantly, with these structures the residuals can be reparametrised as random effects plus independent residuals, i.e. $\Omega = Z\Psi Z^\top + \delta I$ where I is the identity matrix and δ the variance of these independent residuals.

The proportion of explained variance, also called partial R^2 or partial η^2 , is defined as the ratio between sum of squares (e.g. [Lakens \(2013\)](#), equation 12):

$$R^2 = \frac{SSR}{SSR + SSE}$$

```
c(SSR.time/ (SSR.time + SSE),
  SSR.glucagon/ (SSR.glucagon + SSE))
```

```
[1] 0.89959197 0.02357789
```

Computing the SSR for each individual coefficients, taking its squared root, and multiplying by the sign of the corresponding coefficient leads to the partial correlation

```
eCS.R2 <- partialCor(eCS.lmm, R2 = TRUE)
summary(eCS.R2)
```

Partial correlation

	estimate	se	df	lower	upper	p.value
timeB1w	-0.646	0.055	18.6	-0.762	-0.53	5.11e-10
timeA1w	-0.765	0.035	9.5	-0.845	-0.686	2.07e-09

```
timeA3m      -0.946 0.006  2.4 -0.969 -0.923 6.80e-06
glucagon      0.154 0.114 45.3 -0.076  0.383   0.184
-----
```


Columns lower and upper contain 95% pointwise confidence intervals for each coefficient.
Degrees of freedom were computed using a Satterthwaite approximation (column df).

Coefficient of determination (R2)

```
      estimate    se    df  lower upper  p.value
time          0.9 0.011   2.4  0.857 0.942 4.09e-05
glucagon      0.024 0.035 45.3 -0.047 0.094   0.503
global        0.906 0.011   2.3  0.866 0.946 4.51e-05
-----
```

Columns lower and upper contain 95% pointwise confidence intervals for each coefficient.
Degrees of freedom were computed using a Satterthwaite approximation (column df).

Here the line "global" refer to the R2 for all covariates, computed based on the SSR relative to all mean parameters but the intercept.

 **partialCor** will compute values for all types of mixed models. But their interpretation as partial correlation and proportion of explained variance outside the covariance structures mentioned in this section is questionable.

Note: Other software packages like `effectsize::eta_squared` uses another formula to estimate the partial R2:

$$R^2 = \frac{F df_{num}}{F df_{num} + df_{denom}}$$

where F denote the F-statistic, df_{num} (resp. df_{denom}) the degrees of freedom of the numerator (resp. denominator) of this statistic. However since the calculation of degrees of freedom in LMM is approximate, I would expect this approach to be less reliable than the one of **partialCor** based on the SSR and SSE.

```
aCS.aov <- anova(eCS.lmm)$multivariate
setNames(with(aCS.aov, statistic*df.num/(statistic*df.num+df.denom)), aCS.aov$test)
```

```
      time  glucagon
0.92380363 0.03162017
```

4.11 Model diagnostic

The method `residuals` returns the residuals in the wide format:

```
eUN.diagW <- residuals(eUN.lmm, type = "normalized", format = "wide")
colnames(eUN.diagW) <- gsub("normalized.", "", colnames(eUN.diagW))
head(eUN.diagW)
```

	id	r.B3m	r.B1w	r.A1w	r.A3m
1	1	-0.1082872	0.4283943	0.7477306	0.91794015
2	2	1.8182348	-0.3516996	1.5698307	-0.98743171
3	3	-0.9318737	-0.7728221	0.6315751	0.16549699
4	4	0.8408969	1.8695564	0.3485784	-0.09662565
5	5	-0.7882340	NA	-0.6128276	0.09933842
6	6	1.4896141	-1.9727358	-1.9672939	-1.37068983

or in the long format:

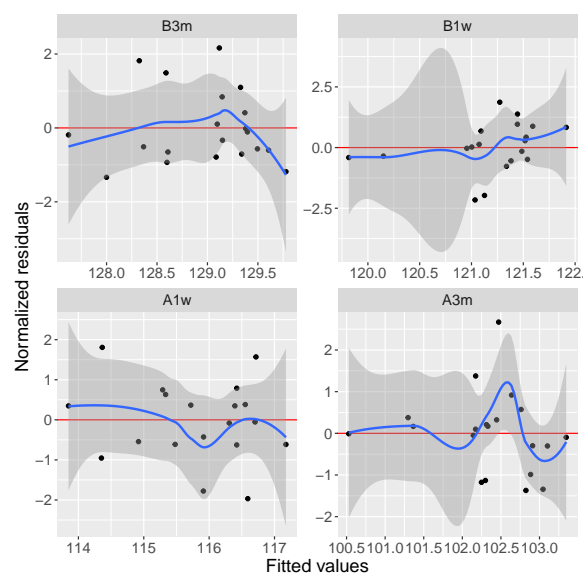
```
eUN.diagL <- residuals(eUN.lmm, type = "normalized", format = "long", keep.data = TRUE)
head(eUN.diagL)
```

	id	visit	time	weight	glucagonAUC	baseline	glucagon	group	fitted	r.normalized
1	1	1	B3m	127.2	5032.50	TRUE	4.034616	1	129.3962	-0.1082872
2	2	1	B3m	165.2	12142.50	TRUE	5.240766	0	128.3247	1.8182348
3	3	1	B3m	109.7	10321.35	TRUE	4.931824	1	128.5992	-0.9318737
4	4	1	B3m	146.2	6693.00	TRUE	4.316306	0	129.1459	0.8408969
5	5	1	B3m	113.1	7090.50	TRUE	4.383738	1	129.0860	-0.7882340
6	6	1	B3m	158.8	10386.00	TRUE	4.942791	0	128.5894	1.4896141

Various type of residuals can be extract but the normalized one are recommended when doing model checking. Diagnostic plots can then be generated by the user, or directly from the `lmm` object via the method `plot` (which internally calls the `residuals` method):

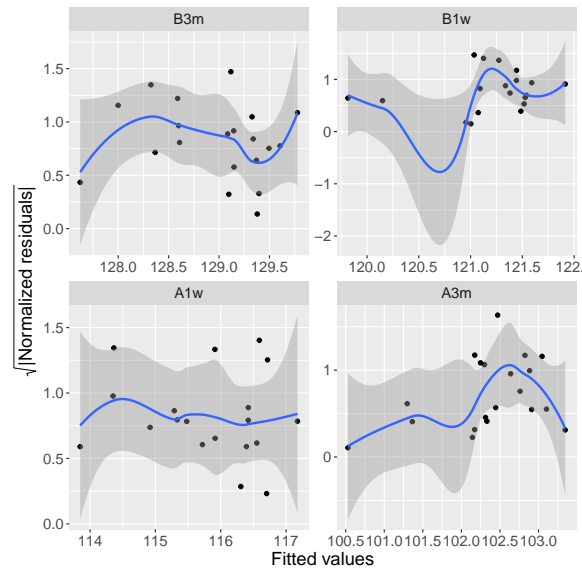
- misspecification of the mean structure

```
plot(eUN.lmm, type = "scatterplot")
```



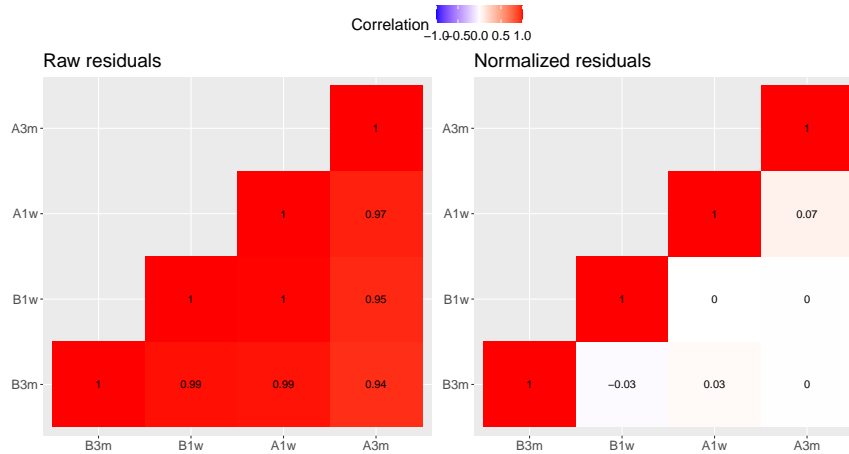
- misspecification of the variance structure

```
plot(eUN.lmm, type = "scatterplot2")
```



- misspecification of the correlation structure

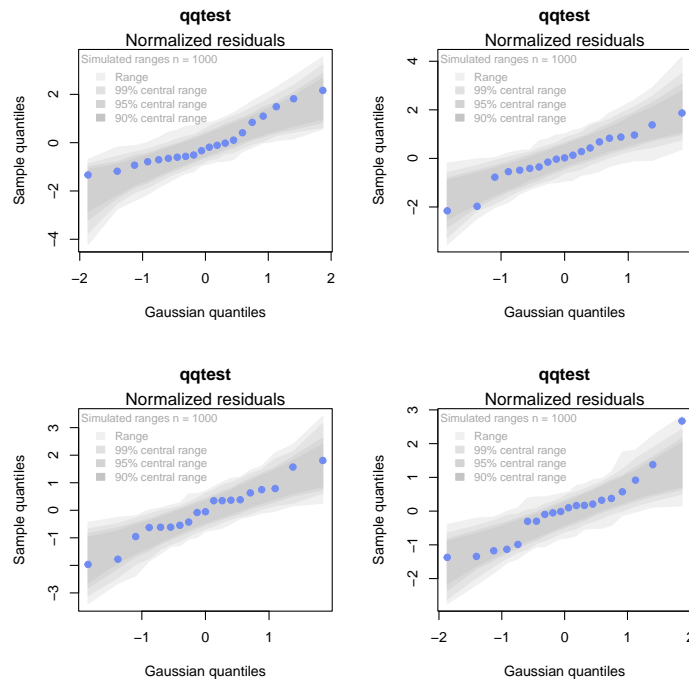
```
plot(eUN.lmm, type = "correlation", type.residual = "response")
plot(eUN.lmm, type = "correlation", type.residual = "normalized")
```



- residual distribution vs. normal distribution ³:

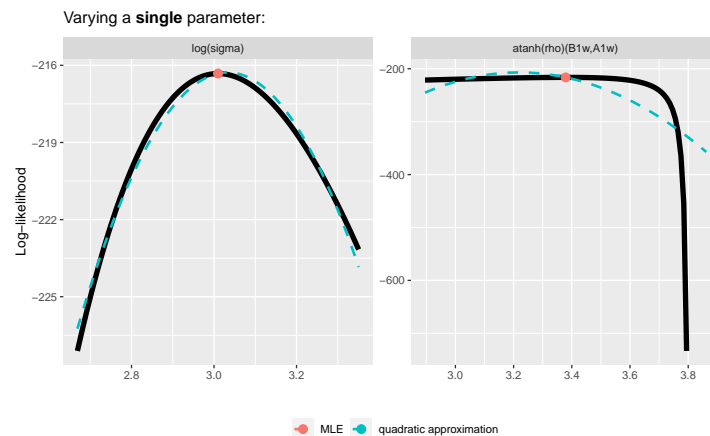
```
plot(eUN.lmm, type = "qqplot", engine.qqplot = "qqtest")
## Note: the qqtest package to be installed to use the argument engine.plot = "qqtest"
```

³see Oldford (2016) for guidance about how to read quantile-quantile plots.



⚠ Deviation from the normal distribution does not necessarily question the validity of the statistical inference. Moreover, for variance and correlation parameters, normally distributed data is not enough to ensure valid statistical inference. Instead one could assess whether the log-likelihood is locally quadratic as this ensures normally distributed estimates in finite samples (Geyer, 2013). Since the likelihood function is a multi-dimensional function this is not an easy task but one can look at specific 'slices' using the `profile` method:

```
plot(profile(eUN.lmm, effects = c("sigma", "rho(B1w,A1w)")))
```

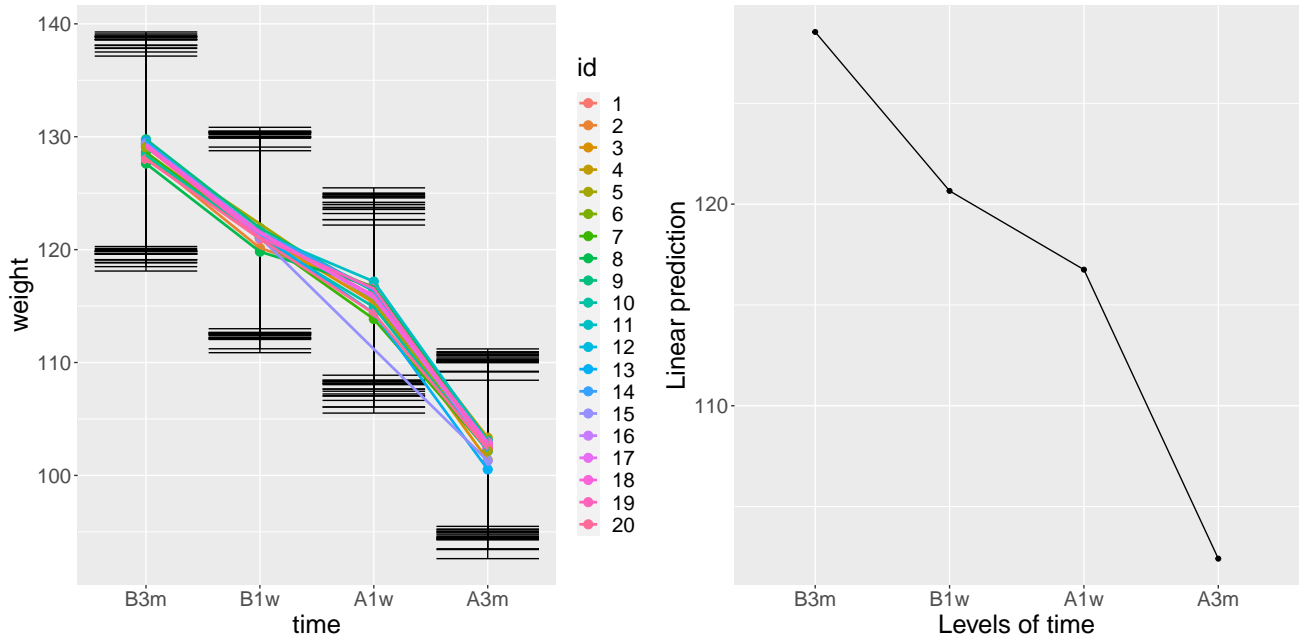


4.12 Model fit

The fitted values can be displayed via the `plot` method or using the `emmeans` package:

```
plot(eUN.lmm, type = "fit", color = "id", ci.alpha = NA, size.text = 20)
```

```
library(emmeans) ## right panel  
emmip(eUN.lmm, ~time) + theme(text = element_text(size=20))
```

