

# Package ‘DynForest’

March 22, 2024

**Title** Random Forest with Multivariate Longitudinal Predictors

**Version** 1.1.3

**Description** Based on random forest principle, 'DynForest' is able to include multiple longitudinal predictors to provide individual predictions. Longitudinal predictors are modeled through the random forest. The methodology is fully described for a survival outcome in: Devaux, Helmer, Genuer & Proust-Lima (2023) [doi:10.1177/09622802231206477](https://doi.org/10.1177/09622802231206477).

**Imports** DescTools, cmprsk, doParallel, doRNG, foreach, ggplot2, lamm, methods, pbapply, pec, prodlim, stringr, survival, zoo

**Depends** R (>= 4.3.0)

**License** LGPL (>= 3)

**LazyData** true

**Encoding** UTF-8

**RoxygenNote** 7.2.3

**URL** <https://github.com/anthonydevaux/DynForest>

**BugReports** <https://github.com/anthonydevaux/DynForest/issues>

**Suggests** knitr, rmarkdown

**VignetteBuilder** knitr

**NeedsCompilation** no

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compute_gVIMP	<i>Compute the grouped importance of variables (gVIMP) statistic</i>
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### Description

Compute the grouped importance of variables (gVIMP) statistic

### Usage

```
compute_gVIMP(
  DynForest_obj,
  IBS.min = 0,
  IBS.max = NULL,
  group = NULL,
  ncores = NULL,
  seed = 1234
)
```

### Arguments

DynForest_obj	DynForest object containing the dynamic random forest used on train data
IBS.min	(Only with survival outcome) Minimal time to compute the Integrated Brier Score. Default value is set to 0.
IBS.max	(Only with survival outcome) Maximal time to compute the Integrated Brier Score. Default value is set to the maximal time-to-event found.
group	A list of groups with the name of the predictors assigned in each group
ncores	Number of cores used to grow trees in parallel. Default value is the number of cores of the computer-1.
seed	Seed to replicate results

**Value**

compute\_gVIMP() function returns a list with the following elements:

Inputs	A list of 3 elements: Longitudinal, Numeric and Factor. Each element contains the names of the predictors
group	A list of each group defined in group argument
gVIMP	A numeric vector containing the gVIMP for each group defined in group argument
tree_oob_err	A numeric vector containing the OOB error for each tree needed to compute the VIMP statistic
IBS.range	A vector containing the IBS min and max

**Examples**

```

data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                    random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                   random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data

```

```

Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Compute gVIMP statistic
res_dyn_gVIMP <- compute_gVIMP(DynForest_obj = res_dyn,
                               group = list(group1 = c("serBilir", "SGOT"),
                                             group2 = c("albumin", "alkaline")),
                               ncores = 2, seed = 1234)

```

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compute_OOBerror	<i>Compute the Out-Of-Bag error (OOB error)</i>
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### Description

Compute the Out-Of-Bag error (OOB error)

### Usage

```
compute_OOBerror(DynForest_obj, IBS.min = 0, IBS.max = NULL, ncores = NULL)
```

### Arguments

DynForest_obj	DynForest object containing the dynamic random forest used on train data
IBS.min	(Only with survival outcome) Minimal time to compute the Integrated Brier Score. Default value is set to 0.
IBS.max	(Only with survival outcome) Maximal time to compute the Integrated Brier Score. Default value is set to the maximal time-to-event found.
ncores	Number of cores used to grow trees in parallel. Default value is the number of cores of the computer-1.

