

# Package ‘LongCART’

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**Type** Package

**Title** Recursive Partitioning for Longitudinal Profiles Using Baseline Covariates

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**Depends** R (>= 3.4.0), nlme, rpart

**Imports** Formula

**Description** Constructs longitudinal tree (i.e., regression tree with heterogeneous longitudinal profile) for continuous longitudinal outcome using baseline covariates as partitioning variables according to the 'LongCART' algorithm as described in Kundu and Harelak (2019) <doi:10.1080/24709360.2018.1557797>.

**License** GPL (>= 2)

**URL** <https://www.r-project.org>

**BugReports** <https://github.com/madanstat/LongCART/issues>

**NeedsCompilation** no

**Repository** CRAN

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ACTG175	<i>Converted AIDS Clinical Trials Group Study 175 (source: speff2trial package)</i>
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### Description

ACTG 175 was a randomized clinical trial to compare monotherapy with zidovudine or didanosine with combination therapy with zidovudine and didanosine or zidovudine and zalcitabine in adults infected with the human immunodeficiency virus type I whose CD4 T cell counts were between 200 and 500 per cubic millimeter.

### Usage

```
data(ACTG175)
```

### Format

A data frame with 6417 observations from 2139 patients on the following 24 variables.

**pidnum** patient ID number  
**age** age in years at baseline  
**wtkg** weight in kg at baseline  
**hemo** hemophilia (0=no, 1=yes)  
**homo** homosexual activity (0=no, 1=yes)  
**drugs** history of intravenous drug use (0=no, 1=yes)  
**karnof** Karnofsky score (on a scale of 0-100)  
**oprior** non-zidovudine antiretroviral therapy prior to initiation of study treatment (0=no, 1=yes)  
**z30** zidovudine use in the 30 days prior to treatment initiation (0=no, 1=yes)  
**zprior** zidovudine use prior to treatment initiation (0=no, 1=yes)  
**preanti** number of days of previously received antiretroviral therapy  
**race** race (0=white, 1=non-white)  
**gender** gender (0=female, 1=male)  
**str2** antiretroviral history (0=naive, 1=experienced)  
**strat** antiretroviral history stratification (1:antiretroviral naive, 2:greater than 1 but less than 52 weeks of prior antiretroviral therapy, 3: greater than 52 weeks)  
**symptom** symptomatic indicator (0=asymptomatic, 1=symptomatic)  
**treat** treatment indicator (0=zidovudine only, 1=other therapies)  
**offtrt** indicator of off-treatment before 96 weeks (0=no,1=yes)  
**r** missing CD4 T cell count at 96 weeks (0=missing, 1=observed)  
**cens** indicator of observing the event in days

**days** number of days until the first occurrence of: (i) a decline in CD4 T cell count of at least 50 (ii) an event indicating progression to AIDS, or (iii) death.

**arms** treatment arm (0=zidovudine, 1=zidovudine and didanosine, 2=zidovudine and zalcitabine, 3=didanosine)

**time** time in weeks

**cd4** CD4 T cell count

## References

Hammer, S.M., et al. (1996), A trial comparing nucleoside monotherapy with combination therapy in HIV-infected adults with CD4 cell counts from 200 to 500 per cubic millimeter. *New England Journal of Medicine*, 335:1081-1090.

---

LongCART

*Longitudinal CART with continuous response via binary partitioning*

---

## Description

Recursive partitioning for linear mixed effects model with continuous univariate response variables per LonCART algorithm based on baseline partitioning variables (Kundu and Harezlak, 2019).

## Usage

```
LongCART(data, patid, fixed, gvars, tgvars, minsplit=40,
          minbucket=20, alpha=0.05, coef.digits=2, print.lme=F)
```

## Arguments

<code>data</code>	name of the dataset. It must contain variable specified for <code>patid</code> (indicating subject id), all the variables specified in the formula and the baseline partitioning variables.
<code>patid</code>	name of the subject id variable.
<code>fixed</code>	a two-sided linear formula object describing the fixed-effects part of the model, with the response on the left of a <code>~</code> operator and the terms, separated by <code>+</code> operators, on the right. Model with <code>-1</code> to the end of right side indicates no intercept. For model with no fixed effect beyond intercept, please specify only <code>1</code> right to the <code>~</code> operator.
<code>gvars</code>	list of partitioning variables of interest. Value of these variables should not change over time. Regarding categorical variables, only numerically coded categorical variables should be specified. For nominal categorical variables or factors, please first create corresponding dummy variable(s) and then pass through <code>gvars</code> .
<code>tgvars</code>	types (categorical or continuous) of partitioning variables specified in <code>gvar</code> . For each of continuous partitioning variables, specify <code>0</code> and for each of the categorical partitioning variables, specify <code>1</code> . Length of <code>tgvars</code> should match to the length of <code>gvars</code>