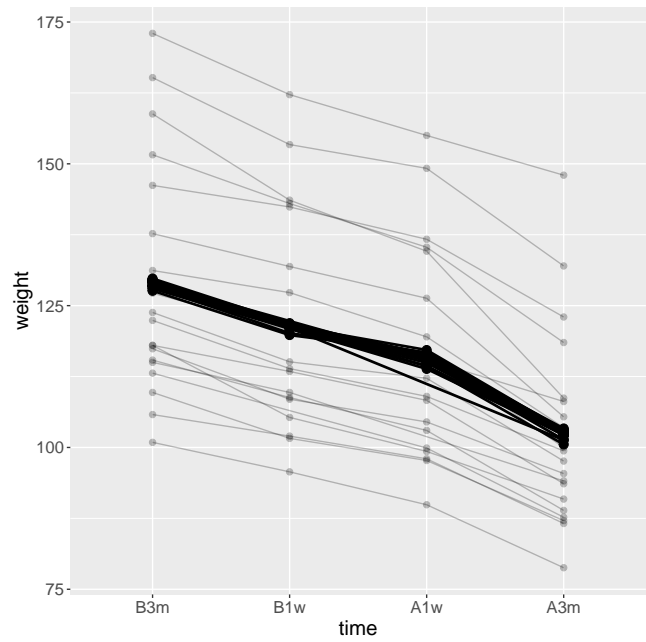
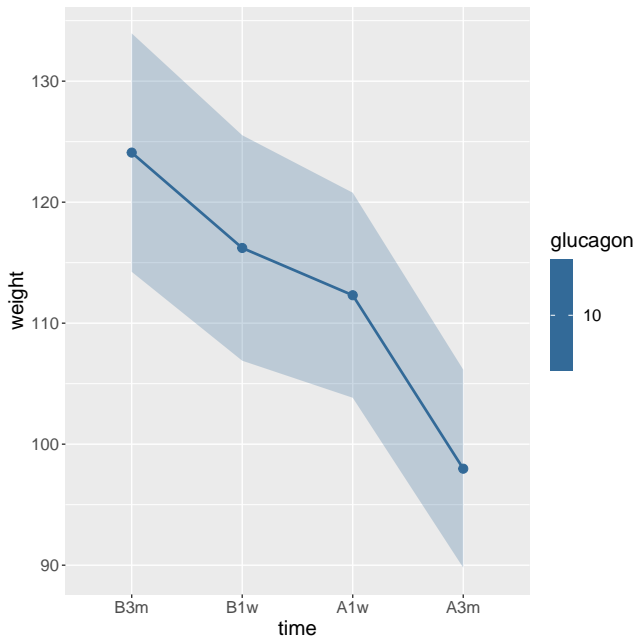


In the first case each possible curve is displayed while in the latter the average curve (over glucagon values). With the `plot` method, it is possible to display a curve specific to a glucagon value via the argument `at`:

```
## left panel  
plot(eUN.lmm, type = "fit", at = data.frame(glucagon = 10), color = "glucagon")
```

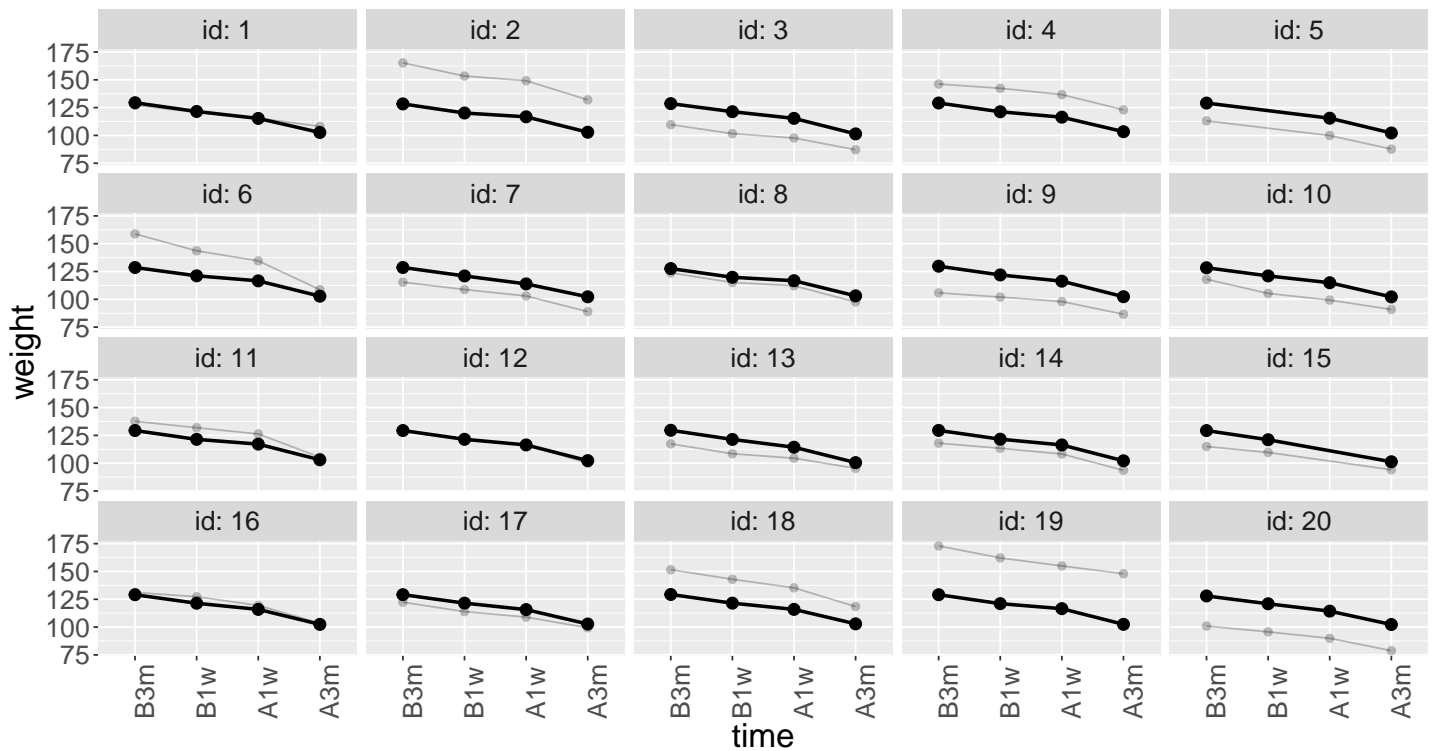
It is also possible to display the observed values along with the fitted values by setting the argument `obs.alpha` to a strictly positive value below or equal to 1. This argument controls the transparency of the color used to display the observed values:

```
## right panel  
gg.spafit <- plot(eUN.lmm, type = "fit", obs.alpha = 0.25, ci = FALSE)$plot
```



The `plot` element output by the `plot` method can be manipulated as a `ggplot` object to modify the visual appearance, e.g. to compare each individual trajectory to the model fit:

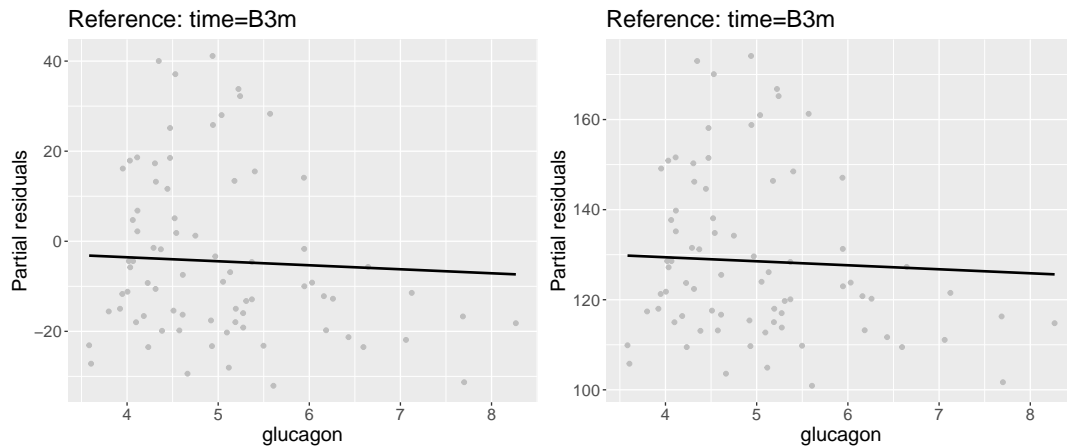
```
gg.traj <- gg.spafit + facet_wrap(~id, labeller = label_both)
gg.traj <- gg.traj + theme(axis.text.x=element_text(angle = 90, hjust = 0))
gg.traj
```



4.13 Partial residuals

Partial residuals can also be displayed via the `plot` method:

```
gg1 <- plot(eUN.lmm, type = "partial", var = "glucagon")$plot
gg2 <- plot(eUN.lmm, type = "partial", var = c("(Intercept)", "glucagon"))$plot
ggarrange(gg1, gg2)
```



Their value can be extracted via the `residuals` method, e.g.:

```
df.pres <- residuals(eUN.lmm, type = "partial", var = "glucagon", keep.data = TRUE)
head(df.pres)
```

	id	visit	time	weight	glucagonAUC	baseline	glucagon	group	fitted	r.partial
1	1	1	B3m	127.2	5032.50	TRUE	4.034616	1	-3.583982	-5.780135
2	2	1	B3m	165.2	12142.50	TRUE	5.240766	0	-4.655415	32.219865
3	3	1	B3m	109.7	10321.35	TRUE	4.931824	1	-4.380979	-23.280135
4	4	1	B3m	146.2	6693.00	TRUE	4.316306	0	-3.834209	13.219865
5	5	1	B3m	113.1	7090.50	TRUE	4.383738	1	-3.894110	-19.880135
6	6	1	B3m	158.8	10386.00	TRUE	4.942791	0	-4.390721	25.819865

This matches manual calculation:

```
m.pres <- dfL$weight - model.matrix(~time, dfL) %*% coef(eUN.lmm)[1:4]
range(df.pres$r.partial - m.pres, na.rm = TRUE)

m.pfit <- model.matrix(~0+glucagon, dfL) %*% coef(eUN.lmm)["glucagon"]
range(df.pres$fitted - m.pfit, na.rm = TRUE)
```

```
[1] -1.065814e-14  1.421085e-14
[1] 0 0
```

The `plot` methods can handle one continuous and one categorical covariate (in addition to the intercept) to display interaction plots. In that case each observation/fitted line is colored according to the categorical covariate.

4.14 Statistical inference (linear)

The `anova` method can be used to test one or several linear combinations of the model coefficients using Wald tests. By default, it will simultaneously test all parameters associated to a variable:

```
anova(eUN.lmm)
```

Multivariate Wald test

	F-statistic	df	p.value
mean: time	86.743 (3,19.0)	2.84e-11	***
: glucagon	13.518 (1,13.7)	0.00257	**

Note that here the p-values are not adjusted for multiple comparisons over variables. It is possible to specify a null hypothesis to be tested: e.g. is there a change in average weight just after taking the treatment:

```
anova(eUN.lmm, effects = c("timeA1w-timeB1w=0"))
```

Multivariate Wald test

	F-statistic	df	p.value
all: 1	43.141 (1,17.9)	3.72e-06	***

One can also simultaneously test several null hypotheses:

```
e.anova <- anova(eUN.lmm, effects = c("timeA1w-timeB1w=0", "timeA3m-timeB1w=0"))
summary(e.anova)
```

Multivariate Wald test

	F-statistic	df	p.value
all: 1	98.651 (2,18.6)	1.23e-10	***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Degrees of freedom were computed using a Satterthwaite approximation (column df).

Univariate Wald test

	estimate	se	df	lower	upper	p.value
timeA1w - timeB1w	-3.906	0.595	17.9	-5.325	-2.487	3e-05 ***
timeA3m - timeB1w	-18.24	1.323	19	-21.397	-15.083	<1e-05 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Columns lower/upper/p.value adjusted for multiple comparisons -- max-test.

(1e+05 samples have been used)

Model-based standard errors are derived from the observed information (column se).

Degrees of freedom were computed using a Satterthwaite approximation (column df).

or return all pairwise comparisons for a given factor using the `mcp` function of the `multcomp` package:

```
library(multcomp)
summary(anova(eUN.lmm, effects = mcp(time = "Tukey")))
```

Singular contrast matrix: contrasts "A1w - B1w" "A3m - B1w" "A3m - A1w" have been removed.

Multivariate Wald test

	F-statistic	df	p.value
all: time	86.743	(3,19.0)	2.84e-11 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Degrees of freedom were computed using a Satterthwaite approximation (column df).

Univariate Wald test

	estimate	se	df	lower	upper	p.value
B1w - B3m	-7.882	0.713	19.2	-9.817	-5.947	<1e-05 ***
A1w - B3m	-11.788	1.018	21.6	-14.549	-9.027	<1e-05 ***
A3m - B3m	-26.122	1.656	18.8	-30.617	-21.628	<1e-05 ***
A1w - B1w	-3.906	0.595	17.9	-5.519	-2.292	<1e-05 ***
A3m - B1w	-18.24	1.323	19	-21.829	-14.651	<1e-05 ***
A3m - A1w	-14.334	1.057	20.3	-17.201	-11.468	<1e-05 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Columns lower/upper/p.value adjusted for multiple comparisons -- max-test.

(1e+05 samples have been used)

Model-based standard errors are derived from the observed information (column se).

Degrees of freedom were computed using a Satterthwaite approximation (column df).

Here the `summary` method prints not only the global test but also the result associated to each hypothesis. When testing transformed variance or correlation parameters, parentheses (as in `log(k).B1w`) cause problem for recognizing parameters:

```
try(
  anova(eUN.lmm,
    effects = c("log(k).B1w=0", "log(k).A1w=0", "log(k).A3m=0"))
)
```

Error in `.anova_Wald(object, effects = effects, robust = robust, rhs = rhs, :`

Possible misspecification of the argument 'effects' as running `mulcomp::glht` lead to the following

Error in `parse(text = ex[i])` : <text>:1:7: uventet symbol

1: log(k).B1w

^

It is then advised to build a contrast matrix, e.g.:

```
name.coef <- rownames(confint(eUN.lmm, effects = "all"))
name.varcoef <- grep("^k",name.coef, value = TRUE)
C <- matrix(0, nrow = 3, ncol = length(name.coef), dimnames = list(name.varcoef, name.coef))
diag(C[name.varcoef,name.varcoef]) <- 1
C[,1:9]
```

```
      (Intercept) timeB1w timeA1w timeA3m glucagon sigma k.B1w k.A1w k.A3m
k.B1w           0         0         0         0         0         0         1         0         0
k.A1w           0         0         0         0         0         0         0         1         0
k.A3m           0         0         0         0         0         0         0         0         1
```

And then call the `anova` method specifying the null hypothesis via the contrast matrix:

```
anova(eUN.lmm, effects = C)
```

Multivariate Wald test

```
      F-statistic      df p.value
all: 1          6.203 (3,18.0) 0.00442 **
```

Note that using the approach of [Pipper et al. \(2012\)](#) it is also possible to adjust for multiple testing across several `lmm` objects. To do so, one first fit the mixed models, then use the `anova` method to indicate which hypotheses are being tested, and combine them using `rbind`. Here is an (artificial) example:

```
Manova <- rbind(anova(eInd.lmm, effects = "glucagon = 0", robust = FALSE),
               anova(eCS.lmm, effects = "glucagon = 0", robust = FALSE),
               anova(eUN.lmm, effects = "glucagon = 0", robust = FALSE),
               name = c("Ind", "CS", "UN"))
summary(Manova)
```

Multivariate Wald test

```
      Chi2-statistic      df p.value
all: 1          8.893 (3,Inf) 6.88e-06 ***
```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Univariate Wald test

```
      estimate      se      df      lower      upper      p.value
Ind: glucagon    -8.27  2.579  34.2  -14.414   -2.126  0.003988 **
CS: glucagon      0.822  0.62  53.8   -0.655    2.299  0.450012
UN: glucagon     -0.888  0.242  13.7   -1.464   -0.313  0.000711 ***
```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1.