### Value

compute\_OOBerror() function return a list with the following elements:

data	A list containing the data used to grow the trees
rf	A table with each tree in column. Provide multiple characteristics about the tree building
type	Outcome type



```

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Compute OOB error
res_dyn_OOB <- compute_OOBError(DynForest_obj = res_dyn, ncores = 2)

```

---

compute\_VIMP

---

*Compute the importance of variables (VIMP) statistic*


---

## Description

Compute the importance of variables (VIMP) statistic

## Usage

```

compute_VIMP(
  DynForest_obj,
  IBS.min = 0,
  IBS.max = NULL,
  ncores = NULL,
  seed = 1234
)

```

## Arguments

DynForest_obj	DynForest object containing the dynamic random forest used on train data
IBS.min	(Only with survival outcome) Minimal time to compute the Integrated Brier Score. Default value is set to 0.
IBS.max	(Only with survival outcome) Maximal time to compute the Integrated Brier Score. Default value is set to the maximal time-to-event found.
ncores	Number of cores used to grow trees in parallel. Default value is the number of cores of the computer-1.
seed	Seed to replicate results

*compute\_VIMP*

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**Value**

`compute_VIMP()` function returns a list with the following elements:

Inputs	A list of 3 elements: Longitudinal, Numeric and Factor. Each element contains the names of the predictors
Importance	A list of 3 elements: Longitudinal, Numeric and Factor. Each element contains a numeric vector of VIMP
tree_oob_err	A numeric vector containing the OOB error for each tree needed to compute the VIMP statistic
IBS.range	A vector containing the IBS min and max

## Examples

```

data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                                "serBilir", "SGOT",
                                "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                    random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                    random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",

```



```
timeVarModel = timeVarModel, Y = Y,  
ntree = 50, nodesize = 5, minsplit = 5,  
cause = 2, ncores = 2, seed = 1234)  
  
# Compute VIMP statistic  
res_dyn_VIMP <- compute_VIMP(DynForest_obj = res_dyn, ncores = 2, seed = 1234)
```

---

data_simu1	<i>data_simu1 dataset</i>
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---

## Description

Simulated dataset 1 with continuous outcome

## Format

Longitudinal dataset with 1200 rows and 13 columns for 200 subjects

**id** Subject identifier  
**time** Time measurement  
**cont\_covar1** Continuous time-fixed predictor 1  
**cont\_covar2** Continuous time-fixed predictor 2  
**bin\_covar1** Binary time-fixed predictor 1  
**bin\_covar2** Binary time-fixed predictor 2  
**marker1** Continuous time-dependent predictor 1  
**marker2** Continuous time-dependent predictor 2  
**marker3** Continuous time-dependent predictor 3  
**marker4** Continuous time-dependent predictor 4  
**marker5** Continuous time-dependent predictor 5  
**marker6** Continuous time-dependent predictor 6  
**Y\_res** Continuous outcome

## Examples

```
data(data_simu1)
```

---

`data_simu2`*data\_simu1 dataset*

---

**Description**

Simulated dataset 2 with continuous outcome

**Format**

Longitudinal dataset with 1200 rows and 13 columns for 200 subjects

**id** Subject identifier

**time** Time measurement

**cont\_covar1** Continuous time-fixed predictor 1

**cont\_covar2** Continuous time-fixed predictor 2

**bin\_covar1** Binary time-fixed predictor 1

**bin\_covar2** Binary time-fixed predictor 2

**marker1** Continuous time-dependent predictor 1

**marker2** Continuous time-dependent predictor 2

**marker3** Continuous time-dependent predictor 3

**marker4** Continuous time-dependent predictor 4

**marker5** Continuous time-dependent predictor 5

**marker6** Continuous time-dependent predictor 6

**Y\_res** Continuous outcome

**Examples**

```
data(data_simu2)
```

---

`DynForest`*Random forest with multivariate longitudinal endogenous covariates*

---

**Description**

Build a random forest using multivariate longitudinal endogenous covariates

**Usage**

```

DynForest(
  timeData = NULL,
  fixedData = NULL,
  idVar = NULL,
  timeVar = NULL,
  timeVarModel = NULL,
  Y = NULL,
  ntree = 200,
  mtry = NULL,
  nodesize = 1,
  minsplit = 2,
  cause = 1,
  nsplit_option = "quantile",
  ncores = NULL,
  seed = 1234,
  verbose = TRUE
)