<code>minsplit</code>	the minimum number of observations that must exist in a node in order for a split to be attempted.
<code>minbucket</code>	the minimum number of observations in any terminal node.
<code>alpha</code>	alpha (i.e., nominal type I error) level for parameter instability test
<code>coef.digits</code>	decimal points for displaying coefficients in the tree structure.
<code>print.lme</code>	if TRUE, then summary of fitted model from <code>lme()</code> will be printed for each node.

### Details

Construct regression tree based on heterogeneity in linear mixed effects models of following type:  $Y_i(t) = W_i(t)\theta + b_i + \epsilon_{it}$  where  $W_i(t)$  is the design matrix,  $\theta$  is the parameter associated with  $W_i(t)$  and  $b_i$  is the random intercept. Also,  $\epsilon_{it} \sim N(0, \sigma^2)$  and  $b_i \sim N(0, \sigma_u^2)$ .

### Value

<code>Treeout</code>	contains summary information of tree fitting for each terminal nodes and non-terminal nodes. Columns of <code>Treeout</code> include "ID", the (unique) node numbers that follow a binary ordering indexed by node depth, <code>n</code> , the number of observations reaching the node, <code>yval</code> , the fitted model of the response at the node, <code>var</code> , a factor giving the names of the variables used in the split at each, <code>index</code> , the cut-off value of splitting variable for binary partitioning, <code>p</code> (Instability), the p-value for parameter instability test for the splitting variable, <code>loglik</code> , the log-likelihood of the node, <code>improve</code> , the improvement in deviance given by this split, and <code>Terminal</code> , indicator (True or False) of terminal node.
<code>p</code>	number of fixed parameters
<code>AIC.tree</code>	AIC of the tree-structured model
<code>AIC.root</code>	AIC at the root node (i.e., without tree structure)
<code>improve.AIC</code>	improvement in AIC due to tree structure ( <code>AIC.tree - AIC.root</code> )
<code>logLik.tree</code>	log-likelihood of the tree-structured model
<code>logLik.root</code>	log-likelihood at the root node (i.e., without tree structure)
<code>Deviance</code>	$2 * (\logLik.tree - \logLik.root)$
<code>LRT.df</code>	degrees of freedom for likelihood ratio test comparing tree-structured model with the model at root node.
<code>LRT.p</code>	p-value for likelihood ratio test comparing tree-structured model with the model at root node.
<code>subj.class</code>	Assigned node for each individual subjects per fitted longitudinal tree
<code>frame</code>	rpart compatible object
<code>splits</code>	rpart compatible object
<code>cptable</code>	rpart compatible object
<code>functions</code>	rpart compatible object

### Author(s)

Madan Gopal Kundu <madan\_g.kundu@yahoo.com>

## References

Kundu, M. G., and Harezlak, J. (2019). Regression trees for longitudinal data with baseline covariates. *Biostatistics & Epidemiology*, 3(1):1-22.

## See Also

[plotLongCART](#), [textLongCART](#), [StabCat](#), [StabCont](#)

## Examples

```
#--- Get the data
data(ACTG175)

#--- Run LongCART()
gvars=c("age", "gender", "wtkg", "hemo", "homo", "drugs",
        "karnof", "oprior", "z30", "zprior", "race",
        "str2", "symptom", "treat", "offtrt")
tgvars=c(1, 0, 1, 0, 0, 0,
        1, 0, 0, 0, 0,
        0, 0, 0, 0)

out<- LongCART(data=ACTG175, patid="pidnum", fixed=cd4~time,
              gvars=gvars, tgvars=tgvars, alpha=0.05,
              minsplit=100, minbucket=50, coef.digits=2)

#--- Plot tree

par(xpd = T)
plot(out, compress = T)
text(out, use.n = T)
```

---

plotLongCART

*Plot an LongCART Object*

---

## Description

Plots an LongCART object on the current graphics device.

