

# Package ‘CFAssay’

April 15, 2019

**Type** Package

**Title** Statistical analysis for the Colony Formation Assay

**Version** 1.16.1

**Date** 2017-02-10

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**Depends** R (>= 2.10.0)

**Description** The package provides functions for calculation of linear-quadratic cell survival curves and for ANOVA of experimental 2-way designs along with the colony formation assay.

**License** LGPL

**biocViews** ImmunoOncology, CellBasedAssays, CellBiology, Regression, Survival

**git\_url** <https://git.bioconductor.org/packages/CFAssay>

**git\_branch** RELEASE\_3\_8

**git\_last\_commit** 9d0521b

**git\_last\_commit\_date** 2019-01-04

**Date/Publication** 2019-04-15

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 cellsurvLQdiff

*Comparison of two linear-quadratic cell survival curves*


---

### Description

The function does an ANOVA test for overall comparison of the parameters *alpha* and *beta* of two linear-quadratic cell survival curves. The parameters are fitted simultaneously to the data with this function, i.e. no other function is necessary to derive the fits.

### Usage

```
cellsurvLQdiff(X, curvevar, method="ml", PEmethod="fit")
```

### Arguments

<code>X</code>	A data frame which contains columns <code>Exp</code> , <code>dose</code> , <code>ncells</code> , <code>ncolonies</code> and a further column containing two different values (character strings), which identify the two curves. Moreover, if there is no 0-value in the <code>dose</code> -column, <code>X</code> has to contain a column <code>pe</code> for plating efficiencies.
<code>curvevar</code>	Character string, which has to be one of the column names of the data frame <code>X</code> , that contains the two different values (character strings that distinguishes between the two curves).
<code>method</code>	Determines the method used for the fit. "ml" is for maximum-likelihood, "ls" for least-squares. "franken" performs weighed least-squares with weights as described in Franken et al. (2006).
<code>PEmethod</code>	Controls the value of the plating efficiencies, i.e. the colony counts for untreated cells. "fit" calculates fitted plating efficiencies as model parameters, "fix" uses fixed ones calculated from the observed zero dose data or from a column named <code>pe</code> in <code>X</code> .

### Details

In the data frame `X`, `Exp` identifies the experimental replicates and may be numeric or non-numeric. `method="ml"` for maximum-likelihood uses R function `glm` with family "quasipoisson" and link function "log". `method="ls"` uses R function `lm`.

### Value

The function returns an object of class `cellsurvLQdiff` containing three elements, `fit1`, `fit2` and `anv`. `fit1` and `fit2` are objects of class `glm` when `method="ml"` or of class `lm` when `method="ls"`. `fit1` has parameters `alpha` and `beta` fitted in common for both cell survival curves. `fit2` has parameters `alpha` and `beta` fitted differently for both curves. `anv` is of class `anova` and contains the F-test. Test results are printed, however, the full result including curve parameters is returned invisibly, i.e. the function has to be used with `print` or assigned to a variable, say for e.g. `fitcomp` as in the example below.

### Author(s)

Herbert Braselmann

**See Also**

[glm](#) and [family](#) with references for generalized linear modelling. [anova](#), [cellsurvLQfit](#).

**Examples**

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\t")
names(datatab) #contains a column "cline"
table(datatab$cline)
fitcomp<- cellsurvLQdiff(datatab, curvevar="cline") #using default options
print(fitcomp)
plot(cellsurvLQfit(subset(datatab, cline=="okf6TERT1")), col=1)
plot(cellsurvLQfit(subset(datatab, cline=="cal33")), col=2, add=TRUE)
legend(0, 0.02, c("okf6TERT1", "cal33"), text.col=1:2)
#using different options:
print(cellsurvLQdiff(datatab, curvevar="cline", method="ls"))
print(cellsurvLQdiff(datatab, curvevar="cline", PEmethod="fix"))
print(cellsurvLQdiff(datatab, curvevar="cline", method="ls", PEmethod="fix"))
print(cellsurvLQdiff(datatab, curvevar="cline", method="franken"))
```

---

 cellsurvLQfit

*Fit the linear-quadratic (LQ) model to cell survival data*


---

**Description**

This function calculates the linear coefficient *alpha* and the coefficient *beta* of the dose-squared term (see manual for this R-package) for colony counts measured for a set of irradiation doses and repeated experiments. The function is a wrapper for the R-functions `glm` or `lm`, which simplifies use of these functions for cell survival data.

**Usage**