Columns lower/upper/p.value adjusted for multiple comparisons -- max-test.

(error when computing the adjusted columns lower/upper/p.value by numerical integration: 1.52e-05)

Model-based standard errors are derived from the observed information (column se).

4.15 Statistical inference (non-linear)

The `estimate` function can be used to test one or several non-linear combinations of model coefficients, using a first order delta method to quantify uncertainty. The combination has to be specified via a function (argument `f`). To illustrate its use consider an ANCOVA analysis:

$$Y_{i1} = \alpha + \beta Y_{i,0} + \gamma X_i + e_i$$

```
e.ANCOVA <- lm(weight4 ~ weight1 + group, data = gastricbypassW)
summary(e.ANCOVA)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-5.9285136	8.78006389	-0.6752244	5.086140e-01
weight1	0.8236279	0.06411563	12.8459772	3.524665e-10
group	4.1404554	2.53335466	1.6343765	1.205604e-01

We can replicate this analysis by first fitting a mixed model:

$$Y_{ij} = \alpha_j + \gamma_j X_i + \varepsilon_{i,j} \text{ where } \varepsilon_i \sim \mathcal{N} \left(\begin{bmatrix} 0 \\ 0 \end{bmatrix}, \begin{bmatrix} \sigma_1^2 & \rho\sigma_1\sigma_2 \\ \rho\sigma_1\sigma_2 & \sigma_2^2 \end{bmatrix} \right)$$

```
dfL14 <- dfL[dfL$visit %in% c(1,4),]
dfL14$time <- droplevels(dfL14$time)
e.lmmANCOVA <- lmm(weight ~ time+time:group, repetition = ~time|id,
  data = dfL14)
```

and then perform a first order delta-method:

```
lava::estimate(e.lmmANCOVA, f = function(p){
  c(Y1 = as.double(p["rho(B3m,A3m)"]*p["k.A3m"]),
    X1 = as.double(p["timeA3m:group"]-p["rho(B3m,A3m)"]*p["k.A3m"]*p["timeB3m:group"]))
})
```

	estimate	se	df	lower	upper	p.value
Y1	0.8236279	0.06230919	9.874633	0.6845551	0.9627007	1.332743e-07
X1	4.1404554	2.46197819	15.161269	-1.1022695	9.3831803	1.130927e-01

Indeed:

$$\begin{aligned} \mathbb{E}[Y_{i2}|Y_{i1}, X_i] &= \alpha_2 + \gamma_2 X_i + \rho \frac{\sigma_2}{\sigma_1} (Y_{i1} - \alpha_1 - \gamma_1 X_i) \\ &= \alpha_2 - \rho \frac{\sigma_2}{\sigma_1} \alpha_1 + \rho \frac{\sigma_2}{\sigma_1} Y_{i1} + \left(\gamma_2 - \rho \frac{\sigma_2}{\sigma_1} \gamma_1 \right) X_i \end{aligned}$$

We obtain identical estimate but different standard-errors/degrees of freedom compared to the univariate linear model approach. The later is to be prefer as it does not rely on approximation. The former is nevertheless useful as it can handle missing data in the outcome variable.

4.16 Baseline adjustment

In clinical trial the group and intervention variable often do not coincide, e.g., in presence of baseline measurement. In our running example, the first two measurement are pre-treatment (i.e. treatment should be "none") while the last two measurements are post-treatment (i.e. treatment should be 1 or 2). The `baselineAdjustment` function can be helpful to:

- define the treatment variable from the time and allocation variable, where baseline has its specific value

```
gastricbypassL$treat <- baselineAdjustment(gastricbypassL, variable = "group",
                                           repetition = ~time|id, constrain = c("B3m","B1w"),
                                           new.level = "none")
table(treat = gastricbypassL$treat, time = gastricbypassL$time, group = gastricbypassL$group)
```

```
, , group = 0
```

```
      time
treat  B3m B1w A1w A3m
  none   10  10   0   0
    0     0   0  10  10
    1     0   0   0   0
```

```
, , group = 1
```

```
      time
treat  B3m B1w A1w A3m
  none   10  10   0   0
    0     0   0   0   0
    1     0   0  10  10
```

- define the treatment variable from the time and allocation variable, where baseline corresponds to the reference group

```
gastricbypassL$treat2 <- baselineAdjustment(gastricbypassL, variable = "group",
                                           repetition = ~time|id, constrain = c("B3m","B1w")
                                           )
table(treat = gastricbypassL$treat2, time = gastricbypassL$time, group = gastricbypassL$group)
```

```
, , group = 0
```

```
      time
treat  B3m B1w A1w A3m
    1   10  10   0   0
    0    0   0  10  10
```



```
, , group = 1
```

```
      time
treat B3m B1w A1w A3m
      1  10  10  10  10
      0   0   0   0   0
```

- define a time varying treatment variable from the time and allocation variable

```
gastricbypassL$timeXtreat <- baselineAdjustment(gastricbypassL, variable = "group",
                                                repetition = ~time|id, constrain = c("B3m", "
B1w"),
                                                collapse.time = ".")

table(treat = gastricbypassL$timeXtreat, time = gastricbypassL$time, group = gastricbypassL$
group)
```

```
, , group = 0
```

```
      time
treat   B3m B1w A1w A3m
B3m      10  0  0  0
B1w       0 10  0  0
A1w.0     0  0 10  0
A3m.0     0  0  0 10
A1w.1     0  0  0  0
A3m.1     0  0  0  0
```

```
, , group = 1
```

```
      time
treat   B3m B1w A1w A3m
B3m      10  0  0  0
B1w       0 10  0  0
A1w.0     0  0  0  0
A3m.0     0  0  0  0
A1w.1     0  0 10  0
A3m.1     0  0  0 10
```

We would then typically like to model group differences only after baseline (i.e. only at 1 week and 3 months after). This can be performed using the time varying treatment variable, e.g.:

```
eC.lmm <- lmm(weight ~ timeXtreat, data = gastricbypassL,
              repetition = ~time|id, structure = "UN")
coef(eC.lmm) ## change from baseline
```

```
(Intercept)    timeXtreatB1w timeXtreatA1w.0 timeXtreatA3m.0 timeXtreatA1w.1 timeXtreatA3m.1
128.97000      -7.73000      -13.38978      -28.52130      -13.15022      -24.68870
```

or

```
eC2.lmm <- lmm(weight ~ 0 + timeXtreat, data = gastricbypassL,
               repetition = ~time|id, structure = "UN")
coef(eC2.lmm) ## absolute value
```

```
timeXtreatB3m    timeXtreatB1w timeXtreatA1w.0 timeXtreatA3m.0 timeXtreatA1w.1 timeXtreatA3m.1
128.9700         121.2400         115.5802         100.4487         115.8198         104.2813
```

The parametrization however does not (directly) output treatment effects. Instead one may be tempted to use a formula like `treatment*time`. However this will lead to a non-indentifiable model. Indeed we are only able to estimate a total of 6 means when constraining the expected baseline value between the two groups to be the same. Therefore can at most identify 6 effects. However the design matrix for the interaction model:

```
colnames(model.matrix(weight ~ treat*time, data = gastricbypassL))
```

```
[1] "(Intercept)"      "treat0"            "treat1"            "timeB1w"           "timeA1w"
[6] "timeA3m"          "treat0:timeB1w"    "treat1:timeB1w"    "treat0:timeA1w"    "treat1:timeA1w"
[11] "treat0:timeA3m"    "treat1:timeA3m"
```

contains 12 parameters (i.e. 6 too many). Fortunately, the `lmm` will drop non-identifiable effects from the model and fit the resulting simplified model:

```
eC3.lmm <- lmm(weight ~ treat2*time, data = gastricbypassL,
               repetition = ~time|id, structure = "UN")
```

Constant values in the design matrix for the mean structure.
Coefficients "treat20" "treat20:timeB1w" relative to interactions "treat2:time" have been removed.

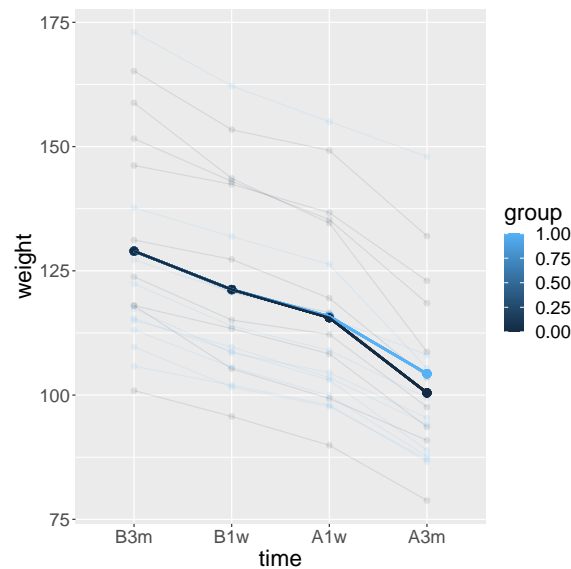
with the following coefficients:

```
model.tables(eC3.lmm)
```

	estimate	se	df	lower	upper	p.value
(Intercept)	128.9700000	4.5323695	18.98130	119.483009	138.4569912	0.000000e+00
timeB1w	-7.7300000	0.6974427	18.97552	-9.189892	-6.2701082	9.938186e-10
timeA1w	-13.1502219	0.8970429	22.87334	-15.006465	-11.2939786	4.058975e-13
timeA3m	-24.6886957	1.7751662	22.25061	-28.367762	-21.0096290	1.863398e-12
treat20:timeA1w	-0.2395562	0.6484895	17.66860	-1.603816	1.1247037	7.162149e-01
treat20:timeA3m	-3.8326086	2.1066817	17.60613	-8.265691	0.6004734	8.592047e-02

One can visualize the baseline adjustment via the `plot` function:

```
plot(eC3.lmm, color = "group", ci = FALSE, size.text = 20, obs.alpha = 0.1)
```



4.17 Marginal means

The `emmeans` package can be used to output marginal means. Consider the following model:

```
dfL$group2 <- as.numeric(dfL$id) %% 3 == 0
e.group <- lmm(glucagon ~ time*group2, data = dfL,
               repetition = ~time|id, structure = "UN")
```

We can for instance compute the average value over time *assuming balanced groups*:

```
emmeans(e.group, specs=~time)
```

NOTE: Results may be misleading due to involvement in interactions

	time	emmean	SE	df	lower.CL	upper.CL
B3m	4.45	0.156	18.0	4.12	4.78	
B1w	4.32	0.131	18.0	4.05	4.60	
A1w	5.95	0.262	18.4	5.40	6.50	
A3m	5.12	0.187	18.0	4.73	5.51	

Results are averaged over the levels of: group2

Confidence level used: 0.95

This differs from the average value over time over the whole sample:

```
df.pred <- predict(e.group, newdata = dfL, keep.newdata = TRUE)
summarize(formula = estimate~time, data = df.pred)
```

	time	observed	missing	mean	sd	min	q1	median	q3	max
1	B3m	20	0	4.514352	0.1502565	4.290643	4.290643	4.610227	4.610227	4.610227
2	B1w	19	0	4.384638	0.1643256	4.149209	4.149209	4.493298	4.493298	4.493298
3	A1w	19	0	6.060587	0.2030012	5.729961	5.954314	6.178668	6.178668	6.178668
4	A3m	20	0	5.057642	0.1465315	4.964144	4.964144	4.964144	5.275805	5.275805

as the groups are not balanced:

```
table(group = dfL$group2, time = dfL$time)
```

```
      time
group  B3m B1w A1w A3m
FALSE  14  13  14  14
TRUE    6   6   5   6
```

The "emmeans" approach gives equal "weight" to the expected value of both group:

```
mu.group1 <- as.double(coef(e.group)["(Intercept)"])
mu.group2 <- as.double(coef(e.group)["(Intercept)"] + coef(e.group)["group2TRUE"])
p.group1 <- 14/20 ; p.group2 <- 6/20
c(emmeans = (mu.group1+mu.group2)/2, predict = mu.group1 * p.group1 + mu.group2 * p.group2)
```

```
emmeans predict
4.450435 4.514352
```

Which one is relevant depends on the application. The `emmeans` function can also be used to display expected value in each group over time:

```
emmeans.group <- emmeans(e.group, specs = ~group2|time)
emmeans.group
```

```
time = B3m:
group2 emmean    SE    df lower.CL upper.CL
FALSE   4.61 0.171 18.0    4.25    4.97
TRUE    4.29 0.262 18.0    3.74    4.84
```

```
time = B1w:
group2 emmean    SE    df lower.CL upper.CL
FALSE   4.49 0.145 18.4    4.19    4.80
TRUE    4.15 0.219 17.9    3.69    4.61
```

```
time = A1w:
group2 emmean    SE    df lower.CL upper.CL
FALSE   6.18 0.277 17.8    5.60    6.76
TRUE    5.73 0.446 18.6    4.80    6.66
```

```
time = A3m:
group2 emmean    SE    df lower.CL upper.CL
FALSE   4.96 0.205 18.0    4.53    5.39
TRUE    5.28 0.313 18.0    4.62    5.93
```

Confidence level used: 0.95

Using the `pair` function displays the differences:

```
epairs.group <- pairs(emmeans.group, reverse = TRUE)
epairs.group
```

```
time = B3m:
  contrast      estimate    SE   df t.ratio p.value
TRUE - FALSE   -0.320 0.313 18.0  -1.022  0.3202
```

```
time = B1w:
  contrast      estimate    SE   df t.ratio p.value
TRUE - FALSE   -0.344 0.262 18.0  -1.311  0.2062
```

```
time = A1w:
  contrast      estimate    SE   df t.ratio p.value
TRUE - FALSE   -0.449 0.525 18.4  -0.855  0.4034
```

```
time = A3m:
  contrast      estimate    SE   df t.ratio p.value
TRUE - FALSE    0.312 0.374 18.0   0.834  0.4153
```

One can adjust for multiple comparison via the `adjust` argument and display confidence intervals setting the argument `infer` to `TRUE`:

```
summary(epairs.group, by = NULL, adjust = "mvt", infer = TRUE)
```

contrast	time	estimate	SE	df	lower.CL	upper.CL	t.ratio	p.value
TRUE - FALSE	B3m	-0.320	0.313	18.0	-1.156	0.517	-1.022	0.6926
TRUE - FALSE	B1w	-0.344	0.262	18.0	-1.046	0.358	-1.311	0.5061
TRUE - FALSE	A1w	-0.449	0.525	18.4	-1.853	0.955	-0.855	0.7960
TRUE - FALSE	A3m	0.312	0.374	18.0	-0.689	1.312	0.834	0.8084

Confidence level used: 0.95

Conf-level adjustment: mvt method for 4 estimates

P value adjustment: mvt method for 4 tests

This should also work when doing baseline adjustment (because of baseline adjustment no difference is expected at the first two timepoints):