## Usage

```
plotLongCART(x, uniform = FALSE, branch = 1, nspace,
            margin = 0, minbranch = 0.3, ...)
```

**Arguments**

x	a fitted object of class "LongCART", containing a linear mixed effects tree.
uniform	similar to plot.rpart; if TRUE, uniform vertical spacing of the nodes is used; this may be less cluttered when fitting a large plot onto a page. The default is to use a non-uniform spacing proportional to the error in the fit.
branch	similar to plot.rpart; controls the shape of the branches from parent to child node. Any number from 0 to 1 is allowed. A value of 1 gives square shouldered branches, a value of 0 give V shaped branches, with other values being intermediate.
nspace	similar to plot.rpart; the amount of extra space between a node with children and a leaf, as compared to the minimal space between leaves. Applies to compressed trees only. The default is the value of branch.
margin	similar to plot.rpart; an extra fraction of white space to leave around the borders of the tree. (Long labels sometimes get cut off by the default computation).
minbranch	similar to plot.rpart; set the minimum length for a branch to minbranch times the average branch length. This parameter is ignored if uniform=TRUE. Sometimes a split will give very little improvement, or even (in the classification case) no improvement at all. A tree with branch lengths strictly proportional to improvement leaves no room to squeeze in node labels.
...	arguments to be passed to or from other methods.

**Details**

This function is a method for the generic function plot, for objects of class LongCART. The y-coordinate of the top node of the tree will always be 1.

**Value**

The coordinates of the nodes are returned as a list, with components x and y.

**Author(s)**

Madan Gopal Kundu <madan\_g.kundu@yahoo.com>

**References**

Kundu, M. G., and Harezlak, J. (2019). Regression trees for longitudinal data with baseline covariates. *Biostatistics & Epidemiology*, 3(1):1-22.

**See Also**

[textLongCART](#), [LongCART](#), [StabCat](#), [StabCont](#)

**Examples**

```

#--- Get the data
data(ACTG175)

#--- Run LongCART()
gvars=c("age", "gender", "wtkg", "hemo", "homo", "drugs",
        "karnof", "oprior", "z30", "zprior", "race",
        "str2", "symptom", "treat", "offtrt")
tgvars=c(1, 0, 1, 0, 0, 0,
         1, 0, 0, 0, 0,
         0, 0, 0, 0)

out<- LongCART(data=ACTG175, patid="pidnum", fixed=cd4~time,
              gvars=gvars, tgvars=tgvars, alpha=0.05,
              minsplit=100, minbucket=50, coef.digits=2)

#--- Plot tree

par(xpd = T)
plot(out, compress = T)
text(out, use.n = T)

```

StabCat

*parameter stability test for categorical partitioning variable***Description**

Performs parameter stability test (Kundu and Harezlak, 2019) with categorical partitioning variable to determine whether the parameters of linear mixed effects model remains same across all distinct values of given categorical partitioning variable.

**Usage**

```
StabCat(data, patid, fixed, splitvar)
```

**Arguments**

<code>data</code>	name of the dataset. It must contain variable specified for <code>patid</code> (indicating subject id) and all the variables specified in the formula and the categorical partitioning variable of interest specified in <code>splitvar</code> . Note that, only numerically coded categorical variable should be specified.
<code>patid</code>	name of the subject id variable.
<code>fixed</code>	a two-sided linear formula object describing the fixed-effects part of the model, with the response on the left of a <code>~</code> operator and the terms, separated by <code>+</code> operators, on the right. Model with <code>-1</code> to the end of right side indicates no intercept. For model with no fixed effect beyond intercept, please specify only <code>1</code> right to the <code>~</code> operator.

`splitvar` the categorical partitioning variable of interest. It's value should not change over time.

### Details

The categorical partitioning variable of interest. It's value should not change over time.

$$Y_i(t) = W_i(t) \theta + b_i + \epsilon_{it}$$

where  $W_i(t)$  is the design matrix,  $\theta$  is the parameter associated with  $W_i(t)$  and  $b_i$  is the random intercept. Also,  $\epsilon_{it} \sim N(0, \sigma^2)$  and  $b_i \sim N(0, \sigma_u^2)$ . Let  $X$  be the baseline categorical partitioning variable of interest. `StabCat()` performs the following omnibus test

$$H_0: \theta_{(g)} = \theta_0 \text{ vs. } H_1: \theta_{(g)} \neq \theta_0, \text{ for all } g$$

where,  $\theta_{(g)}$  is the true value of  $\theta$  for subjects with  $X=C_g$  where  $C_g$  is the any value realized by  $X$ .

### Value

`p` It returns the p-value for parameter instability test

### Author(s)

Madan Gopal Kundu <madan\_g.kundu@yahoo.com>

### References

Kundu, M. G., and Harezlak, J. (2019). Regression trees for longitudinal data with baseline covariates. *Biostatistics & Epidemiology*, 3(1):1-22.