```
cellsurvLQfit(X, method="ml", PEmethod="fit")
```

**Arguments**

<code>X</code>	A data frame which contains at least columns <code>Exp</code> , <code>dose</code> , <code>ncells</code> , <code>ncolonies</code> and if there is no 0-value in the dose-column, <code>X</code> has to contain a further column <code>pe</code> for plating efficiencies.
<code>method</code>	Determines the method used for the fit. <code>"ml"</code> is for maximum-likelihood, <code>"ls"</code> for least-squares. <code>"franken"</code> performs weighed least-squares with weights as described in Franken et al. (2006).
<code>PEmethod</code>	Controls the value of the plating efficiencies, i.e. the colony counts for untreated cells. <code>"fit"</code> calculates fitted plating efficiencies as model parameters, <code>"fix"</code> uses fixed ones calculated from the observed zero dose data or from a column named <code>pe</code> in <code>X</code> .

## Details

In the data frame *X*, *Exp* identifies the experimental replicates and may be numeric or non-numeric. `method="ml"` uses R function `glm` with quasipoisson family and link function "log". `method="ls"` uses R function `lm`. `PEmethod="fit"` fits plating efficiencies for every experiments. `PEmethod="fix"` uses observed plating efficiencies. If there is no 0-value in the dose-column, `PEmethod` is overwritten with "fix" and *X* has to contain a further column `pe` containing the plating efficiencies, i.e. `ncolonies/ncells` from untreated cells, not per hundred or percent.

## Value

The function returns an object of class `cellsurvLQfit`, which is similar to classes `glm` or `lm`, however containing two additional entries, `type` and `PEmethod`, which are used for printing and plotting. The full result is returned invisibly, i.e. the function has to be used with `print` or `plot` or assigned to a variable, say for e.g. `fit` as in the example below.

## Author(s)

Herbert Braselmann

## References

Franken NAP, Rodermond HM, Stap J, et al. Clonogenic assay of cells in vitro. *Nature Protoc* 2006;1:2315-19.

## See Also

`glm` and `family` with references for generalized linear modelling, `lm`

## Examples

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\t")
X<- subset(datatab, cline=="okf6TERT1")
fit<- cellsurvLQfit(X) #using default options
print(fit)
print(fit$type)
print(fit$PEmethod)
#Using other options:
print(cellsurvLQfit(X, method="ls"))
print(cellsurvLQfit(X, PEmethod="fix"))
print(cellsurvLQfit(X, method="ls", PEmethod="fix"))
print(cellsurvLQfit(X, method="franken"))
```

## Description

The function does an ANOVA of cell survival data from experimental 2-way designs where a treatment factor is tested on a control and on an altered cell line or where two different simultaneous treatments are tested on cells from a common unaltered clone. The function is a wrapper for the R-function `glm`. quasipoisson family is used with link function "log", i.e. dependency of treatment factors is considered as logarithmically additive.

**Usage**

```
cfa2way(X, A, B, param="A/B", method="ml")
```

**Arguments**

X	a data frame which contains columns Exp, ncells, ncolonies and two further columns for the treatment variables, see details.
A	a character string containing the name of a treatment or cell line variable (first factor in the model)
B	a character string containing the name of a treatment or cell line variable (second factor in the model)
param	Controls the parametrization of the model. Options are "A/B" for B nested in A, "B/A" for A nested in B and "A*B" for interaction term.
method	determines the method used for the fit. "ml" is for maximum-likelihood, "ls" for least-squares.

**Details**

In the data frame X, Exp identifies the experimental replicates and may be numeric or non-numeric. The two treatment or cell line columns should have numeric values 0, 1, ... for 2, 3, ... levels. For e.g. if a column describes clonal alteration (transfection, knock-down etc.) by a gene then 0 means unaltered or control and 1 means altered. Similar if a column describes treatment with one dose then 0 means untreated and 1 treated. 2 would indicate another dose level from the same treatment drug without taking it as a continuous covariate as for cell survival curves for radiation.

**Value**

The function returns an object of class cfa2way containing three elements, fit1, fit2 and anv. fit1 and fit2 are objects of class glm when method="ml" or of class lm when method="ls". fit1 has logarithmic additive parameters without interaction. fit2 has logarithmic additive parameters and interaction. anv is of class anova and contains the F-test. The full result is returned invisibly, i.e. the function has to be used with print or assigned to a variable, say for e.g. fitcomp as in the example below.

**Author(s)**

Herbert Braselmann

**See Also**

[glm](#) and [family](#) with references for generalized linear modelling.