```

**Arguments**

timeData	A data.frame containing the id and time measurements variables and the time-dependent predictors.
fixedData	A data.frame containing the id variable and the time-fixed predictors. Categorical variables should be characterized as factor.
idVar	A character indicating the name of variable to identify the subjects
timeVar	A character indicating the name of time variable
timeVarModel	A list for each time-dependent predictors containing a list of formula for fixed and random part from the mixed model
Y	A list of output which should contain: type defines the nature of the outcome, can be "surv", "numeric" or "factor"; .
ntree	Number of trees to grow. Default value set to 200.
mtry	Number of candidate variables randomly drawn at each node of the trees. This parameter should be tuned by minimizing the OOB error. Default is defined as the square root of the number of predictors.
nodesize	Minimal number of subjects required in both child nodes to split. Cannot be smaller than 1.
minsplit	(Only with survival outcome) Minimal number of events required to split the node. Cannot be smaller than 2.
cause	(Only with competing events) Number indicates the event of interest.
nsplit_option	A character indicates how the values are chosen to build the two groups for the splitting rule (only for continuous predictors). Values are chosen using deciles (nsplit_option="quantile") or randomly (nsplit_option="sample"). Default value is "quantile".

ncores	Number of cores used to grow trees in parallel. Default value is the number of cores of the computer-1.
seed	Seed to replicate results
verbose	A logical controlling the function progress. Default is TRUE

### Details

The function currently supports survival (competing or single event), continuous or categorical outcome.

FUTUR IMPLEMENTATIONS:

- Continuous longitudinal outcome
- Functional data analysis

### Value

DynForest function returns a list with the following elements:

data	A list containing the data used to grow the trees
rf	A table with each tree in column. Provide multiple characteristics about the tree building
type	Outcome type
times	A numeric vector containing the time-to-event for all subjects
cause	Indicating the cause of interest
causes	A numeric vector containing the causes indicator
Inputs	A list of 3 elements: Longitudinal, Numeric and Factor. Each element contains the names of the p
Longitudinal.model	A list of longitudinal markers containing the formula used for modeling in the random forest
param	A list containing the hyperparameters
comput.time	Computation time

### Author(s)

Anthony Devaux (<anthony.devauxbarault@gmail.com>)

### References

- Devaux A., Helmer C., Genuer R., Proust-Lima C. (2023). Random survival forests with multivariate longitudinal endogenous covariates. SMMR [doi:10.1177/09622802231206477](https://doi.org/10.1177/09622802231206477)



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getTree	<i>Extract some information about the split for a tree by user</i>
---------	--

---

**Description**

Extract some information about the split for a tree by user

**Usage**

```
getTree(DynForest_obj, tree)
```

**Arguments**

DynForest_obj	DynForest object containing the dynamic random forest used on train data
tree	Integer indicating the tree identifier

**Value**

A table sorted by the node/leaf identifier with each row representing a node/leaf. Each column provides information about the splits:

type	The nature of the predictor ( <code>Longitudinal</code> for longitudinal predictor, <code>Numeric</code> for continuous predictor or <code>Factor</code> for categorical predictor)
var_split	The predictor used for the split defined by its order in <code>timeData</code> and <code>fixedData</code>
feature	The feature used for the split defined by its position in random statistic
threshold	The threshold used for the split (only with <code>Longitudinal</code> and <code>Numeric</code> ). No information is returned for <code>Factor</code>
N	The number of subjects in the node/leaf
Nevent	The number of events of interest in the node/leaf (only with survival outcome)
depth	the depth level of the node/leaf

**See Also**

[DynForest summary.DynForest](#)

**Examples**

```
data(pbc2)

# Get Gaussian distribution for longitudinal predictors
```

```

pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id","time",
                                "serBilir","SGOT",
                                "albumin","alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                     random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                 random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                    random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id","age","drug","sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id","years","event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Extract split information from tree 4
res_tree4 <- getTree(DynForest_obj = res_dyn, tree = 4)