```
summary(pairs(emmeans(eC3.lmm , specs = ~treat2|time), reverse = TRUE), by = NULL)
```

Note: `adjust = "tukey"` was changed to `"sidak"`

because `"tukey"` is only appropriate for one set of pairwise comparisons

contrast	time	estimate	SE	df	t.ratio	p.value
treat20 - treat21	B3m	0.00	0.000	Inf	NaN	NaN
treat20 - treat21	B1w	0.00	0.000	Inf	NaN	NaN
treat20 - treat21	A1w	-0.24	0.648	18	-0.369	0.9195
treat20 - treat21	A3m	-3.83	2.107	18	-1.819	0.1645

P value adjustment: sidak method for 2 tests

4.18 Predictions

Two types of predictions can be performed with the `predict` method:

- **static predictions** that are only conditional on the covariates:

```
news <- dfL[dfL$id==1,]  
news$glucagon <- 0  
predict(eUN.lmm, newdata = news)
```

```
      estimate      se      df      lower      upper  
1 132.9801 4.664247 19.75815 123.24305 142.7172  
2 125.0979 4.388294 19.91418 115.94155 134.2543  
3 121.1922 4.214230 20.55331 112.41660 129.9678  
4 106.8577 3.942058 20.95499  98.65871 115.0568
```

which can be computing by creating a design matrix:

```
X.12 <- model.matrix(formula(eUN.lmm), news)  
X.12
```

```
      (Intercept) timeB1w timeA1w timeA3m glucagon  
1              1         0         0         0         0  
21             1         1         0         0         0  
41             1         0         1         0         0  
61             1         0         0         1         0  
attr(,"assign")  
[1] 0 1 1 1 2  
attr(,"contrasts")  
attr(,"contrasts")$time  
[1] "contr.treatment"
```

and then multiplying it with the regression coefficients:

```
X.12 %*% coef(eUN.lmm)
```

```
      [,1]  
1 132.9801  
21 125.0979  
41 121.1922  
61 106.8577
```

- **dynamic predictions** that are conditional on the covariates and the outcome measured at other timepoints. Consider two subjects for who we would like to predict the weight 1 week before the intervention based on the weight 3 months before the intervention:

```
newd <- rbind(
  data.frame(id = 1, time = "B3m", weight = coef(eUN.lmm)["(Intercept)"], glucagon = 0),
  data.frame(id = 1, time = "B1w", weight = NA, glucagon = 0),
  data.frame(id = 2, time = "B3m", weight = 100, glucagon = 0),
  data.frame(id = 2, time = "B1w", weight = NA, glucagon = 0)
)
predict(eUN.lmm, newdata = newd, type = "dynamic", keep.newdata = TRUE)
```

	id	time	weight	glucagon	estimate	se	df	lower	upper
1	1	B3m	132.9801	0	NA	NA	Inf	NA	NA
2	1	B1w	NA	0	125.09790	0.6362754	Inf	123.85083	126.3450
3	2	B3m	100.0000	0	NA	NA	Inf	NA	NA
4	2	B1w	NA	0	94.47017	7.2279385	Inf	80.30367	108.6367

The first subjects has the average weight while the second has a much lower weight. The predicted weight for the first subject is then the average weight one week before while it is lower for the second subject due to the positive correlation over time. The predicted value is computed using the formula of the conditional mean for a Gaussian vector:

```
mu1 <- coef(eUN.lmm)[1]
mu2 <- sum(coef(eUN.lmm)[1:2])
Omega_11 <- sigma(eUN.lmm)["B3m","B3m"]
Omega_21 <- sigma(eUN.lmm)["B1w","B3m"]
as.double(mu2 + Omega_21 * (100 - mu1) / Omega_11)
```

```
[1] 94.47017
```

5 Equivalence with other statistical methods

5.1 T-test

A t-test:

```
t.test(weight4 ~ group, data = gastricbypassW)
```

Welch Two Sample t-test

```
data: weight4 by group
t = 0.59144, df = 17.679, p-value = 0.5617
alternative hypothesis: true difference in means between group 0 and group 1 is not equal to 0
95 percent confidence interval:
 -11.73582  20.91582
sample estimates:
mean in group 0 mean in group 1
      104.66      100.07
```

is equivalent to an independent covariance pattern with a different variable for each group:

```
e.ttest4 <- lmm(weight4 ~ group, structure = IND(~group),
               data = gastricbypassW, trace = FALSE)
model.tables(e.ttest4)
```

	estimate	se	df	lower	upper	p.value
(Intercept)	104.66	5.104469	9.00180	93.11324	116.20676	7.270954e-09
group	-4.59	7.760674	17.68244	-20.91558	11.73558	5.617090e-01

Multiple t-tests:

```
e.ttest1 <- lmm(weight1 ~ group, structure = IND(~group),
               data = gastricbypassW, trace = FALSE)
e.ttest2 <- lmm(weight2 ~ group, structure = IND(~group),
               data = gastricbypassW, trace = FALSE)
e.ttest3 <- lmm(weight3 ~ group, structure = IND(~group),
               data = gastricbypassW, trace = FALSE)
```

can be adjusted for multiple comparison by first using the `anova` function to specify the parameter of interest and combining the results using `rbind`:

```
e.mttest <- rbind(anova(e.ttest1, effects = "group=0"),
                  anova(e.ttest2, effects = "group=0"),
                  anova(e.ttest3, effects = "group=0"),
                  anova(e.ttest4, effects = "group=0"))
model.tables(e.mttest, method = "bonferroni")
```


	estimate	se	df	lower	upper	p.value
weight1: group	-10.60	8.971747	17.96464	-35.49775	14.29775	1
weight2: group	-9.50	8.395143	17.98540	-32.79464	13.79464	1
weight3: group	-8.92	8.129458	17.95876	-31.48110	13.64110	1
weight4: group	-4.59	7.760674	17.68244	-26.16472	16.98472	1

⚠ efficient adjustment for multiple comparisons (like "single-step") will not be valid as the correlation structure has not been specified. To do so it is more convenient to work with the long format:

```
e.mttest2 <- mlmm(weight ~ group, structure = IND(~group),
  data = gastricbypassL, trace = FALSE,
  effects = "group=0", by = "time", repetition = ~time|id)
model.tables(e.mttest2, method = "single-step2")
```

	by parameter	estimate	se	df	lower	upper	p.value
1 B3m	group	-10.60	8.971747	17.96464	-30.90465	9.704648	0.3166268
2 B1w	group	-9.50	8.395143	17.98540	-28.49969	9.499691	0.3400966
3 A1w	group	-8.92	8.129458	17.95876	-27.31840	9.478400	0.3566864
4 A3m	group	-4.59	7.760674	17.68244	-22.15378	12.973775	0.6673833

or call the dedicated function `mt.test`:

```
mt.test(weight1+weight2+weight3+weight4~group, data = gastricbypassW)
```

	by parameter	estimate	se	df	lower	upper	p.value
1 weight1	group	-10.60	8.971747	17.96464	-30.92628	9.726280	0.3200568
2 weight2	group	-9.50	8.395143	17.98540	-28.51993	9.519932	0.3433166
3 weight3	group	-8.92	8.129458	17.95876	-27.33800	9.498000	0.3602064
4 weight4	group	-4.59	7.760674	17.68244	-22.17249	12.992487	0.6654633

5.2 Linear regression on the change

A widely spread approach to analyze longitudinal data is to reduce the number of repetitions to 1 by working on the change and then apply 'usual' statistical methods. For instance one could compare the pre- and post- operation values using:

```
gastricbypassW$changeG41 <- gastricbypassW$glucagonAUC4-gastricbypassW$glucagonAUC1
e.change41 <- lm(changeG41 ~ weight1, data = gastricbypassW)
summary(e.change41)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	17865.953	9292.61106	1.922598	0.07050076
weight1	-113.696	71.22173	-1.596367	0.12781371

This turns out to be equivalent to the following mixed model:

```
gastricbypassL41 <- gastricbypassL[gastricbypassL$visit %in% c(1,4),]
gastricbypassL41$time <- droplevels(gastricbypassL41$time)
gastricbypassL41$weight1 <- gastricbypassW$weight1[gastricbypassL41$id]

e.lmm41 <- lmm(glucagonAUC ~ time + time*weight1,
               repetition =~ time|id, structure = "UN",
               data = gastricbypassL41)
model.tables(e.lmm41)
```

	estimate	se	df	lower	upper	p.value
(Intercept)	7730.051990	5737.22268	18.00298	-4323.26268	19783.36666	0.19458155
timeA3m	17865.953183	9292.61106	18.00104	-1657.01749	37388.92385	0.07049983
weight1	1.011014	43.97202	18.00298	-91.36968	93.39171	0.98190941
timeA3m:weight1	-113.695981	71.22173	18.00104	-263.32666	35.93469	0.12781271

This equivalence only holds as there is no missing data.

```
index.missing41 <- which(is.na(gastricbypassW$changeG41))
index.missing41
```

```
integer(0)
```

5.3 Correlation between changes

In some studies, one is interested in studying the relation between two evolutions. Say weight and glucagon before and after the operation:

```
gastricbypassW$changeG41 <- gastricbypassW$glucagonAUC4-gastricbypassW$glucagonAUC1
gastricbypassW$changeW41 <- gastricbypassW$weight4-gastricbypassW$weight1
```

One can evaluate their correlation:

```
cor.test(gastricbypassW$changeW41, gastricbypassW$changeG41)
```

Pearson's product-moment correlation

```
data: gastricbypassW$changeW41 and gastricbypassW$changeG41
t = 1.8667, df = 18, p-value = 0.07831
alternative hypothesis: true correlation is not equal to 0
95 percent confidence interval:
 -0.0484149  0.7174011
sample estimates:
      cor
0.4027343
```

or regress one against the other:

```
e2.change41 <- lm(changeG41 ~ changeW41, data = gastricbypassW)
summary(e2.change41)$coef
```

```
      Estimate Std. Error t value Pr(>|t|)
(Intercept) 13321.9427   5592.8058  2.381978 0.02845909
changeW41     380.3556    203.7541  1.866738 0.07831464
```

This problem can be recast using all measurement as outcomes:

```
keep.col <- c("id","weight1","weight4","glucagonAUC1","glucagonAUC4")
gastricbypassL4 <- reshape(gastricbypassW[,keep.col], direction = "long",
                          idvar = "id", varying = 2:5, timevar = "type", v.names = "value")
gastricbypassL4$type <- factor(gastricbypassL4$type, labels = keep.col[-1])
gastricbypassL4 <- gastricbypassL4[order(gastricbypassL4$id),]
head(gastricbypassL4)
```

```
      id      type  value
1.1  1  weight1  127.20
1.2  1  weight4  108.10
1.3  1 glucagonAUC1 5032.50
1.4  1 glucagonAUC4 9249.45
2.1  2  weight1  165.20
2.2  2  weight4  132.00
```

fitting an unstructured mixed model:

```
e.lmm4 <- lmm(value ~ type,
              repetition = ~type|id, structure = "UN",
              data = gastricbypassL4)
```

extract the residual covariance matrix:

```
sigma.lmm4 <- sigma(e.lmm4)
sigma.lmm4
```

```
          weight1    weight4 glucagonAUC1 glucagonAUC4
weight1    410.8475    326.8357      415.3727    -46296.33
weight4    326.8357    290.8350     -5983.5871    -34434.03
glucagonAUC1 415.3727 -5983.5871 14299430.9269 -4229230.69
glucagonAUC4 -46296.3339 -34434.0320 -4229230.6877 20065722.32
```

Deduce the residual covariance matrix for the change:

```
Mcon <- cbind(c(-1,1,0,0),c(0,0,-1,1))
sigmeChange.lmm4 <- t(Mcon) %*% sigma.lmm4 %*% Mcon
dimnames(sigmeChange.lmm4) <- list(c("d.weight","d.glucagonAUC"),
                                   c("d.weight","d.glucagonAUC"))
sigmeChange.lmm4
```

```
          d.weight d.glucagonAUC
d.weight    48.01103    18261.26
d.glucagonAUC 18261.26175    42823614.62
```

and the correlation or covariance:

```
cov2cor(sigmeChange.lmm4)[1,2]
sigmeChange.lmm4[1,2]/sigmeChange.lmm4[1,1]
```

```
[1] 0.4027343
[1] 380.3556
```

The uncertainty can be quantified using a delta method:

```
estimate(e.lmm4, function(p){
  Sigma.change <- t(Mcon) %*% sigma(e.lmm4, p = p) %*% Mcon
  c(cor = cov2cor(Sigma.change)[1,2],
    beta = Sigma.change[1,2]/Sigma.change[1,1])
})
```

```
      estimate      se      df      lower      upper  p.value
cor    0.4027343 0.1922078 2.660595 -0.2555602  1.061029 0.1386265
beta 380.3555798 198.3453360 2.798661 -277.3518013 1038.062961 0.1575655
```

The standard errors and degrees of freedom do not match the univariate analysis, suggesting probably poor small sample properties of this technic.

