Package: hdcuremodels (via r-universe)

May 17, 2026

Title High-Dimensional Cure Models

Version 0.0.6

Date 2025-12-01

Description Provides functions for fitting various penalized parametric and semi-parametric mixture cure models with different penalty functions, testing for a significant cure fraction, and testing for sufficient follow-up as described in Fu et al (2022)<doi:10.1002/sim.9513> and Archer et al (2024)<doi:10.1186/s13045-024-01553-6>. False discovery rate controlled variable selection is provided using model-X knock-offs.

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Encoding UTF-8

Depends R (>= 4.2.0)

Imports doParallel, flexsurv, flexsurvcure, foreach, ggplot2, ggpubr, glmnet, knockoff, mvnfast, parallel, plyr, methods, survival, withr

Roxygen list(markdown = TRUE, roclets = c (``namespace", ``rd"))

RoxygenNote 7.3.3

Remotes ropensci-review-tools/srr

Suggests knitr, Rdsdp, rmarkdown, roxygen2, srr, testthat (>= 3.0.0)

VignetteBuilder knitr

LazyData true

URL `https://github.com/ropensci/hdcuremodels`

BugReports `https://github.com/ropensci/hdcuremodels/issues`

Config/testthat/edition 3

Config/pak/sysreqs cmake make libicu-dev

Repository https://ropensci.r-universe.dev

Date/Publication 2025-12-03 20:10:56 UTC

RemoteUrl https://github.com/ropensci/hdcuremodels

RemoteRef main

RemoteSha 466902cdc609628f1d9f0fb42e70f7dfb73eda72

Contents

`amltest` *AML test data*

Description

Duration of complete response for 40 cytogenetically normal AML patients and a subset of 320 transcript expression from RNA-sequencing.

Usage

```
amltest
```

Format

A data frame with 40 rows (subjects) and 322 columns:

cryr duration of complete response in years

relapse.death censoring indicator: 1 = relapsed or died; 0 = alive at last follow=up

ENSG00000001561 normalized expression for indicated transcript

ENSG00000005249 normalized expression for indicated transcript

ENSG00000006757 normalized expression for indicated transcript

ENSG00000007062 normalized expression for indicated transcript

ENSG00000007968 normalized expression for indicated transcript

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ENSG00000273018	normalized expression for indicated transcript
ENSG00000273033	normalized expression for indicated transcript

Source

doi:10.1186/s13045-024-01553-6

`amltrain` *AML training data*

Description

Duration of complete response for 306 cytogenetically normal AML patients and a subset of 320 transcript expression from RNA-sequencing.

Usage

```
amltrain
```

Format

A data frame with 306 rows (subjects) and 322 columns:

cryr	duration of complete response in years
relapse.death	censoring indicator: 1 = relapsed or died; 0 = alive at last follow=up
ENSG00000001561	normalized expression for indicated transcript
ENSG00000005249	normalized expression for indicated transcript
ENSG00000006757	normalized expression for indicated transcript
ENSG00000007062	normalized expression for indicated transcript
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ENSG00000179630	normalized expression for indicated transcript
ENSG00000179639	normalized expression for indicated transcript

ENSG00000179820	normalized expression for indicated transcript
ENSG00000180096	normalized expression for indicated transcript
ENSG00000180596	normalized expression for indicated transcript
ENSG00000180902	normalized expression for indicated transcript
ENSG00000181104	normalized expression for indicated transcript
ENSG00000182866	normalized expression for indicated transcript
ENSG00000182871	normalized expression for indicated transcript
ENSG00000183087	normalized expression for indicated transcript
ENSG00000183091	normalized expression for indicated transcript
ENSG00000184371	normalized expression for indicated transcript
ENSG00000185129	normalized expression for indicated transcript
ENSG00000185201	normalized expression for indicated transcript
ENSG00000185245	normalized expression for indicated transcript
ENSG00000185291	normalized expression for indicated transcript
ENSG00000185304	normalized expression for indicated transcript
ENSG00000185710	normalized expression for indicated transcript
ENSG00000185883	normalized expression for indicated transcript
ENSG00000185986	normalized expression for indicated transcript
ENSG00000186130	normalized expression for indicated transcript
ENSG00000186854	normalized expression for indicated transcript
ENSG00000187010	normalized expression for indicated transcript
ENSG00000187627	normalized expression for indicated transcript
ENSG00000187653	normalized expression for indicated transcript
ENSG00000187837	normalized expression for indicated transcript
ENSG00000188002	normalized expression for indicated transcript
ENSG00000188107	normalized expression for indicated transcript
ENSG00000188211	normalized expression for indicated transcript
ENSG00000188636	normalized expression for indicated transcript
ENSG00000188738	normalized expression for indicated transcript
ENSG00000188856	normalized expression for indicated transcript
ENSG00000189164	normalized expression for indicated transcript
ENSG00000189223	normalized expression for indicated transcript
ENSG00000196155	normalized expression for indicated transcript
ENSG00000196189	normalized expression for indicated transcript
ENSG00000196415	normalized expression for indicated transcript
ENSG00000196565	normalized expression for indicated transcript
ENSG00000197081	normalized expression for indicated transcript