**Examples**

```
datatab<- read.table(system.file("doc", "exp2_2waycfa.txt", package="CFAssay"), header=TRUE, sep="\t")
names(datatab) # has columns "x5fuCis" and "siRNA"
fitcomp<- cfa2way(datatab, A="siRNA", B="x5fuCis", param="A/B")
print(fitcomp, labels=c(A="siRNA",B="x5fuCis"))
print(cfa2way(datatab, A="siRNA", B="x5fuCis", param="A/B", method="ls"))
```

---

pes	<i>Calculation of plating efficiencies from a curve data set containing one specified curve</i>
-----	---

---

### Description

The function calculates plating efficiencies, i.e. fractions of colonies per cell of untreated cells, for every experimental replicate in a data frame with one specified curve. For that, lines with zero dose ( $dose = 0$ ) are extracted from the data frame.

### Usage

```
pes(X)
```

### Arguments

X                    A data frame which contains columns Exp, dose, ncells, ncolonies.

### Details

In the data frame X, Exp identifies the experimental replicates and may be numeric or non-numeric.

### Value

The function returns a data frame with three columns Exp, pe and S0, containing experiment identifiers (biological replicates), measured plating efficiencies and plating efficiencies fitted separately for each repeated experiments. Rows of the data frame are named for the different experiments.

### Author(s)

Herbert Braselmann

### Examples

```
datatab <- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")
X <- subset(datatab, cline=="okf6TERT1") #Specification of curve
pes(X)
pes(subset(datatab, cline=="cal33") )
```

---

plot.cellsurvLQfit	<i>Plot of an LQ model fit</i>
--------------------	--------------------------------

---

### Description

This function plots a cell survival curve derived from fitting an LQ model with function cellsurvLQfit

### Usage

```
## S3 method for class 'cellsurvLQfit'
plot(x, xlim = NULL, ylim = c(0.008, 1), xlab = "Dose (Gy)", ylab = "Survival (1 = 100%)", col=1, pch=1)
```

**Arguments**

x	an object of class <code>cellsurvLQfit</code> resulting from function <code>cellsurvLQfit</code> .
xlim	plot range for the x-axis. Default is the dose range of the data.
ylim	plot range for the y-axis. Default is from 0.008 to 1.0
xlab	label for the x-axis. Default is "Dose (Gy)".
ylab	label for the y-axis. Default is "Survival (1 = 100%)".
col	colour for plot. Default is <code>col = 1</code> .
pch	symbol for plotting points. Default is <code>pch = 1</code> .
add	logical; if TRUE add to an already existing plot, see <code>curve</code> .
...	further arguments to pass to R function <code>plot</code> .

**Author(s)**

Herbert Braselmann

**See Also**

`cellsurvLQfit`

**Examples**

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")
X<- subset(datatab, cline=="okf6TERT1")
fit<- cellsurvLQfit(X)
plot(fit)
S0 <- pes(X)$S0
names(S0) <- pes(X)$Exp
sfpmean(X, S0) #values of plotted mean survival fractions and error bars
# add second plot
plot(cellsurvLQfit(subset(datatab, cline=="cal33")), col=2, add=TRUE)
```

---

plotExp

*Generic plotting of experimental repeats*

---

**Description**

Generic plotting of experimental repeats of cell survival data in separated plots. `plotExp` methods are defined for objects resulting from `CFAssay` functions `cellsurvLQfit` and `cfa2way`.

**Usage**

```
plotExp(x, ...)
```

**Arguments**

x	should be an object of class <code>cellsurvLQfit</code> or <code>cfa2way</code> , see details.
...	further arguments to pass to generic function <code>plotExp</code> , dependent of the class of the argument <code>x</code> .

**Details**

For other objects than of class `cellsurvLQfit` or `cfa2way`, `plot.default` will be called. In this version of `CFAssay` this will give an error message and a hint to use one of the two defined classes.

**Author(s)**

Herbert Braselmann

**See Also**

[plotExp.cellsurvLQfit](#) and [plotExp.cfa2way](#)

**Examples**

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")
X<- subset(datatab, cline=="okf6TERT1")
fit<- cellsurvLQfit(X)
#pdf("okf6TERT1_experimental_plots.pdf")
  plotExp(fit)
#dev.off()
## Not run:
x <- 1
plotExp(X) #yields an error for this data type

## End(Not run)
```

---

`plotExp.cellsurvLQfit` *Diagnostic plots of LQ model versus experiments*

---

**Description**

This function plots the fit of an LQ model versus the fits of each experiment in a series of plots. It is recommended to direct it in a pdf-file.