6 Missing values and imputation

We reconsider the example of the previous section, but now in presence of missing values. The `summarize` function can be used to describe the amount of missing data at each repetition:

```
sss <- summarize(glucagon ~ time, data = gastricbypassL, na.rm = TRUE)
cbind(sss[,1:4], pc = paste0(100 * sss$missing / (sss$missing + sss$observed), "%"))
```

```
  outcome time observed missing pc
1 glucagon B3m      20       0 0%
2 glucagon B1w      19       1 5%
3 glucagon A1w      19       1 5%
4 glucagon A3m      20       0 0%
```

For more detail about the missing data patterns, see the `summarizeNA` function:

```
summarizeNA(data = gastricbypassL, repetition = ~ time|id)
```

```
  variable frequency missing.pattern n.missing id B3m B1w A1w A3m
  visit          20          00000           0 0 0 0 0 0
  weight          20          00000           0 0 0 0 0 0
glucagonAUC       18          00000           0 0 0 0 0 0
               1          00100           1 0 0 1 0 0
               1          00010           1 0 0 0 1 0
  baseline        20          00000           0 0 0 0 0 0
  glucagon         18          00000           0 0 0 0 0 0
               1          00100           1 0 0 1 0 0
               1          00010           1 0 0 0 1 0
  group           20          00000           0 0 0 0 0 0
```

To begin with we will only consider 1 week before and 1 week after surgery:

```
## long format
gastricbypassL32 <- gastricbypassL[gastricbypassL$visit %in% c(3,2),]
gastricbypassL32$time <- droplevels(gastricbypassL32$time)
gastricbypassL32$weight1 <- gastricbypassW$weight1[gastricbypassL32$id]
## wide format
gastricbypassW$changeG32 <- gastricbypassW$glucagonAUC3-gastricbypassW$glucagonAUC2
```

6.1 Full information approach

LMM uses a full information approach:

```
e.lmm32 <- lmm(glucagonAUC ~ time + time*weight1,
              repetition =~ time|id, structure = "UN",
              data = gastricbypassL32)
model.tables(e.lmm32)
```

	estimate	se	df	lower	upper	p.value
(Intercept)	2226.30678	4973.21491	17.01148	-8265.72037	12718.33393	0.6600471546
timeA1w	37469.89400	8950.26818	17.88948	18657.74792	56282.04008	0.0005612515
weight1	37.90933	37.87004	17.01113	-41.98548	117.80414	0.3308362562
timeA1w:weight1	-213.20181	68.15807	17.71309	-356.56304	-69.84058	0.0058968630

whereas a linear model would perform a complete case approach:

```
e.change32 <- lm(changeG32 ~ weight1, data = gastricbypassW)
summary(e.change32)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	38101.9400	9417.61506	4.045816	0.0009373023
weight1	-217.2672	71.25218	-3.049271	0.0076504030

In the former the likelihood is evaluated using all observations, even those from individuals with some (but not all) missing outcome values: baseline is used even if follow-up is missing. In the later the likelihood is only evaluated on individuals with no missing outcome values: if follow-up is missing then baseline is not used. Indeed:


```
coef(lm(changeG32 ~ weight1, data = gastricbypassW[-c(5,15),]))
```

(Intercept)	weight1
38101.9400	-217.2672

The estimates of the LMM can be retrived using a linear model where we have imputed the conditional expectation of the missing values given the observed value and the estimated model parameters: (see section 6.3 for a graphical representation)

```
gastricbypassWA <- fitted(e.lmm32, impute = TRUE, format = "wide")
gastricbypassWA$change32 <- gastricbypassWA$glucagonAUC_A1w - gastricbypassWA$glucagonAUC_B1w
gastricbypassWA$weight1 <- gastricbypassW$weight1[match(gastricbypassW$id,gastricbypassWA$id)]
coef(lm(change32 ~ weight1, data = gastricbypassWA))
```

(Intercept)	weight1
37469.8940	-213.2018

 Standard errors, confidence intervals, and p-values from this linear model should not be trusted as they do not account for the uncertainty in the imputed values.

6.2 Complete case approach

The `lmmCC` can be used to obtain the LMM that is equivalent to a linear regression. In the case of the comparing the change between groups, the `repetition` argument should indicate how the change has been computed:

```
e.lmmCC <- lmmCC(e.change32, repetition = changeG32 ~ glucagonAUC3-glucagonAUC2|id)
model.tables(e.lmmCC)
```

Remove 2 clusters (4 observations)

- 2 observations with missing data (2 clusters)
- 0 missing repetitions (0 clusters)

	estimate	se	df	lower	upper	p.value
(Intercept)	-36283.0356	12841.66795	15.99997	-63506.15929	-9059.91191	0.0121849728
time	38101.9400	9417.61506	16.00030	18137.51789	58066.36212	0.0009372705
weight1	257.7767	97.15802	15.99997	51.81085	463.74249	0.0173566171
time:weight1	-217.2672	71.25218	16.00030	-368.31484	-66.21956	0.0076502759

As output, the data from two clusters (i.e. 4 observations) has been excluded before fitting the LMM (instead of just the 2 observations with missing values for the full information approach). The interaction term of the LMM matches the regression coefficient of the linear model:

```
summary(e.change32)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	38101.9400	9417.61506	4.045816	0.0009373023
weight1	-217.2672	71.25218	-3.049271	0.0076504030

In the case of regressing two changes:

```
gastricbypassW$changeW32 <- gastricbypassW$weight3 - gastricbypassW$weight2
e2g.change32 <- lm(changeG32 ~ changeW32 + group, data = gastricbypassW)
summary(e2g.change32)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	2720.5540	6930.588	0.3925430	0.7001787
changeW32	-783.8895	1122.541	-0.6983171	0.4956633
group	6059.7378	3525.140	1.7190062	0.1061756

the `repetition` argument should indicate how each change has been computed:

```
e2.lmmCC <- lmmCC(e2g.change32, repetition = list(changeG32 ~ glucagonAUC3-glucagonAUC2|id,
                                                    changeW32 ~ weight3-weight2|id))
model.tables(e2.lmmCC)
```

Remove 2 clusters (8 observations)

- 2 observations with missing data (2 clusters)
- 0 missing repetitions (0 clusters)

	estimate	se	df	lower	upper	p.value
cor	-0.1774435	0.2416113	1.714523	-1.401851	1.046964	0.5499188
beta	-783.8895353	1081.5567036	2.338122	-4848.482516	3280.703446	0.5342415

We retrieve the same estimate for the effect of change in weights but the uncertainty (standard error, confidence intervals, p.value) do not match. They should be asymptotically correct but may not have very good small sample properties.

6.3 Imputation

When fitting a linear mixed model on a dataset with missing values:

```
eUN.lmmNA <- lmm(glucagon ~ time, repetition = ~time|id, data = gastricbypassL)
nobs(eUN.lmmNA)
```

obs	cluster	missing.obs	missing.cluster
78	20	2	0

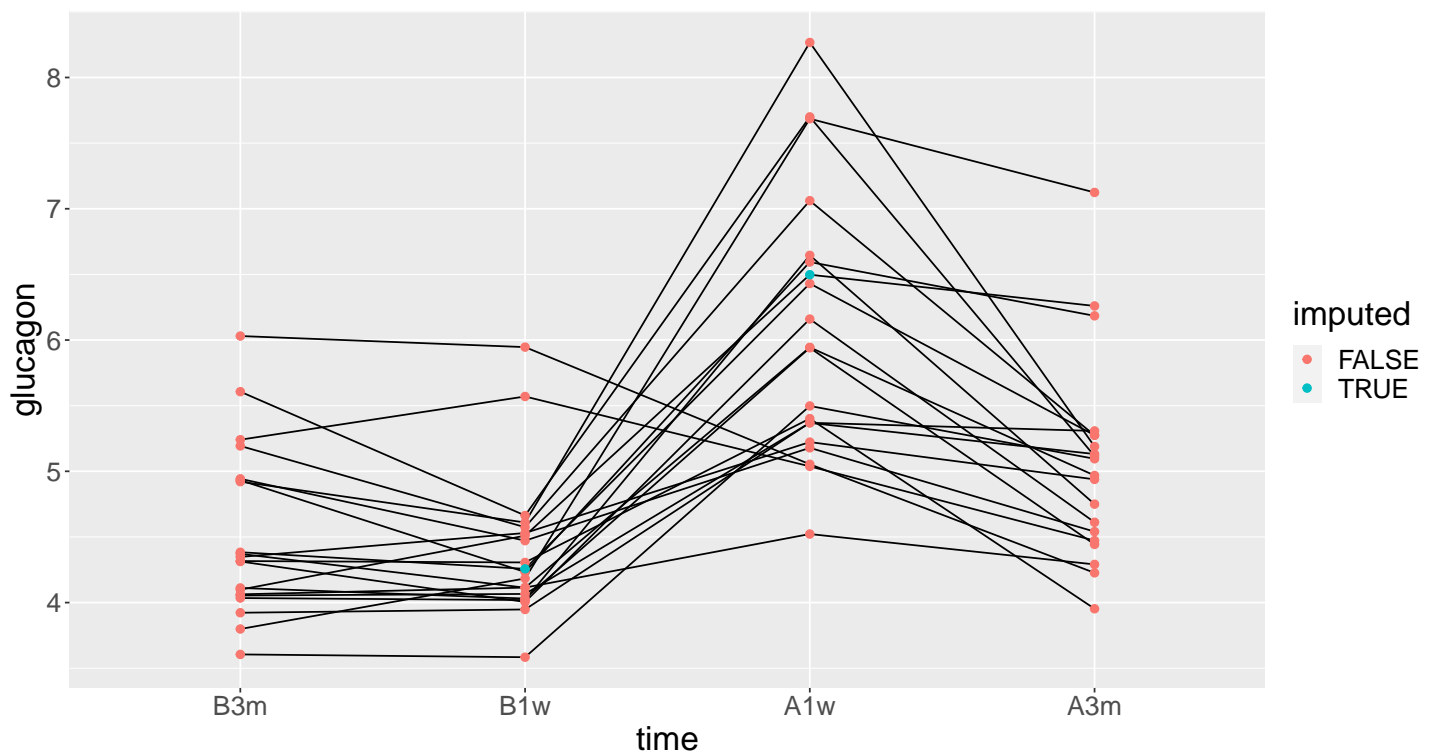
It is possible to extract the most likely value for these missing observations using the `fitted` function with argument `impute=TRUE`:

```
eData <- fitted(eUN.lmmNA, impute = TRUE, keep.newdata = TRUE)
eData$treat <- eData$treat2 <- eData$timeXtreat <- NULL
eData[eData$id %in% eData[eData$imputed, "id"],]
```

	id	visit	time	weight	glucagonAUC	baseline	glucagon	group	imputed
5	5	1	B3m	113.1	7090.5	TRUE	4.383738	1	FALSE
15	15	1	B3m	115.0	5410.5	TRUE	4.098741	1	FALSE
25	5	2	B1w	105.6	NA	TRUE	4.256984	1	TRUE
35	15	2	B1w	109.7	7833.0	TRUE	4.509697	1	FALSE
45	5	3	A1w	99.9	19155.0	FALSE	6.430376	1	FALSE
55	15	3	A1w	103.5	NA	FALSE	6.497856	1	TRUE
65	5	4	A3m	87.7	12345.0	FALSE	5.275118	1	FALSE
75	15	4	A3m	94.1	18148.5	FALSE	6.259632	1	FALSE

Missing outcome values in the dataset have been replaced by its most likely value (which is the same as the dynamic prediction, described previously). A column `imputed` has also been added to differentiate between the modeled and observed value. Visually:

```
ggplot(eData, aes(x=time,y=glucagon, group=id)) + geom_line() + geom_point(aes(color=imputed))
```

It is possible to sample from the estimated distribution of the missing value instead of using the most likely value, e.g. accounting for residual variance and uncertainty related to parameter estimation:

```
set.seed(10)
index.na <- which(is.na(gastricbypassL$glucagonAUC))
fitted(eUN.lmmNA, impute = TRUE, se.impute = "total")[index.na]
fitted(eUN.lmmNA, impute = TRUE, se.impute = "total")[index.na]
fitted(eUN.lmmNA, impute = TRUE, se.impute = "total")[index.na]
```

```
[1] 4.262434 6.305287
```

```
[1] 3.858267 5.871642
```

```
[1] 4.342624 6.905246
```

6.4 Multiple imputation

The `mlmm` function can be used to perform stratified analyses, typically useful when performing multiple imputations. Consider the wide format of the dataset where a few values are missing:

```
data(gastricbypassW, package = "LMMstar")
colSums(is.na(gastricbypassW))
```

```
      id      weight1      weight2      weight3      weight4 glucagonAUC1 glucagonAUC2
      0              0              0              0              0              0              1
glucagonAUC3 glucagonAUC4
      1              0
```

We use `mice` to generate a number of imputed datasets (here 5):

```
library(mice)
set.seed(10)
gastricbypassW.mice <- mice(gastricbypassW, m = 5, printFlag = FALSE)
gastricbypassW.NNA <- complete(gastricbypassW.mice, action = "long")
table(gastricbypassW.NNA$.imp)
```

```
Advarselsbesked:
Number of logged events: 110
```

```
 1  2  3  4  5
20 20 20 20 20
```

We can then use `mlmm` to perform a separate linear regression per dataset:

```
e.mlmm <- mlmm(glucagonAUC3~glucagonAUC2+weight2, data=gastricbypassW.NNA,
               by = ".imp", effects = "weight2=0", trace = FALSE)
model.tables(e.mlmm)
```

```
by parameter estimate      se      df      lower      upper      p.value
1  1  weight2 -204.6291 62.88617 17.0034 -337.3053 -71.95289 0.004670840
2  2  weight2 -194.4004 62.31006 17.0034 -325.8611 -62.93968 0.006231893
3  3  weight2 -211.9042 65.51654 17.0034 -350.1299 -73.67848 0.004872354
4  4  weight2 -199.8417 62.12071 17.0034 -330.9029 -68.78041 0.005058119
5  5  weight2 -199.9269 62.16057 17.0034 -331.0722 -68.78152 0.005065662
```

and pool the results using Rubin's rule:

```
model.tables(e.mlmm, method = "pool.rubin")
```

```
      estimate      se      df      lower      upper      p.value
<1, 5> -202.1404 63.4192 15.09811 -337.2388 -67.04208 0.006078676
```

This matches⁴ the results obtained with the `mice` package:

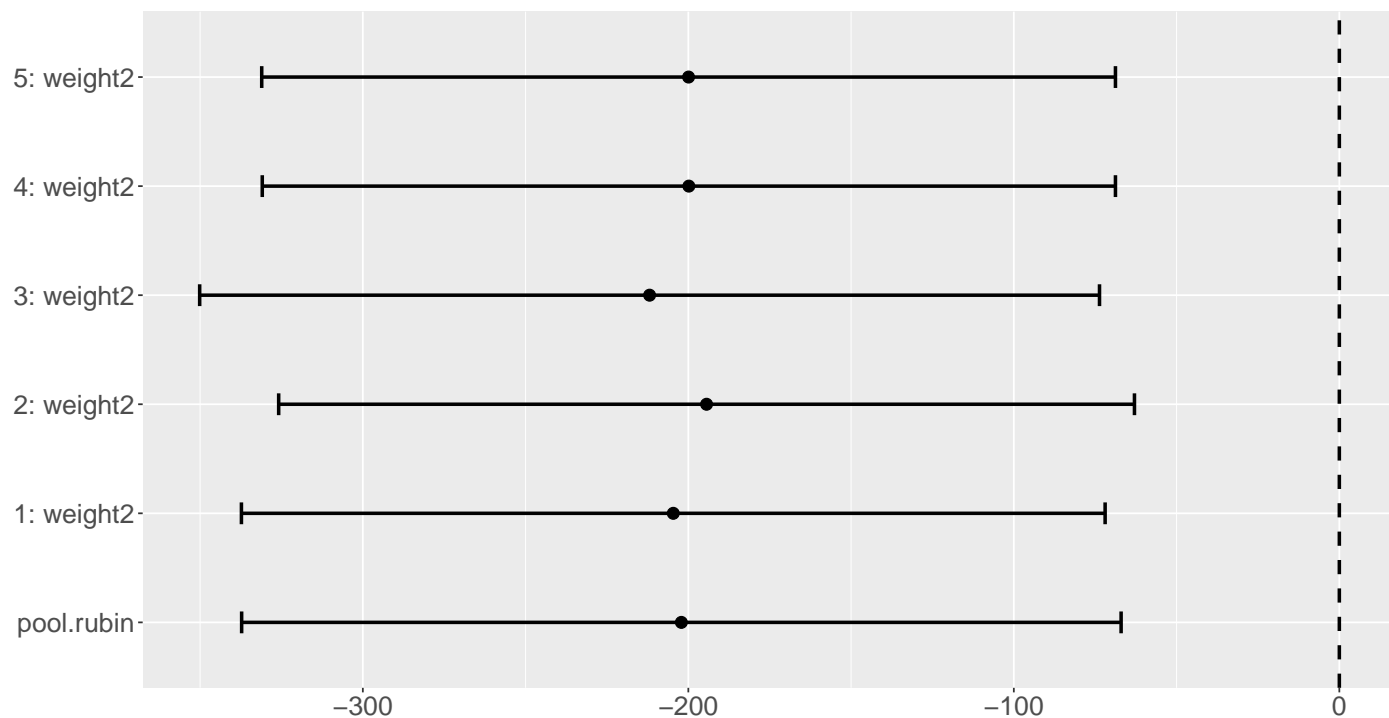
⁴almost exactly, only the degrees of freedom are a little different