ENSG00000197121	normalized expression for indicated transcript
ENSG00000197253	normalized expression for indicated transcript
ENSG00000197256	normalized expression for indicated transcript
ENSG00000197321	normalized expression for indicated transcript
ENSG00000197561	normalized expression for indicated transcript
ENSG00000197728	normalized expression for indicated transcript
ENSG00000197860	normalized expression for indicated transcript
ENSG00000197937	normalized expression for indicated transcript
ENSG00000197951	normalized expression for indicated transcript
ENSG00000198743	normalized expression for indicated transcript
ENSG00000198838	normalized expression for indicated transcript
ENSG00000199347	normalized expression for indicated transcript
ENSG00000200243	normalized expression for indicated transcript
ENSG00000201801	normalized expression for indicated transcript
ENSG00000203872	normalized expression for indicated transcript
ENSG00000204172	normalized expression for indicated transcript
ENSG00000205571	normalized expression for indicated transcript
ENSG00000205593	normalized expression for indicated transcript
ENSG00000208772	normalized expression for indicated transcript
ENSG00000213085	normalized expression for indicated transcript
ENSG00000213261	normalized expression for indicated transcript
ENSG00000213626	normalized expression for indicated transcript
ENSG00000213722	normalized expression for indicated transcript
ENSG00000213906	normalized expression for indicated transcript
ENSG00000213967	normalized expression for indicated transcript
ENSG00000214016	normalized expression for indicated transcript
ENSG00000214425	normalized expression for indicated transcript
ENSG00000216316	normalized expression for indicated transcript
ENSG00000220008	normalized expression for indicated transcript
ENSG00000223345	normalized expression for indicated transcript
ENSG00000224080	normalized expression for indicated transcript
ENSG00000225138	normalized expression for indicated transcript
ENSG00000226471	normalized expression for indicated transcript
ENSG00000227097	normalized expression for indicated transcript
ENSG00000227191	normalized expression for indicated transcript
ENSG00000227615	normalized expression for indicated transcript
ENSG00000228049	normalized expression for indicated transcript

ENSG00000229153	normalized expression for indicated transcript
ENSG00000230076	normalized expression for indicated transcript
ENSG00000231160	normalized expression for indicated transcript
ENSG00000231721	normalized expression for indicated transcript
ENSG00000233927	normalized expression for indicated transcript
ENSG00000233974	normalized expression for indicated transcript
ENSG00000234883	normalized expression for indicated transcript
ENSG00000236876	normalized expression for indicated transcript
ENSG00000237298	normalized expression for indicated transcript
ENSG00000237892	normalized expression for indicated transcript
ENSG00000238160	normalized expression for indicated transcript
ENSG00000239437	normalized expression for indicated transcript
ENSG00000241399	normalized expression for indicated transcript
ENSG00000241489	normalized expression for indicated transcript
ENSG00000241529	normalized expression for indicated transcript
ENSG00000244405	normalized expression for indicated transcript
ENSG00000247627	normalized expression for indicated transcript
ENSG00000249592	normalized expression for indicated transcript
ENSG00000250116	normalized expression for indicated transcript
ENSG00000250251	normalized expression for indicated transcript
ENSG00000251079	normalized expression for indicated transcript
ENSG00000253210	normalized expression for indicated transcript
ENSG00000253276	normalized expression for indicated transcript
ENSG00000254415	normalized expression for indicated transcript
ENSG00000259276	normalized expression for indicated transcript
ENSG00000260727	normalized expression for indicated transcript
ENSG00000261377	normalized expression for indicated transcript
ENSG00000264885	normalized expression for indicated transcript
ENSG00000264895	normalized expression for indicated transcript
ENSG00000267136	normalized expression for indicated transcript
ENSG00000267551	normalized expression for indicated transcript
ENSG00000267702	normalized expression for indicated transcript
ENSG00000268001	normalized expression for indicated transcript
ENSG00000268573	normalized expression for indicated transcript
ENSG00000270554	normalized expression for indicated transcript
ENSG00000270562	normalized expression for indicated transcript
ENSG00000271646	normalized expression for indicated transcript
ENSG00000273018	normalized expression for indicated transcript
ENSG00000273033	normalized expression for indicated transcript

Source

doi:10.1186/s13045-024-01553-6

`auc_mcm` *AUC for cure prediction using mean score imputation*

Description

This function calculates the AUC for cure prediction using the mean score imputation (MSI) method proposed by Asano et al (2014).