**Usage**

```
## S3 method for class 'cellsurvLQfit'
plotExp(x, xlim = NULL, ylim = c(0.001, 1.5), xlab = "Dose (Gy)", ylab = "Survival (1 = 100%)", ...)
```

**Arguments**

<code>x</code>	an object of class <code>cellsurvLQfit</code> resulting from function <a href="#">cellsurvLQfit</a> .
<code>xlim</code>	range of x axis (dose).
<code>ylim</code>	range of y-axis.
<code>xlab</code>	label of x-axis.
<code>ylab</code>	label of y-axis.
<code>...</code>	further arguments to pass to generic function <code>plotExp</code> .

**Author(s)**

Herbert Braselmann



**See Also**

cellsurvLQfit

**Examples**

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\t")
X<- subset(datatab, cline=="okf6TERT1")
fit<- cellsurvLQfit(X)
print(fit)
#pdf("okf6TERT1_experimental_plots.pdf")
  plotExp(fit)
#dev.off()
```

---

plotExp.cfa2way

*Diagnostic plot of experimental replicates for two-way analysis of cell survival data*

---

**Description**

This function plots the estimated means of a two-way ANOVA for each experiment in a series of plots. It is recommended to direct it in a pdf-file.

**Usage**

```
## S3 method for class 'cfa2way'
plotExp(x, labels=c(A="A",B="B"), ...)
```

**Arguments**

`x` an R object of class `cfa2way` and resulting from function `cfa2way`.  
`labels` Labels for output description. These describe the meaning of A and of B.  
`...` further arguments to pass to generic function `plotExp`.

**Author(s)**

Herbert Braselmann

**See Also**

[cfa2way](#)

**Examples**

```
datatab<- read.table(system.file("doc", "exp2_2waycfa.txt", package="CFAssay"), header=TRUE, sep="\t")
names(datatab) # has columns "x5fuCis" and "siRNA"
fitcomp<- cfa2way(datatab, A="siRNA", B="x5fuCis", param="A/B")
print(fitcomp, labels=c(A="siRNA",B="x5fuCis"))
pdf("TwoWay_experimental_plots.pdf")
  plotExp(fitcomp, labels=c(A="siRNA", B="x5fuCis"))
dev.off()
```

---

plotExp.default	<i>Default function for plotting of experimental repeats</i>
-----------------	--

---

## Description

plotExp.default is the formal default method of the generic `plotExp` function.

## Usage

```
## Default S3 method:  
plotExp(x, ...)
```

## Arguments

<code>x</code>	should be an object of class <code>cellsurvLQfit</code> or <code>cfa2way</code> , see details.
<code>...</code>	further arguments to pass to generic function <code>plotExp</code> , dependent of the class of the argument <code>x</code> .

## Details

In this version of `CFAssay` other objects than of class `cellsurvLQfit` or `cfa2way` this will give an error message and a hint to use one of the two defined classes.

## Author(s)

Herbert Braselmann

## See Also

[plotExp.cellsurvLQfit](#) and [plotExp.cfa2way](#)

## Examples

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")  
X<- subset(datatab, cline=="okf6TERT1")  
fit<- cellsurvLQfit(X)  
#pdf("okf6TERT1_experimental_plots.pdf")  
  plotExp(fit)  
#dev.off()  
## Not run:  
x <- 1  
plotExp(X) #yields an error for this data type  
  
## End(Not run)
```

---

```
print.cellsurvLQdiff Print test results of comparison of two linear-quadratic cell survival curves
```

---

### Description

The function prints linear coefficients alpha and dose-squared coefficients beta of calculated with function `cellsurvLQdiff`. In addition quality statistics of the fit and ANOVA F-test results for overall comparison of the coefficients are printed.

### Usage

```
## S3 method for class 'cellsurvLQdiff'
print(x, ...)
```

### Arguments

`x` an object of class `cellsurvLQdiff` resulting from function `cellsurvLQdiff`.  
`...` further arguments to pass to R function `print`.