```
e.mice <- with(data=gastricbypassW.mice,exp=lm(glucagonAUC3~glucagonAUC2+weight2))
summary(pool(e.mice))
```

	term	estimate	std.error	statistic	df	p.value
1	(Intercept)	4.119699e+04	7674.2675772	5.3681988	15.08457	7.675819e-05
2	glucagonAUC2	7.038742e-02	0.3689445	0.1907805	15.23549	8.512165e-01
3	weight2	-2.021404e+02	63.4191998	-3.1873698	15.09481	6.080058e-03

One can use the `plot` function to obtain a forest plot of the individual estimates along with the pooled estimate:

```
plot(e.mlmm, method = c("pool.rubin", "none"))
```



7 Data generation

Simulate some data in the wide format:

```
set.seed(10) ## ensure reproductibility
n.obs <- 100
n.times <- 4
mu <- rep(0,4)
gamma <- matrix(0, nrow = n.times, ncol = 10) ## add interaction
gamma[,6] <- c(0,1,1.5,1.5)
dW <- sampleRem(n.obs, n.times = n.times, mu = mu, gamma = gamma, format = "wide")
head(round(dW,3))
```

	id	X1	X2	X3	X4	X5	X6	X7	X8	X9	X10	Y1	Y2	Y3	Y4
1	1	1	0	1	1	0	-0.367	1.534	-1.894	1.729	0.959	1.791	2.429	3.958	2.991
2	2	1	0	1	2	0	-0.410	2.065	1.766	0.761	-0.563	2.500	4.272	3.002	2.019
3	3	0	0	2	1	0	-1.720	-0.178	2.357	1.966	1.215	-3.208	-5.908	-4.277	-5.154
4	4	0	0	0	1	0	0.923	-2.089	0.233	1.307	-0.906	-2.062	0.397	1.757	-1.380
5	5	0	0	2	1	0	0.987	5.880	0.385	0.028	0.820	7.963	7.870	7.388	8.609
6	6	0	0	1	1	2	-1.075	0.479	2.202	0.900	-0.739	0.109	-1.602	-1.496	-1.841

Simulate some data in the long format:

```
set.seed(10) ## ensure reproductibility
dL <- sampleRem(n.obs, n.times = n.times, mu = mu, gamma = gamma, format = "long")
head(dL)
```

	id	visit	Y	X1	X2	X3	X4	X5	X6	X7	X8	X9	X10
1	1	1	1.791444	1	0	1	1	0	-0.3665251	1.533815	-1.894425	1.7288665	0.9592499
2	1	2	2.428570	1	0	1	1	0	-0.3665251	1.533815	-1.894425	1.7288665	0.9592499
3	1	3	3.958350	1	0	1	1	0	-0.3665251	1.533815	-1.894425	1.7288665	0.9592499
4	1	4	2.991198	1	0	1	1	0	-0.3665251	1.533815	-1.894425	1.7288665	0.9592499
5	2	1	2.500179	1	0	1	2	0	-0.4097541	2.065413	1.765841	0.7613348	-0.5630173
6	2	2	4.272357	1	0	1	2	0	-0.4097541	2.065413	1.765841	0.7613348	-0.5630173

8 Modifying default options

The `LMMstar.options` method enable to get and set the default options used by the package. For instance, the default option for the information matrix is:

```
LMMstar.options("type.information")
```

```
$type.information  
[1] "observed"
```

To change the default option to "expected" (faster to compute but less accurate p-values and confidence intervals in small samples) use:

```
LMMstar.options(type.information = "expected")
```

To restore the original default options do:

```
LMMstar.options(reinitialise = TRUE)
```

9 R session

Details of the R session used to generate this document:

```
sessionInfo()
```

```
R version 4.2.0 (2022-04-22 ucrt)
```

```
Platform: x86_64-w64-mingw32/x64 (64-bit)
```

```
Running under: Windows 10 x64 (build 19045)
```

```
Matrix products: default
```

```
locale:
```

```
[1] LC_COLLATE=Danish_Denmark.utf8 LC_CTYPE=Danish_Denmark.utf8
```

```
LC_MONETARY=Danish_Denmark.utf8
```

```
[4] LC_NUMERIC=C
```

```
LC_TIME=Danish_Denmark.utf8
```

```
attached base packages:
```

```
[1] parallel grid stats graphics grDevices utils datasets methods base
```

```
other attached packages:
```

```
[1] mice_3.14.0 sandwich_3.0-2 scales_1.2.1 rlang_1.1.1
[5] pbapply_1.7-0 numDeriv_2016.8-1.1 nlme_3.1-158 lava_1.7.2.1
[9] doSNOW_1.0.20 snow_0.4-4 iterators_1.0.14 foreach_1.5.2
[13] copula_1.1-2 lme4_1.1-29 Matrix_1.5-1 LMMstar_1.0.0
[17] ggpubr_0.4.0 multcomp_1.4-22 TH.data_1.1-1 MASS_7.3-57
[21] survival_3.3-1 mvtnorm_1.2-3 qqtest_1.2.0 emmeans_1.8.8-090002
[25] ggplot2_3.4.3
```

```
loaded via a namespace (and not attached):
```

```
[1] butils.base_1.2 minqa_1.2.4 colorspace_2.1-0 ggsignif_0.6.3
[5] ellipsis_0.3.2 estimability_1.4.1 parameters_0.18.2 fs_1.6.3
[9] listenv_0.9.0 farver_2.1.1 remotes_2.4.2 gsl_2.1-8
[13] fansi_1.0.4 codetools_0.2-18 splines_4.2.0 doParallel_1.0.17
[17] cachem_1.0.8 pkgload_1.3.0 nloptr_2.0.3 broom_0.8.0
[21] stabledist_0.7-1 effectsize_0.7.0.5 shiny_1.7.2 compiler_4.2.0
[25] backports_1.4.1 fastmap_1.1.1 cli_3.6.1 later_1.3.0
[29] htmltools_0.5.6 prettyunits_1.1.1 tools_4.2.0 lmerTest_3.1-3
[33] coda_0.19-4 gtable_0.3.4 glue_1.6.2 reshape2_1.4.4
[37] dplyr_1.1.3 Rcpp_1.0.11 carData_3.0-5 vctr_0.6.3
[41] insight_0.18.4 stringr_1.5.0 globals_0.16.2 ps_1.7.1
[45] mime_0.12 miniUI_0.1.1.1 lifecycle_1.0.3 devtools_2.4.4
[49] rstatix_0.7.0 future_1.31.0 zoo_1.8-11 promises_1.2.0.1
[53] memoise_2.0.1 gridExtra_2.3 stringi_1.7.12 bayestestR_0.13.0
[57] pcaPP_2.0-3 boot_1.3-28 pkgbuild_1.3.1 pkgconfig_2.0.3
[61] lattice_0.20-45 purrr_1.0.2 htmlwidgets_1.6.2 labeling_0.4.3
[65] cowplot_1.1.1 tidyselect_1.2.0 processx_3.6.1 parallelly_1.34.0
[69] plyr_1.8.7 magrittr_2.0.3 R6_2.5.1 generics_0.1.3
[73] profvis_0.3.7 ADGofTest_0.3 pillar_1.9.0 withr_2.5.1
```

[77] mgcv_1.8-40	datawizard_0.6.1	abind_1.4-5	pspline_1.0-19
[81] tibble_3.2.1	future.apply_1.10.0	crayon_1.5.1	car_3.1-0
[85] utf8_1.2.3	urlchecker_1.0.1	usethis_2.1.6	data.table_1.14.2
[89] callr_3.7.2	digest_0.6.33	xtable_1.8-4	tidyr_1.3.0
[93] httpuv_1.6.5	stats4_4.2.0	munsell_0.5.0	sessioninfo_1.2.2

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Appendix A Likelihood in a linear mixed model

Denote by \mathbf{Y} a vector of m outcomes, \mathbf{X} a vector of p covariates, $\mu(\boldsymbol{\Theta}, \mathbf{X})$ the modeled mean, and $\Omega(\boldsymbol{\Theta}, \mathbf{X})$ the modeled residual variance-covariance. We consider n replicates (i.e. $\mathbf{Y}_1, \dots, \mathbf{Y}_n$) and $VX_1, \dots, \mathbf{X}_n$ along with a vector of weights $\omega = (w_1, \dots, w_n)$, which are by default all equal to 1.

A.1 Log-likelihood

The restricted log-likelihood in a linear mixed model can then be written:

$$\begin{aligned} \mathcal{L}(\boldsymbol{\Theta}|\mathbf{Y}, \mathbf{X}) = & \frac{p}{2} \log(2\pi) - \frac{1}{2} \log \left(\left| \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top \right| \right) \\ & + \sum_{i=1}^n w_i \left(-\frac{m}{2} \log(2\pi) - \frac{1}{2} \log |\Omega_i(\boldsymbol{\Theta})| - \frac{1}{2} (\mathbf{Y}_i - \mu(\boldsymbol{\Theta}, \mathbf{X}_i)) \Omega_i(\boldsymbol{\Theta})^{-1} (\mathbf{Y}_i - \mu(\boldsymbol{\Theta}, \mathbf{X}_i))^\top \right) \end{aligned} \quad (\text{A})$$

This is what the `logLik` method is computing for the REML criteria. The red term is specific to the REML criteria and prevents from computing individual contributions to the likelihood⁵. The blue term is what `logLik` outputs for the ML criteria when setting the argument `indiv` to `TRUE`.

A.2 Score

Using that $\partial \log(\det(X)) = \text{tr}(X^{-1} \partial(X))$, the score is obtained by derivating once the log-likelihood, i.e., for $\theta \in \boldsymbol{\Theta}$:

$$\begin{aligned} \mathcal{S}(\theta) = & \frac{\partial \mathcal{L}(\boldsymbol{\Theta}|\mathbf{Y}, \mathbf{X})}{\partial \theta} = \frac{1}{2} \text{tr} \left(\left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top \right)^{-1} \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \Omega_i(\boldsymbol{\Theta})^{-1} \mathbf{X}_i^\top \right) \right) \\ & + \sum_{i=1}^n w_i \left(-\frac{1}{2} \text{tr} \left(\Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \right) + \frac{\partial \mu(\boldsymbol{\Theta}, \mathbf{X}_i)}{\partial \theta} \Omega_i(\boldsymbol{\Theta})^{-1} (\mathbf{Y}_i - \mu(\boldsymbol{\Theta}, \mathbf{X}_i))^\top \right. \\ & \quad \left. + \frac{1}{2} (\mathbf{Y}_i - \mu(\boldsymbol{\Theta}, \mathbf{X}_i)) \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \Omega_i(\boldsymbol{\Theta})^{-1} (\mathbf{Y}_i - \mu(\boldsymbol{\Theta}, \mathbf{X}_i))^\top \right). \end{aligned}$$

This is what the `score` method is computing for the REML criteria. The red term is specific to the REML criteria and prevents from computing the score relative to each cluster. The blue term is what `score` outputs for the ML criteria when setting the argument `indiv` to `TRUE`.

⁵The REML is the likelihood of the observations divided by the prior on the estimated mean parameters $\hat{\boldsymbol{\Theta}}_\mu \sim \mathcal{N}(\mu, (\mathbf{X} \Omega^{-1}(\boldsymbol{\Theta}) \mathbf{X}^\top)^{-1})$. This corresponds to $\frac{1}{\sqrt{2\pi^p} |(\sum_{i=1}^n \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top)^{-1}|} \exp \left(-(\hat{\boldsymbol{\Theta}}_\mu - \mu) (2 \sum_{i=1}^n \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top)^{-1} (\hat{\boldsymbol{\Theta}}_\mu - \mu)^\top \right)$. Since μ will be estimated to be $\boldsymbol{\Theta}_\mu$, the exponential term equals 1 and thus does not contribute to the log-likelihood. One divided by the other term gives $\sqrt{2\pi^p} |(\sum_{i=1}^n \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top)|^{-1}$. The log of this term equals the red term

A.3 Hessian

Derivating a second time the log-likelihood gives the hessian, $\mathcal{H}(\Theta)$, with element⁶:

$$\begin{aligned}\mathcal{H}(\theta, \theta') &= \frac{\partial^2 \mathcal{L}(\Theta | \mathbf{Y}, \mathbf{X})}{\partial \theta \partial \theta'} = \frac{\partial \mathcal{S}(\theta)}{\partial \theta'} \\ &= \frac{1}{2} \text{tr} \left(\left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \mathbf{X}_i^\top \right)^{-1} \left\{ \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \left(\frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} - 2 \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \right) \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right. \right. \\ &\quad \left. \left. + \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right) \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \mathbf{X}_i^\top \right)^{-1} \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right) \right\} \right) \\ &\quad + \sum_{i=1}^n w_i \left(\frac{1}{2} \text{tr} \left(\Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} - \Omega_i(\Theta)^{-1} \frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} \right) \right. \\ &\quad \left. - \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \boldsymbol{\varepsilon}_i(\Theta)^\top - \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta'} \right. \\ &\quad \left. + \frac{1}{2} \boldsymbol{\varepsilon}_i(\Theta) \Omega_i(\Theta)^{-1} \left(\frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} - \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} - \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \right) \Omega_i(\Theta)^{-1} \boldsymbol{\varepsilon}_i(\Theta)^\top \right).\end{aligned}$$

where $\boldsymbol{\varepsilon}_i(\Theta) = \mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i)$.

The `information` method will (by default) return the (observed) information which is the opposite of the hessian. So multiplying the previous formula by -1 gives what `information` output for the REML criteria. The red term is specific to the REML criteria and prevents from computing the information relative to each cluster. The blue term is what `information` outputs for the ML criteria (up to a factor -1) when setting the argument `indiv` to `TRUE`.

A possible simplification is to use the expected hessian at the maximum likelihood. Indeed for any deterministic matrix A :

- $\mathbb{E}[A(\mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i))^\top | \mathbf{X}_i] = 0$
- $\mathbb{E}[(\mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i))A(\mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i))^\top | \mathbf{X}_i] = \text{tr}(A \text{Var}(\mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i)))$

when $\mathbb{E}[\mathbf{Y}_i - \mu(\Theta, \mathbf{X}_i)] = 0$. This leads to:

$$\begin{aligned}\mathbb{E}[\mathcal{H}(\theta, \theta') | \mathbf{X}] &= \frac{1}{2} \text{tr} \left(\left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \mathbf{X}_i^\top \right)^{-1} \left\{ \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \left(\frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} - 2 \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \right) \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right. \right. \\ &\quad \left. \left. + \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right) \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \mathbf{X}_i^\top \right)^{-1} \left(\sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \mathbf{X}_i^\top \right) \right\} \right) \\ &\quad + \sum_{i=1}^n w_i \left(-\frac{1}{2} \text{tr} \left(\Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} \right) - \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta'} \right) \quad (\text{B})\end{aligned}$$

This is what `information` output when the argument `type.information` is set to "expected" (up to a factor -1).

⁶if one is relative to the mean and the other to the variance then they are respectively θ and θ'

A.4 Degrees of freedom

Degrees of freedom are computed using a Satterthwaite approximation, i.e. for an estimate coefficient $\hat{\beta} \in \hat{\Theta}$ with standard error $\sigma_{\hat{\beta}}$, the degree of freedom is:

$$df(\sigma_{\hat{\beta}}) = \frac{2\sigma_{\hat{\beta}}^4}{\text{Var}[\hat{\sigma}_{\hat{\beta}}]}$$

Using a first order Taylor expansion we can approximate the variance term as:

$$\begin{aligned} \text{Var}[\hat{\sigma}_{\hat{\beta}}] &\approx \frac{\partial \hat{\sigma}_{\hat{\beta}}}{\partial \Theta} \Sigma_{\Theta} \frac{\partial \hat{\sigma}_{\hat{\beta}}}{\partial \Theta}^{\top} \\ &\approx c_{\beta} (\hat{\mathcal{I}}_{\hat{\Theta}})^{-1} \frac{\partial \hat{\mathcal{I}}_{\hat{\Theta}}}{\partial \Theta} (\hat{\mathcal{I}}_{\hat{\Theta}})^{-1} c_{\beta}^{\top} \Sigma_{\Theta} c_{\beta} (\hat{\mathcal{I}}_{\hat{\Theta}})^{-1} \frac{\partial \hat{\mathcal{I}}_{\hat{\Theta}}}{\partial \Theta}^{\top} (\hat{\mathcal{I}}_{\hat{\Theta}})^{-1} c_{\beta} \end{aligned}$$

where Σ_{Θ} is the variance-covariance matrix of all model coefficients, \mathcal{I}_{Θ} the information matrix for all model coefficients, c_{β} a matrix used to select the element relative to β in the first derivative of the information matrix, and $\frac{\partial}{\partial \Theta}$ denotes the vector of derivatives with respect to all model coefficients.

The derivative of the information matrix (i.e. negative hessian) can then be computed using numerical derivatives or using analytical formula. To obtain the later we first notice that:

$$\begin{aligned} \mathcal{H}(\theta, \theta') &= \mathbb{E}[\mathcal{H}(\theta, \theta') | \mathbf{X}] \\ &+ \sum_{i=1}^n w_i \left(\text{tr} \left(\Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} - \Omega_i(\Theta)^{-1} \frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} \right) \right. \\ &\quad - \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \epsilon_i(\Theta)^{\top} \\ &\quad \left. + \frac{1}{2} \epsilon_i(\Theta) \Omega_i(\Theta)^{-1} \left(\frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} - \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} - \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \right) \Omega_i(\Theta)^{-1} \epsilon_i(\Theta)^{\top} \right) \end{aligned} \quad (\text{C})$$

where

$$\begin{aligned} \mathbb{E}[\mathcal{H}(\theta, \theta') | \mathbf{X}] &= \frac{1}{2} \text{tr} \left(A(\Theta)^{-1} \left(\sum_{i=1}^n w_i b_i(\Theta) B_i(\Theta) b_i^{\top}(\Theta) + C(\Theta) A(\Theta)^{-1} C^{\top}(\Theta) \right) \right) + \sum_{i=1}^n w_i E_i(\Theta) \\ E_i(\Theta) &= \frac{1}{2} \text{tr} \left(\Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \Omega_i(\Theta)^{-1} \frac{\partial \Omega_i(\Theta)}{\partial \theta} \right) - \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta} \Omega_i(\Theta)^{-1} \frac{\partial \mu(\Theta, \mathbf{X}_i)}{\partial \theta'}^{\top} \\ A(\Theta) &= \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \mathbf{X}_i^{\top} \\ B_i(\Theta) &= \frac{\partial^2 \Omega_i(\Theta)}{\partial \theta \partial \theta'} - 2 \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta'} \\ b_i(\Theta) &= \mathbf{X}_i \Omega_i^{-1} \\ C(\Theta) &= \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\Theta) \frac{\partial \Omega_i(\Theta)}{\partial \theta} \Omega_i(\Theta)^{-1} \mathbf{X}_i^{\top} \end{aligned}$$

So we will first derive the derivative of $\mathbb{E}[\mathcal{H}(\theta, \theta') | \mathbf{X}]$ and then the one of the blue term in Equation C. To simplify the derivation of the formula we will only derive them at the maximum likelihood, i.e. when

$\mathbb{E} \left[\frac{\partial \mathcal{H}(\theta, \theta' | \mathbf{X})}{\partial \theta''} \right] = \frac{\partial \mathbb{E}[\mathcal{H}(\theta, \theta' | \mathbf{X})]}{\partial \theta''}$ where the expectation is taken over \mathbf{X} . We first notice that the derivative with respect to the mean parameters is 0. So we just need to compute the derivative with respect to a variance parameter θ'' :

$$\begin{aligned} & \frac{\partial A(\boldsymbol{\Theta})^{-1} \left(\sum_{i=1}^n w_i b_i(\boldsymbol{\Theta}) B_i(\boldsymbol{\Theta}) b_i^\top(\boldsymbol{\Theta}) + C(\boldsymbol{\Theta}) A(\boldsymbol{\Theta})^{-1} C^\top(\boldsymbol{\Theta}) \right)}{\partial \theta''} \\ &= A(\boldsymbol{\Theta})^{-1} \frac{\partial A(\boldsymbol{\Theta})}{\partial \theta''} A(\boldsymbol{\Theta})^{-1} \left(\sum_{i=1}^n w_i b_i(\boldsymbol{\Theta}) B_i(\boldsymbol{\Theta}) b_i^\top(\boldsymbol{\Theta}) + C(\boldsymbol{\Theta}) A(\boldsymbol{\Theta})^{-1} C^\top(\boldsymbol{\Theta}) \right) \\ &+ A(\boldsymbol{\Theta})^{-1} \left(\sum_{i=1}^n w_i \left(\frac{\partial b_i(\boldsymbol{\Theta})}{\partial \theta''} B_i(\boldsymbol{\Theta}) b_i^\top(\boldsymbol{\Theta}) + b_i(\boldsymbol{\Theta}) \frac{\partial B_i(\boldsymbol{\Theta})}{\partial \theta''} b_i^\top(\boldsymbol{\Theta}) + b_i(\boldsymbol{\Theta}) B_i(\boldsymbol{\Theta}) \frac{\partial b_i^\top(\boldsymbol{\Theta})}{\partial \theta''} \right. \right. \\ &\quad \left. \left. + \frac{\partial C(\boldsymbol{\Theta})}{\partial \theta''} A^{-1}(\boldsymbol{\Theta}) C^\top(\boldsymbol{\Theta}) + C(\boldsymbol{\Theta}) A^{-1} \frac{\partial A(\boldsymbol{\Theta})}{\partial \theta''} A^{-1} C^\top(\boldsymbol{\Theta}) + C(\boldsymbol{\Theta}) A^{-1}(\boldsymbol{\Theta}) \frac{\partial C^\top(\boldsymbol{\Theta})}{\partial \theta''} \right) \right) \end{aligned}$$

and

$$\begin{aligned} \frac{\partial E(\boldsymbol{\Theta})}{\partial \theta''} &= \sum_{i=1}^n w_i \left(-\frac{1}{2} \text{tr} \left(-2 \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta'} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \right. \right. \\ &\quad \left. \left. + \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial^2 \Omega_i(\boldsymbol{\Theta})}{\partial \theta' \partial \theta''} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} + \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta'} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial^2 \Omega_i(\boldsymbol{\Theta})}{\partial \theta \partial \theta''} \right) \right. \\ &\quad \left. + \frac{\partial \mu(\boldsymbol{\Theta}, \mathbf{X}_i)}{\partial \theta} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i(\boldsymbol{\Theta})^{-1} \frac{\partial \mu(\boldsymbol{\Theta}, \mathbf{X}_i)^\top}{\partial \theta'} \right) \end{aligned}$$

where:

$$\begin{aligned} \frac{\partial A(\boldsymbol{\Theta})}{\partial \theta''} &= \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top \\ \frac{\partial b_i(\boldsymbol{\Theta})}{\partial \theta''} &= \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i^{-1}(\boldsymbol{\Theta}) \\ \frac{\partial B_i(\boldsymbol{\Theta})}{\partial \theta''} &= \frac{\partial^3 \Omega_i(\boldsymbol{\Theta})}{\partial \theta' \partial \theta''} \\ &\quad - 2 \left(\frac{\partial^2 \Omega_i(\boldsymbol{\Theta})}{\partial \theta \partial \theta''} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta'} + \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta'} + \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial^2 \Omega_i(\boldsymbol{\Theta})}{\partial \theta' \partial \theta''} \right) \\ \frac{\partial C(\boldsymbol{\Theta})}{\partial \theta''} &= \sum_{i=1}^n w_i \mathbf{X}_i \Omega_i^{-1}(\boldsymbol{\Theta}) \left(\frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} + \frac{\partial^2 \Omega_i(\boldsymbol{\Theta})}{\partial \theta \partial \theta''} + \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta} \Omega_i^{-1}(\boldsymbol{\Theta}) \frac{\partial \Omega_i(\boldsymbol{\Theta})}{\partial \theta''} \right) \Omega_i^{-1}(\boldsymbol{\Theta}) \mathbf{X}_i^\top \end{aligned}$$

Appendix B Likelihood ratio test with the REML criterion

The blue term of Equation A in the log-likelihood is invariant to re-parameterisation while the red term is not. This means that a re-parametrisation of X into $\tilde{X} = BX$ with B invertible would not change the likelihood when using ML but would decrease the log-likelihood by $\log(|B|)$ when using REML.

Let's take an example:

```
## data(dfL, package = "LMMstar")
dfTest <- dfL
dfTest$glucagon2 <- dfTest$glucagon*2
```

where we multiply one column of the design matrix by 2. As mentioned previously this does not affect the log-likelihood when using ML:

```
eML.lmmUN <- lmm(weight ~ time+glucagon, data = dfTest, repetition = ~time|id, method = "ML")
eML.lmmUN2 <- lmm(weight ~ time+glucagon2, data = dfTest, repetition = ~time|id, method = "ML")
```

```
logLik(eML.lmmUN)
logLik(eML.lmmUN2)
```

```
[1] -218.71
[1] -218.71
```

but it does when using REML:

```
eREML.lmmUN <- lmm(weight ~ time + glucagon, data = dfTest, repetition = ~time|id, method = "REML")
eREML.lmmUN2 <- lmm(weight ~ time + glucagon2, data = dfTest, repetition = ~time|id, method = "REML")
```

```
logLik(eREML.lmmUN)-logLik(eREML.lmmUN2)
log(2)
```

```
[1] 0.6931472
[1] 0.6931472
```

Therefore, when comparing models with different mean effects there is a risk that the difference (or part of it) in log-likelihood is due to a new parametrisation and not only to a difference in model fit. This would typically be the case when adding an interaction where we can have a smaller restricted log-likelihood when considering a more complex model:

```
set.seed(5)
dfTest$ff <- rbinom(NROW(dfTest), size = 1, prob = 0.5)
logLik(lmm(weight ~ time+glucagon, data = dfTest, repetition = ~time|id, method = "REML"))
logLik(lmm(weight ~ time+glucagon*ff, data = dfTest, repetition = ~time|id, method = "REML"))
```

```
[1] -216.3189
[1] -216.8425
```

This is quite counter-intuitive as more complex model should lead to better fit and would never happen when using ML:

```
logLik(lmm(weight ~ time + glucagon, data = dfTest, repetition = ~time|id, method = "ML"))  
logLik(lmm(weight ~ time + glucagon*ff, data = dfTest, repetition = ~time|id, method = "ML"))
```

```
[1] -218.71
```

```
[1] -218.6259
```

This is why, unless one knows what he/she is doing, it is not recommended to use likelihood ratio test to assess relevance of mean parameters in mixed models estimated with REML.

Appendix C Sum of squares in a linear mixed model

All mixed models implemented in LMMstar can be written as:

$$Y_{it} = X_{it}\beta + \varepsilon_{it} \text{ where } \varepsilon_i \sim \mathcal{N}(0, \Omega)$$

where Y denote the outcome repeatedly measured within each cluster i where t indexes the repetitions. X denotes the covariates, β the mean parameters, ε the residuals, and Ω the residual variance-covariance matrix. Ω must be positive definite so there must exist a square positive definite matrix $\Omega^{1/2}$ such that $\Omega^{1/2}\Omega^{1/2} = \Omega$. Therefore the previous model is equivalent to:

$$Y_{it}^* = X_{it}^*\beta + \varepsilon_{it}^* \text{ where } \varepsilon_i \sim \mathcal{N}(0, I_T)$$

where $Y_i^* = \Omega^{-1/2}Y_i$, $X_i^* = \Omega^{-1/2}X_i$, $\varepsilon_i^* = \Omega^{-1/2}\varepsilon_i$, and I_x is the identity matrix with x rows and columns. One can then introduce the projectors $H = X(X^\top\Omega^{-1}X)^{-1}X^\top\Omega^{-1}$ and $H^* = X^*(X^{*\top}X^*)^{-1}X^{*\top}$ onto the space spanned by X and X^* respectively. We can now define the "normalized" residual sum of squares as the squared sum of the normalized residuals:

$$\begin{aligned} SSE^* &= \varepsilon^{*\top}\varepsilon^* = Y^{*\top}(I_{nT} - H^*)Y^* \\ &= Y^\top\Omega^{-1}Y - Y^\top\Omega^{-1}X(X^\top\Omega^{-1}X)^{-1}X^\top\Omega^{-1}Y \\ &= Y^\top(I_{nT} - H^\top)\Omega^{-1}(I_{nT} - H)Y \end{aligned}$$

The previous to last line uses that: $(I_{nT} - H^\top)\Omega^{-1}(I_{nT} - H) = \Omega^{-1} - H^\top\Omega^{-1} - \Omega^{-1}H + H^\top\Omega^{-1}H = \Omega^{-1} - H^\top\Omega^{-1}$ as $H^\top\Omega^{-1}H = \Omega^{-1}HH = \Omega^{-1}H$ since H is a projector. Note that compared to the "traditional" SSE defined for linear regression and random effect models (e.g. see [Christensen \(2011\)](#) section 2.7), $SSE = \delta SSE^*$ where δ is the residual variance conditional on any random effects, i.e. SSE^* are the residual degrees of freedom. This is because the same definition for the sum of squares is used except that $\varepsilon_i \sim \mathcal{N}(0, \delta\Omega)$.

We can also define the "normalized" regression sum of squares:

$$\begin{aligned} SSR^* &= (X^*\beta)^\top X^*\beta = (H^*Y^*)^\top H^*Y^* = Y^{*\top}H^*Y^* \\ &= Y^\top H^\top\Omega^{-1}Y^* = Y^\top H^\top H^\top\Omega^{-1}Y^* = Y^\top H^\top\Omega^{-1}HY^* \\ &= \hat{\beta}X^\top\Omega^{-1}X\hat{\beta} \end{aligned}$$

where $\hat{\beta} = (X^\top\Omega^{-1}X)^{-1}X^\top\Omega^{-1}Y$. Note that when using the expected information $SSR^* = \hat{\beta}\Sigma_{\hat{\beta}}^{-1}\hat{\beta}$, i.e. it is the F-statistics times the number of parameters. Again the "traditional" SSR defined for linear regression and random effect models is proportional to this normalized SSR: $SSR = \delta SSR^*$.

The proportion of explained variance of p parameters can thus be re-expressed as:

$$R^2 = \frac{SSR}{SSR + SSE} = \frac{SSR^*}{SSR^* + SSE^*} = \frac{Fp}{Fp + df}$$

where df denotes the residual degrees of freedom, typically $n - p$ in a univariate linear model fitted with n observations.

⚠ In practice df is estimated using the Satterthwaite approximation of the degrees of freedom of the regression coefficient. This is only equivalent to the "SSR/SSE" formula in univariate linear regression.

Illustration for a univariate linear model:

Data without missing values:

```
df.aov <- dfL[!is.na(dfL$glucagon),]
```

Traditional anova decomposition:

```
e.lm <- lm(weight ~ time + glucagon, data = df.aov)
car::Anova(e.lm, type = "II")
```

Anova Table (Type II tests)

```
Response: weight
          Sum Sq Df F value    Pr(>F)
time      6367.3  3  6.4308 0.0006329 ***
glucagon   1964.8  1  5.9531 0.0171207 *
Residuals 24093.1 73
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Fit lmm:

```
e.lmm <- lmm(weight ~ time + glucagon, data = df.aov)
```

Residual sum of squares (SSE):

```
SSEstar <- crossprod(residuals(e.lmm, type = "normalized"))
c(SSEstar = SSEstar, SSE = SSEstar * sigma(e.lmm))
```

```
SSEstar      SSE
73.00 24093.11
```

The normalized SSE can also be obtained using the `df.residual` method:

```
df.residual(e.lmm)
```

```
[1] 73
```

Regression sum of squares (SSR):

```
eBeta.lmm <- coef(e.lmm)
eVcov.lmm <- vcov(e.lmm, type.information = "expected")

SSRstar.glucagon <- eBeta.lmm[5] %*% solve(eVcov.lmm[5,5]) %*% eBeta.lmm[5]
SSRstar.time <- eBeta.lmm[2:4] %*% solve(eVcov.lmm[2:4,2:4]) %*% eBeta.lmm[2:4]
c(SSR.glucagon = SSRstar.glucagon * sigma(e.lmm),
  SSR.time = SSRstar.time * sigma(e.lmm),
  F.glucagon = SSRstar.glucagon,
  F.time = SSRstar.time/3)
```

SSR.glucagon	SSR.time	F.glucagon	F.time
1964.764452	6367.324429	5.953062	6.430810

So the proportion of explained variance is:

```
R2.glucagon <- SSRstar.glucagon/(SSRstar.glucagon+SSEstar)
R2.glucagon
```

```
      [,1]
[1,] 0.07540002
```

and the corresponding partial correlation is:

```
sign(coef(e.lmm)["glucagon"])*sqrt(R2.glucagon)
```

```
      [,1]
[1,] -0.2745906
```

which matches the output of `partialCor`:

```
summary(partialCor(e.lmm, R2 = TRUE))
```

Partial correlation

	estimate	se	df	lower	upper	p.value
timeB1w	-0.153	0.113	73	-0.378	0.072	0.1796
timeA1w	-0.038	0.117	73	-0.27	0.195	0.7475
timeA3m	-0.413	0.088	73	-0.589	-0.236	1.36e-05
glucagon	-0.275	0.104	73	-0.482	-0.067	0.0102

Columns lower and upper contain 95% pointwise confidence intervals for each coefficient. Degrees of freedom were computed using a Satterthwaite approximation (column df).

Coefficient of determination (R2)

	estimate	se	df	lower	upper	p.value
time	0.209	0.075	73	0.059	0.359	0.006976
glucagon	0.075	0.057	73	-0.038	0.189	0.191156
global	0.285	0.076	73	0.134	0.435	0.000328

Columns lower and upper contain 95% pointwise confidence intervals for each coefficient. Degrees of freedom were computed using a Satterthwaite approximation (column df).

Appendix D Equivalence with other R packages

D.1 nlme package

The model class obtained with the `lmm` function overlaps the model class of the `lme` and `gls` functions from the `nlme` package.

```
library(nlme)
```

For instance, the compound symmetry is equivalent to `corCompSymm` correlation structure, or to a random intercept model (when the within subject correlation is positive):

```
eCS.gls <- gls(weight ~ time + glucagon, correlation = corCompSymm(form=~time|id),
              data = dfL, na.action = na.omit)
eCS.lme <- lme(weight ~ time + glucagon, random = ~1|id,
              data = dfL, na.action = na.omit)
logLik(eCS.lme)
logLik(eCS.gls)
logLik(eCS.lmm)
```

```
'log Lik.' -243.6005 (df=7)
'log Lik.' -243.6005 (df=7)
[1] -243.6005
```

The estimated random effect also match:

```
range(ranef(eCS.lmm)$estimate-ranef(eCS.lme))
```

```
[1] -3.136991e-08  2.384372e-08
```

Unstructured residual covariance matrix can also be obtained with `gls`:

```
eUN.gls <- gls(weight ~ time + glucagon,
              correlation = corSymm(form=~as.numeric(time)|id),
              weights = varIdent(form=~1|time),
              data = dfL, na.action = na.omit)
logLik(eUN.gls)
logLik(eUN.lmm)
```

```
'log Lik.' -216.3189 (df=15)
[1] -216.3189
```

D.2 lme4 package

The model class obtained with the `lmm` function overlaps the model class of the `lmer` function from the `lme4` package.

```
library(lme4)
library(lmerTest)
```

For instance, the compound symmetry is equivalent to a random intercept model (when the within subject correlation is positive):

```
eRI.lmer <- lmer(weight ~ time + glucagon + (1|id),
               data = dfL)
logLik(eRI.lmer)
logLik(eRI.lmm)
```

```
'log Lik.' -243.6005 (df=7)
[1] -243.6005
```

The estimated random effects match:

```
range(ranef(eRI.lmm)$estimate-ranef(eRI.lmer)$id)
```

```
[1] -3.167867e-08  2.406756e-08
```

Nested random effects correspond to block unstructured:

```
eNRI.lmer <- lmer(weight ~ time*group + (1|id/baseline),
               data = dfL)
logLik(eNRI.lmer)
logLik(eNRI.lmm)
```

```
'log Lik.' -230.5328 (df=11)
[1] -230.5328
```

And the estimated random effects still match:

```
eRanefNRI.lmm <- ranef(eNRI.lmm)
eRanefNRI.lmer <- ranef(eNRI.lmer)
## id
range(eRanefNRI.lmm[eRanefNRI.lmm$variable=="id","estimate"]-eRanefNRI.lmer$id)
## baseline
range(eRanefNRI.lmm[eRanefNRI.lmm$variable!="id","estimate"]-ranef(eNRI.lmer)$'baseline:id')
```

```
[1] -7.457484e-05  1.182242e-04
[1] -0.0001493705  0.0001080902
```

An unstructure residual covariance matrix can also be obtained using random slopes:

```
eUN.lmer <- lmer(weight ~ time + glucagon + (0 + time|id),
                 data = dfL, control = lmerControl(check.nobs.vs.nRE = "ignore"))
logLik(eUN.lmer)
logLik(eUN.lmm)
```

```
'log Lik.' -216.3189 (df=16)
[1] -216.3189
```

Note that however the uncertainty is quantified in a slightly different way, e.g.:

```
anova(eUN.lmm)
```

Multivariate Wald test

	F-statistic	df	p.value
mean: time	86.743 (3,19.0)	2.84e-11	***
: glucagon	13.518 (1,13.7)	0.00257	**

do not match

```
anova(eUN.lmer)
```

Type III Analysis of Variance Table with Satterthwaite's method

	Sum Sq	Mean Sq	NumDF	DenDF	F value	Pr(>F)
time	114.275	38.092	3	20.483	87.242	7.784e-12 ***
glucagon	10.125	10.125	1	16.784	23.191	0.0001671 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

I think this is because lmer base uncertainty computation on the expected information (instead of the observed information). Doing so leads to more similar results:

```
eUN2.lmm <- lmm(weight ~ time + glucagon, repetition = ~time|id,
                structure = "UN", data = dfL, type.information = "expected")
suppressWarnings(anova(eUN2.lmm))
```

Multivariate Wald test

	F-statistic	df	p.value
mean: time	87.253 (3,22.5)	1.48e-12	***
: glucagon	23.198 (1,19.4)	0.000114	***

It is also possible to fit cross-random effects such as:

```
data("Penicillin")
eCRI.lmer <- lmer(diameter ~ 1 + (1|plate) + (1|sample), Penicillin)
logLik(eCRI.lmer)
```

```
'log Lik.' -165.4303 (df=4)
```

using lmm:

```
Penicillin$index <- paste(Penicillin$sample, Penicillin$plate, sep=".")
Penicillin$id <- 1

eCRI.lmm <- lmm(diameter ~ 1 + (1|plate) + (1|sample), data = Penicillin)
logLik(eCRI.lmm)
```

```
[1] -165.4303
```

Despite being significantly slower, the loglikelihood and random effect still match:

```
range(ranef(eCRI.lmm)$estimate-rbind(ranef(eCRI.lmer)$plate, ranef(eCRI.lmer)$sample))
```

```
[1] -4.381305e-07 6.017161e-07
```

D.3 mmrm package

The package `mmrm` is an alternative implementation of mixed models specified via covariance structures:

```
library(mmrm)
e.mmrm <- mmrm(
  formula = FEV1 ~ RACE + SEX + ARMCD * AVISIT + us(AVISIT | USUBJID),
  data = fev_data
)
```

`mmrm()` registered as `emmeans` extension

It leads nearly identical results compared to `lmm`:

```
e.lmm <- lmm(
  formula = FEV1 ~ RACE + SEX + ARMCD * AVISIT,
  repetition = ~ AVISIT | USUBJID, structure = "UN",
  data = fev_data, type.information = "expected"
)
```

Advarselsbesked:

```
I .lmmNormalizeData(as.data.frame(data)[unique(stats::na.omit(var.all))], :
  3 clusters have been removed.
```

```
logLik(e.mmrm) - logLik(e.lmm)
range(coef(e.mmrm) - coef(e.lmm))
range(vcov(e.mmrm) - vcov(e.lmm))
```

```
[1] -2.541278e-06
[1] -0.0001830095  0.0001626755
[1] -0.0003971008  0.0002047941
```

The main differences are:

- `mmrm` uses the expected information matrix to quantify uncertainty instead of the observed information matrix.
- `mmrm` implements the Kenward and Roger method for computing the degrees of freedom and not only the Satterthwaite approximation
- `mmrm` implements different covariance patterns
- `mmrm` is faster and probably more memory efficient
- `mmrm` has currently fewer post-processing methods (e.g. adjustment multiple comparisons when testing several model parameters). This being said, the latest version of the package (0.3.7) included several additional extractor of model feature so this may be improved in the future.

D.4 effectsize package (R^2 or η^2)

Partial η^2 can be computed based on `lmer` using the `effectsize` package:

```
library(effectsize)
eta_squared(eCS.lmer)
cat("\n")
```

```
# Effect Size for ANOVA (Type III)
```

Parameter	Eta2 (partial)	95% CI
time	0.92	[0.89, 1.00]
glucagon	0.03	[0.00, 1.00]

```
- One-sided CIs: upper bound fixed at [1.00].>
```

and are approximately equal to what one can compute "manually":

```
eCS.Wald <- anova(eCS.lmm)$multivariate
eCS.Wald$df.num*eCS.Wald$statistic/(eCS.Wald$df.num*eCS.Wald$statistic+eCS.Wald$df.denom)
```

```
[1] 0.92380363 0.03162017
```

The will not be true for heteroschedastic models:

```
eUN.Wald <- anova(eUN.lmm)$multivariate
eUN.Wald$df.num*eUN.Wald$statistic/(eUN.Wald$df.num*eUN.Wald$statistic+eUN.Wald$df.denom)
```

```
[1] 0.9319379 0.4965135
```

compared to:

```
eta_squared(eUN.lmer)
cat("\n")
```

```
# Effect Size for ANOVA (Type III)
```

```
Parameter | Eta2 (partial) |      95% CI
-----|-----|-----
time      | 0.93 | [0.87, 1.00]
glucagon  | 0.58 | [0.29, 1.00]
```

```
- One-sided CIs: upper bound fixed at [1.00].>
```

But in that case both may be misleading as the proportion of explained variance is not clearly defined.

D.5 MuMIn package (R^2)

```
library(MuMIn)
r.squaredGLMM(eCS.lmer)
cat("\n")
```

```
           R2m      R2c
[1,] 0.2163302 0.9764382
```

To reproduce these R^2 , we extract from the random intercept model:

- the residual variance

```
sigmaW <- sigma(eCS.lmm)[1,1]-sigma(eCS.lmm)[1,2]
```

- the variance of the random effect

```
sigmaB <- sigma(eCS.lmm)[1,2]
```

- the variance of the fitted values:

```
sigma2_XB <- var(fitted(eCS.lmm))
```

and evaluate the ratios:

```
c(R2m = sigma2_XB/(sigmaW + sigmaB + sigma2_XB),
  R2c = (sigma2_XB + sigmaB)/(sigmaW + sigmaB + sigma2_XB))
```

```
           R2m      R2c
0.2163302 0.9764382
```


D.6 stats package (partial residuals)

The function `residuals.lm` can be used to extract partial residuals from `lm` objects:

```
eIID.lm <- lm(weight ~ time + glucagon, data = dfL)
pRes.lm <- residuals(eIID.lm, type = "partial")
head(pRes.lm)
```

```
      time  glucagon
1  3.359543  1.475108
2 49.419060 39.475108
3 -8.145299 -16.024892
4 24.241798 20.475108
5 -8.407618 -12.624892
6 41.027985 33.075108
```

Those generally differ (by a constant) from the one provided by `residuals.lmm`:

```
eIID.lmm <- lmm(weight ~ time + glucagon, data = dfL)
head(residuals(eIID.lmm, type = "partial", var = "glucagon"))
```

```
[1] -31.9349871  6.0650129 -49.4349871 -12.9349871 -46.0349871 -0.3349871
```

Indeed, `residuals.lm` centers the design matrix of the variable relative to which the partial residuals are computed:

```
m.pres2 <- dfL$weight - cbind(model.matrix(~time,dfL), mean(dfL$glucagon)) %*% coef(eIID.lmm)
range(pRes.lm[, "glucagon"] - m.pres2, na.rm = TRUE)
```

```
[1] -3.348433e-13  4.654055e-13
```

For continuous variable with a linear effect, these residuals can be obtained by setting the `type` argument to `"partial-center"`:

```
eIID.lmm <- lmm(weight ~ time + glucagon, data = dfL)
pRes.lmm <- residuals(eIID.lmm, type = "partial-center", var = "glucagon")
range(pRes.lm[, "glucagon"] - pRes.lmm)
```

```
[1] -3.330669e-13  4.725109e-13
```

⚠ When evaluating the partial residuals relative to categorical variables, interactions, or non-linear terms, the output obtained with `partial-center` will not match the one of `residuals.lm`. Indeed `partial-center` will, when numeric, center the original variable whereas `residuals.lm` will center the column relative to the coefficient in the design matrix.