Usage

```
auc_mcm(object, newdata, cure_cutoff = 5, model_select = "AIC")
```

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.
`newdata`	an optional data.frame that minimally includes the incidence and/or latency variables to use for predicting the response. If omitted, the training data are used.
`cure_cutoff`	cutoff value for cure, used to produce a proxy for the unobserved cure status (default is 5 representing 5 years). Users should be careful to note the time scale of their data and adjust this according to the time scale and clinical application.
`model_select`	either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are: • `"AIC"` for the minimum AIC (default). • `"mAIC"` for the minimum modified AIC. • `"cAIC"` for the minimum corrected AIC. • `"BIC"`, for the minimum BIC. • `"mBIC"` for the minimum modified BIC. • `"EBIC"` for the minimum extended BIC. • `"logLik"` for the step that maximizes the log-likelihood. • `n` where n is any numeric value from the solution path. This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.

Value

Returns the AUC value for cure prediction using the mean score imputation (MSI) method.

References

Asano, J., Hirakawa, H., Hamada, C. (2014) Assessing the prediction accuracy of cure in the Cox proportional hazards cure model: an application to breast cancer data. *Pharmaceutical Statistics*, **13**:357–363.

See Also

`concordance_mcm`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
testing <- temp$testing
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
auc_mcm(fit, model_select = "cAIC")
auc_mcm(fit, newdata = testing)
```

`coef.mixturecure` *Extract model coefficients from a fitted mixturecure object*

Description

`coef.mixturecure` is a generic function which extracts the model coefficients from a fitted `mixturecure` model object fit using `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.

Usage

```
## S3 method for class 'mixturecure'
coef(object, model_select = "AIC", ...)
```

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.
`model_select`	either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are: • `"AIC"` for the minimum AIC (default).

- `"mAIC"` for the minimum modified AIC.
- `"cAIC"` for the minimum corrected AIC.
- `"BIC"`, for the minimum BIC.
- `"mBIC"` for the minimum modified BIC.
- `"EBIC"` for the minimum extended BIC.
- `"logLik"` for the step that maximizes the log-likelihood.
- `n` where n is any numeric value from the solution path.

This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.

`...`	other arguments.

Value

`rate`	estimated rate parameter when fitting a Weibull or exponential mixture cure model.
`shape`	estimated shape parameter when fitting a Weibull mixture cure model.
`b0`	estimated intercept for the incidence portion of the mixture cure model.
`beta_inc`	the vector of coefficient estimates for the incidence portion of the mixture cure model.
`beta_lat`	the vector of coefficient estimates for the latency portion of the mixture cure model.
`p_uncured`	a vector of probabilities from the incidence portion of the fitted model representing the P(uncured).

See Also

`curegmifs`, `cureem`, `summary.mixturecure`, `plot.mixturecure`, `predict.mixturecure`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
coef(fit)
```

`concordance_mcm` *C-statistic for mixture cure models*

Description

This function calculates the C-statistic using the cure status weighting (CSW) method proposed by Asano and Hirakawa (2017).

Usage

```
concordance_mcm(object, newdata, cure_cutoff = 5, model_select = "AIC")
```

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.
`newdata`	an optional data.frame that minimally includes the incidence and/or latency variables to use for predicting the response. If omitted, the training data are used.
`cure_cutoff`	cutoff value for cure, used to produce a proxy for the unobserved cure status (default is 5 representing 5 years). Users should be careful to note the time scale of their data and adjust this according to the time scale and clinical application.
`model_select`	either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are: • `"AIC"` for the minimum AIC (default). • `"mAIC"` for the minimum modified AIC. • `"cAIC"` for the minimum corrected AIC. • `"BIC"`, for the minimum BIC. • `"mBIC"` for the minimum modified BIC. • `"EBIC"` for the minimum extended BIC. • `"logLik"` for the step that maximizes the log-likelihood. • `model_select = n` where `n` is any numeric value from the solution path. This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.

Value

value of C-statistic for the cure models.

References

Asano, J. and Hirakawa, H. (2017) Assessing the prediction accuracy of a cure model for censored survival data with long-term survivors: Application to breast cancer data. *Journal of Biopharmaceutical Statistics*, **27**:6, 918–932.