### Author(s)

Herbert Braselmann

### See Also

[cellsurvLQdiff](#)

### Examples

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")
fitcomp<- cellsurvLQdiff(datatab, curvevar="cline") #using default options
print(fitcomp)
#using different options for cellsurvLQdiff:
print(cellsurvLQdiff(datatab, curvevar="cline", method="ls"))
print(cellsurvLQdiff(datatab, curvevar="cline", PEmethod="fix"))
print(cellsurvLQdiff(datatab, curvevar="cline", method="ls", PEmethod="fix"))
print(cellsurvLQdiff(datatab, curvevar="cline", method="franken"))
```

---

```
print.cellsurvLQfit Print summary of an LQ-model fit for cell survival data
```

---

### Description

The function prints the results of an LQ-model fit for radiation dose dependent cell survival.

### Usage

```
## S3 method for class 'cellsurvLQfit'
print(x, ...)
```

**Arguments**

`x` an object of class `cellsurvLQfit` resulting from `cellsurvLQfit`.  
`...` further arguments to pass to R function `print`.

**Details**

In this version of `CFAssay` the class argument `x` is checked by its entry `fit$type` and results in an error, when `x` results from an independent use of `glm` or `lm`.

**Author(s)**

Herbert Braselmann

**See Also**

`cellsurvLQfit`

**Examples**

```
datatab<- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="\n")
X<- subset(datatab, cline=="okf6TERT1")
fit<- cellsurvLQfit(X) #using default options
print(fit)
print(fit$type)
print(fit$PEmethod)
#using other options
print(cellsurvLQfit(X, method="ls"))
print(cellsurvLQfit(X, PEmethod="fix"))
print(cellsurvLQfit(X, method="ls", PEmethod="fix"))
print(cellsurvLQfit(X, method="franken"))
```

---

```
print.cfa2way
```

*Print summary of two-way analysis for cell survival data.*

---

**Description**

The function prints a summary of two-way analysis for cell survival data.

**Usage**

```
## S3 method for class 'cfa2way'
print(x, labels=c(A="A",B="B"), ...)
```

**Arguments**

`x` an R object of class `cfa2way` and resulting from function `cfa2way`.  
`labels` labels for output description. These describe the meaning of A and of B.  
`...` further arguments to pass to R function `print`.

**Author(s)**

Herbert Braselmann

**See Also**[cfa2way](#)**Examples**

```
datatab<- read.table(system.file("doc", "exp2_2waycfa.txt", package="CFAssay"), header=TRUE, sep="\t")
names(datatab) # has columns "x5fuCis" and "siRNA"
fitcomp<- cfa2way(datatab, A="siRNA", B="x5fuCis", param="A/B")
print(fitcomp, labels=c(A="siRNA",B="x5fuCis"))
```

---

sfpmean	<i>Pointwise mean survival fractions for curves with several experimental replicates</i>
---------	--

---

**Description**

The function calculates mean survival fractions for curves averaged over experimental replicates. The function is employed by function `plot.cellsurvLQfit` for plotting observed means

**Usage**

```
sfpmean(X, S0=NULL)
```

**Arguments**

X	A data frame which contains columns <code>Exp</code> , <code>dose</code> , <code>ncells</code> , <code>ncolonies</code> and if <code>S0=NULL</code> , X has to contain a further column <code>pe</code> for plating efficiencies.
S0	If not NULL, a named numerical vector of length equal to the number of different experiments, i.e. <code>length(S0)==length(unique(X\$Exp))</code> has to be TRUE. Default is <code>S0=NULL</code> , i.e. undefined.

**Details**

In the data frame X, `Exp` identifies the experimental replicates and may be numeric or non-numeric. `S0` may contain plating efficiencies for each replicate, resulting from function [pes](#) or from [cellsurvLQfit](#) (fitted). When `S0=NULL`, X must have a column with name `pe`, containing the plating efficiencies.

**Value**

A numerical matrix with two rows, the first row containing the survival fractions for each radiation dose, second row the standard deviations.

**Author(s)**

Herbert Braselmann

**See Also**

[pes](#), [cellsurvLQfit](#), [plot.cellsurvLQfit](#)

**Examples**

```
datatab <- read.table(system.file("doc", "expl1_cellsurvcurves.txt", package="CFAssay"), header=TRUE, sep="")
X <- subset(datatab, cline=="okf6TERT1")
S0 <- pes(X)$pe #observed plating efficiencies
length(S0)==length(unique(X$Exp)) #length ok?
names(S0) <- pes(X)$Exp
sfpmean(X, S0)
fit <- cellsurvLQfit(X)
fit$coef #contains fitted log-pe
grep("Exp",names(fit$coef))
S01 <- exp(fit$coef[1:8]) #fitted pe
sfpmean(X, S01)
## Not run: sfpmean(X) #yields an error for this data set
```

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