### See Also

[StabCont](#), [LongCART](#), [plotLongCART](#), [textLongCART](#)

### Examples

```
#--- Get the data
data(ACTG175)

#--- Run StabCat()
out<- StabCat(data=ACTG175, patid="pidnum", fixed=cd4~time, splitvar="gender")
out$pval
```



---

StabCont *parameter stability test for continuous partitioning variable*

---

### Description

Performs parameter stability test (Kundu and Harezlak, 2019) with continuous partitioning variable to determine whether the parameters of linear mixed effects model remains same across all distinct values of given continuous partitioning variable.

### Usage

```
StabCont(data, patid, fixed, splitvar)
```

### Arguments

data	name of the dataset. It must contain variable specified for patid (indicating subject id) and all the variables specified in the formula and the StabCont(data, fixed, splitvar)partitioning variable of interest specified in splitvar.
patid	name of the subject id variable.
fixed	a two-sided linear formula object describing the fixed-effects part of the model, with the response on the left of a ~ operator and the terms, separated by + operators, on the right. Model with -1 to the end of right side indicates no intercept. For model with no fixed effect beyond intercept, please specify only 1 right to the ~ operator.
splitvar	the continuous partitioning variable of interest. It's value should not change over time.

### Details

The continuous partitioning variable of interest. It's value should not change over time.

$$Y_i(t) = W_i(t)\theta + b_i + \epsilon_{it}$$

where  $W_i(t)$  is the design matrix,  $\theta$  is the parameter associated with  $W_i(t)$  and  $b_i$  is the random intercept. Also,  $\epsilon_{it} \sim N(0, \sigma^2)$  and  $b_i \sim N(0, \sigma_u^2)$ . Let  $X$  be the baseline continuous partitioning variable of interest. StabCont() performs the following omnibus test

$$H_0: \theta_{(g)} = \theta_0 \text{ vs. } H_1: \theta_{(g)} \neq \theta_0, \text{ for all } g$$

where,  $\theta_{(g)}$  is the true value of  $\theta$  for subjects with  $X=C_g$  where  $C_g$  is the any value realized by  $X$ .

### Value

p It returns the p-value for parameter instability test

### Author(s)

Madan Gopal Kundu <madan\_g.kundu@yahoo.com>

## References

Kundu, M. G., and Harezlak, J. (2019). Regression trees for longitudinal data with baseline covariates. *Biostatistics & Epidemiology*, 3(1):1-22.

## See Also

[StabCont](#), [LongCART](#), [plotLongCART](#), [textLongCART](#)

## Examples

```
#--- Get the data
data(ACTG175)

#--- Run StabCont()
out<- StabCont(data=ACTG175, patid="pidnum", fixed=cd4~time, splitvar="age")
out$pval
```

---

textLongCART

*Place text on LongCART tree*

---

## Description

Labels the current plot of the tree generated from LongCART object with text.

## Usage

```
textLongCART(x, splits = TRUE, all = FALSE,
             use.n = FALSE, minlength = 1L, ...)
```

## Arguments

x	a fitted object of class "LongCART", containing a linear mixed effects tree.
splits	similar to plot.rpart; logical flag. If TRUE (default), then the splits in the tree are labeled with the criterion for the split.
all	Logical. If TRUE, all nodes are labeled, otherwise just terminal nodes.
use.n	Logical. If TRUE, adds n to label.
minlength	the length to use for factor labels. A value of 1 causes them to be printed as 'a', 'b', .... Larger values use abbreviations of the label names. See the labels.rpart function for details.
...	arguments to be passed to or from other methods.

## Author(s)

Madan Gopal Kundu <madan\_g.kundu@yahoo.com>

## References

Kundu, M. G., and Harezlak, J. (2019). Regression trees for longitudinal data with baseline covariates. *Biostatistics & Epidemiology*, 3(1):1-22.

## See Also

[plotLongCART](#), [LongCART](#), [StabCat](#), [StabCont](#)

## Examples

```
#--- Get the data
data(ACTG175)

#--- Run LongCART()
gvars=c("age", "gender", "wtkg", "hemo", "homo", "drugs",
        "karnof", "oprior", "z30", "zprior", "race",
        "str2", "symptom", "treat", "offtrt")
tgvars=c(1, 0, 1, 0, 0, 0,
        1, 0, 0, 0, 0,
        0, 0, 0, 0)

out<- LongCART(data=ACTG175, patid="pidnum", fixed=cd4~time,
              gvars=gvars, tgvars=tgvars, alpha=0.05,
              minsplit=100, minbucket=50, coef.digits=2)

#--- Plot tree

par(xpd = T)
plot(out, compress = T)
text(out, use.n = T)
```

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