```

---

getTreeNodes

*Extract nodes identifiers for a given tree*


---

## Description

Extract nodes identifiers for a given tree

**Usage**

```
getTreeNodes(DynForest_obj, tree = NULL)
```

**Arguments**

```
DynForest_obj  A DynForest object from DynForest() function
tree           Integer indicating the tree identifier
```

**Value**

Extract nodes identifiers for a given tree

**Examples**

```
data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                    random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                   random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))
```



```
# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Extract nodes identifiers for a given tree
getTreeNodees(DynForest_obj = res_dyn, tree = 1)
```

pbc2

*pbc2 dataset*


---

## Description

pbc2 data from Mayo clinic

## Format

Longitudinal dataset with 1945 rows and 19 columns for 312 patients

**id** Patient identifier

**time** Time measurement

**ascites** Presence of ascites (Yes/No)

**hepatomegaly** Presence of hepatomegaly (Yes/No)

**spiders** Blood vessel malformations in the skin (Yes/No)

**edema** Edema levels (No edema/edema no diuretics/edema despite diuretics)

**serBilir** Level of serum bilirubin

**serChol** Level of serum cholesterol

**albumin** Level of albumin

**alkaline** Level of alkaline phosphatase

**SGOT** Level of aspartate aminotransferase

**platelets** Platelet count

**prothrombin** Prothrombin time

**histologic** Histologic stage of disease

**drug** Drug treatment (D-penicillmain/Placebo)

**age** Age at enrollment

**sex** Sex of patient

**years** Time-to-event in years

**event** Event indicator: 0 (alive), 1 (transplanted) and 2 (dead)

**Source**

```
pb2c2 joinRML
```

**Examples**

```
data(pbc2)
```

---

plot.DynForest	<i>Plot function in DynForest</i>
----------------	-----------------------------------

---

**Description**

This function displays a plot of CIF for a given node and tree (for class `DynForest`), the most predictive variables with the minimal depth (for class `DynForestVarDepth`), the variable importance (for class `DynForestVIMP`) or the grouped variable importance (for class `DynForestgVIMP`).

**Usage**

```
## S3 method for class 'DynForest'
plot(x, tree = NULL, nodes = NULL, id = NULL, max_tree = NULL, ...)

## S3 method for class 'DynForestVarDepth'
plot(x, plot_level = c("predictor", "feature"), ...)

## S3 method for class 'DynForestVIMP'
plot(x, PCT = FALSE, ordering = TRUE, ...)

## S3 method for class 'DynForestgVIMP'
plot(x, PCT = FALSE, ...)

## S3 method for class 'DynForestPred'
plot(x, id = NULL, ...)
```

**Arguments**

x	Object inheriting from classes <code>DynForest</code> , <code>DynForestVarDepth</code> , <code>DynForestVIMP</code> or <code>DynForestgVIMP</code> , to respectively plot the CIF, the minimal depth, the variable importance or grouped variable importance.
tree	For <code>DynForest</code> class, integer indicating the tree identifier
nodes	For <code>DynForest</code> class, identifiers for the selected nodes
id	For <code>DynForest</code> and <code>DynForestPred</code> classes, identifier for a given subject
max_tree	For <code>DynForest</code> class, integer indicating the number of tree to display while using id argument
...	Optional parameters to be passed to the low level function

plot_level	For DynForestVarDepth class, compute the statistic at predictor (plot_level="predictor") or feature (plot_level="feature") level
PCT	For DynForestVIMP or DynForestgVIMP class, display VIMP statistic in percentage. Default value is FALSE.
ordering	For DynForestVIMP class, order predictors according to VIMP value. Default value is TRUE.

### Value

plot() function displays:

With DynForestVarDepth	the minimal depth for each predictor/feature
With DynForestVIMP	the VIMP for each predictor
With DynForestgVIMP	the grouped-VIMP for each given group

### See Also

[DynForest](#) [var\\_depth](#) [compute\\_VIMP](#) [compute\\_gVIMP](#)

### Examples

```
data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                   random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
```

```

                                random = ~ time),
alkaline = list(fixed = alkaline ~ time,
                random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Plot estimated CIF at nodes 17 and 32
plot(x = res_dyn, tree = 1, nodes = c(17,32))

# Run var_depth function
res_varDepth <- var_depth(res_dyn)

# Plot minimal depth
plot(x = res_varDepth, plot_level = "feature")

# Compute VIMP statistic
res_dyn_VIMP <- compute_VIMP(DynForest_obj = res_dyn, ncores = 2)

# Plot VIMP
plot(x = res_dyn_VIMP, PCT = TRUE)

# Compute gVIMP statistic
res_dyn_gVIMP <- compute_gVIMP(DynForest_obj = res_dyn,
                              group = list(group1 = c("serBilir", "SGOT"),
                                             group2 = c("albumin", "alkaline")),
                              ncores = 2)

# Plot gVIMP
plot(x = res_dyn_gVIMP, PCT = TRUE)

# Sample 5 subjects to predict the event
set.seed(123)
id_pred <- sample(id, 5)

# Create predictors objects
pbc2_pred <- pbc2[which(pbc2$id%in%id_pred),]
timeData_pred <- pbc2_pred[,c("id", "time", "serBilir", "SGOT", "albumin", "alkaline")]
fixedData_pred <- unique(pbc2_pred[,c("id", "age", "drug", "sex")])

# Predict the CIF function for the new subjects with landmark time at 4 years
pred_dyn <- predict(object = res_dyn,

```

```

timeData = timeData_pred, fixedData = fixedData_pred,
idVar = "id", timeVar = "time",
t0 = 4)

# Plot predicted CIF for subjects 26 and 110
plot(x = pred_dyn, id = c(26, 110))

```

---

predict.DynForest      *Prediction using dynamic random forests*

---

## Description

Prediction using dynamic random forests

## Usage

```

## S3 method for class 'DynForest'
predict(
  object,
  timeData = NULL,
  fixedData = NULL,
  idVar,
  timeVar,
  t0 = NULL,
  ...
)

```

## Arguments

object	DynForest object containing the dynamic random forest used on train data
timeData	A data.frame containing the id and time measurements variables and the time-dependent predictors.
fixedData	A data.frame containing the id variable and the time-fixed predictors. Non-continuous variables should be characterized as factor.
idVar	A character indicating the name of variable to identify the subjects
timeVar	A character indicating the name of time variable
t0	Landmark time
...	Optional parameters to be passed to the low level function

## Value

Return the outcome of interest for the new subjects: matrix of probability of event of interest in survival mode, average value in regression mode and most likely value in classification mode

**Examples**

```

data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                                "serBilir", "SGOT",
                                "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                     random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                 random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                    random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Sample 5 subjects to predict the event
set.seed(123)
id_pred <- sample(id, 5)

# Create predictors objects
pbc2_pred <- pbc2[which(pbc2$id%in%id_pred),]

```

```

timeData_pred <- pbc2_pred[,c("id", "time", "serBilir", "SGOT", "albumin", "alkaline")]
fixedData_pred <- unique(pbc2_pred[,c("id", "age", "drug", "sex")])

# Predict the CIF function for the new subjects with landmark time at 4 years
pred_dyn <- predict(object = res_dyn,
                    timeData = timeData_pred, fixedData = fixedData_pred,
                    idVar = "id", timeVar = "time",
                    t0 = 4)

```

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print.DynForest	<i>Print function</i>
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### Description

This function displays a brief summary regarding the trees (for class `DynForest`), a data frame with variable importance (for class `DynForestVIMP`) or the grouped variable importance (for class `DynForestgVIMP`).

### Usage

```

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

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

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

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

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

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

```

### Arguments

<code>x</code>	Object inheriting from classes <code>DynForest</code> , <code>DynForestVIMP</code> or <code>DynForestgVIMP</code> .
<code>...</code>	Optional parameters to be passed to the low level function

### See Also

[DynForest](#) [var\\_depth](#) [compute\\_VIMP](#) [compute\\_gVIMP](#) [compute\\_OOBerror](#)

**Examples**

```

data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                     random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                   random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Print function
print(res_dyn)

# Compute VIMP statistic
res_dyn_VIMP <- compute_VIMP(DynForest_obj = res_dyn, ncores = 2, seed = 1234)

```



```
# Print function
print(res_dyn_VIMP)

# Compute gVIMP statistic
res_dyn_gVIMP <- compute_gVIMP(DynForest_obj = res_dyn,
                              group = list(group1 = c("serBilir", "SGOT"),
                                           group2 = c("albumin", "alkaline")),
                              ncores = 2, seed = 1234)

# Print function
print(res_dyn_gVIMP)

# Run var_depth function
res_varDepth <- var_depth(res_dyn)

# Print function
print(res_varDepth)
```

---

summary.DynForest      *Display the summary of DynForest*

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## Description

Display the summary of DynForest

## Usage

```
## S3 method for class 'DynForest'
summary(object, ...)

## S3 method for class 'DynForest00B'
summary(object, ...)
```

## Arguments

object            DynForest or DynForest00B object  
...                Optional parameters to be passed to the low level function

## Value

Return some information about the random forest

**Examples**

```

data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]

timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                    random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                   random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Compute OOB error
res_dyn_OOB <- compute_OOBerror(DynForest_obj = res_dyn, ncores = 2)

# DynForest summary
summary(object = res_dyn_OOB)

```

---

var_depth	<i>Extract characteristics from the trees building process</i>
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---

**Description**

Extract characteristics from the trees building process

**Usage**

```
var_depth(DynForest_obj)
```

**Arguments**

DynForest\_obj DynForest object

**Value**

var\_depth function return a list with the following elements:

min\_depth A table providing for each feature in row: the average depth and the rank

var\_node\_depth A table providing for each tree in column the minimal depth for each feature in row. NA indicates that the

var\_count A table providing for each tree in column the number of times where the feature is used (in row). 0 value

**See Also**

[DynForest](#)

**Examples**

```
data(pbc2)

# Get Gaussian distribution for longitudinal predictors
pbc2$serBilir <- log(pbc2$serBilir)
pbc2$SGOT <- log(pbc2$SGOT)
pbc2$albumin <- log(pbc2$albumin)
pbc2$alkaline <- log(pbc2$alkaline)

# Sample 100 subjects
set.seed(1234)
id <- unique(pbc2$id)
id_sample <- sample(id, 100)
id_row <- which(pbc2$id%in%id_sample)

pbc2_train <- pbc2[id_row,]
```

```
timeData_train <- pbc2_train[,c("id", "time",
                               "serBilir", "SGOT",
                               "albumin", "alkaline")]

# Create object with longitudinal association for each predictor
timeVarModel <- list(serBilir = list(fixed = serBilir ~ time,
                                     random = ~ time),
                    SGOT = list(fixed = SGOT ~ time + I(time^2),
                                 random = ~ time + I(time^2)),
                    albumin = list(fixed = albumin ~ time,
                                    random = ~ time),
                    alkaline = list(fixed = alkaline ~ time,
                                    random = ~ time))

# Build fixed data
fixedData_train <- unique(pbc2_train[,c("id", "age", "drug", "sex")])

# Build outcome data
Y <- list(type = "surv",
          Y = unique(pbc2_train[,c("id", "years", "event")]))

# Run DynForest function
res_dyn <- DynForest(timeData = timeData_train, fixedData = fixedData_train,
                    timeVar = "time", idVar = "id",
                    timeVarModel = timeVarModel, Y = Y,
                    ntree = 50, nodesize = 5, minsplit = 5,
                    cause = 2, ncores = 2, seed = 1234)

# Run var_depth function
res_varDepth <- var_depth(res_dyn)
```

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