See Also

auc_mcm

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
testing <- temp$testing
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
concordance_mcm(fit, model_select = "cAIC")
concordance_mcm(fit, newdata = testing, model_select = "cAIC")
```

`cure_estimate` *Estimate cured fraction*

Description

Estimates the cured fraction using a Kaplan-Meier fitted object.

Usage

```
cure_estimate(object)
```

Arguments

`object`	a `survfit` object.

Value

estimated proportion of cured observations

See Also

survfit, sufficient_fu_test, nonzerocure_test

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
km_fit <- survfit(Surv(Time, Censor) ~ 1, data = training)
cure_estimate(km_fit)
```

`cureem` *Fit penalized mixture cure model using the E-M algorithm*

Description

Fits penalized parametric and semi-parametric mixture cure models (MCM) using the E-M algorithm with user-specified penalty parameters. The lasso (L1), MCP, and SCAD penalty are supported for the Cox MCM while only lasso is currently supported for parametric MCMs.

Usage

```
cureem(
  formula,
  data,
  subset,
  x_latency = NULL,
  model = c("cox", "weibull", "exponential"),
  penalty = c("lasso", "MCP", "SCAD"),
  penalty_factor_inc = NULL,
  penalty_factor_lat = NULL,
  thresh = 0.001,
  scale = TRUE,
  maxit = NULL,
  inits = NULL,
  lambda_inc = 0.1,
  lambda_lat = 0.1,
  gamma_inc = 3,
  gamma_lat = 3,
  na.action = na.omit,
  ...
)
```

Arguments

`formula`	an object of class "`formula`" (or one that can be coerced to that class): a symbolic description of the model to be fitted. The response must be a survival object as returned by the `Surv` function while the variables on the right side of the formula are the covariates that are included in the incidence portion of the model.
`data`	a data.frame in which to interpret the variables named in the `formula` or in the `subset` argument. Rows with missing data are omitted (only `na.action = na.omit` is operational) therefore users may want to impute missing data prior to calling this function.
`subset`	an optional expression indicating which subset of observations to be used in the fitting process, either a numeric or factor variable should be used in subset, not a character variable. All observations are included by default.

`x_latency`	specifies the variables to be included in the latency portion of the model and can be either a matrix of predictors, a model formula with the right hand side specifying the latency variables, or the same data.frame passed to the `data` parameter. Note that when using the model formula syntax for `x_latency` it cannot handle `x_latency = ~ .`.
`model`	type of regression model to use for the latency portion of mixture cure model. Can be "cox", "weibull", or "exponential" (default is "cox").
`penalty`	type of penalty function. Can be "lasso", "MCP", or "SCAD" (default is "lasso").
`penalty_factor_inc`	vector of binary indicators representing the penalty to apply to each incidence coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all incidence variables.
`penalty_factor_lat`	vector of binary indicators representing the penalty to apply to each latency coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all latency variables.
`thresh`	small numeric value. The iterative process stops when the differences between successive expected penalized complete-data log-likelihoods for both incidence and latency components are less than this specified level of tolerance (default is 10^-3).
`scale`	logical, if TRUE the predictors are centered and scaled.
`maxit`	integer specifying the maximum number of passes over the data for each lambda. If not specified, 100 is applied when `penalty = "lasso"` and 1000 is applied when `penalty = "MCP"` or `penalty = "SCAD"`.
`inits`	an optional list specifiying the initial values. This includes: • `itct` the incidence intercept. • `b_u` a numeric vector for the unpenalized incidence coefficients for the incidence portion of the model. • `beta_u` a numeric vector for unpenalized latency coefficients in the incidence portion of the model. • `lambda` a numeric value for the rate parameter when fitting either a Weibull or exponential MCM using `model = "weibull"` or `model = "exponential"`. • `alpha` a numeric value for the shape parameter when fitting a Weibull MCM using `model = "weibull"`. • `survprob` a numeric vector for the latency survival probabilities $S_u(t_i \mid w_i)$ for i=1,...,N when fitting a Cox MCM `model = "cox"`. Penalized coefficients are initialized to zero. If `inits` is not specified or improperly specified, initialization is automatically provided by the function.
`lambda_inc`	numeric value for the penalization parameter λ for variables in the incidence portion of the model.
`lambda_lat`	numeric value for the penalization parameter λ for variables in the latency portion of the model.
`gamma_inc`	numeric value for the penalization parameter γ for variables in the incidence portion of the model when `penalty = "MCP"` or `penalty = "SCAD"` (default is 3).

`gamma_lat`	numeric value for the penalization parameter γ for variables in the latency portion of the model when `penalty = "MCP"` or `penalty = "SCAD"` (default is 3).
`na.action`	this function requires complete data so `"na.omit"` is invoked. Users can impute missing data as an alternative prior to model fitting.
`...`	additional arguments.

Value

`b_path`	Matrix representing the solution path of the coefficients in the incidence portion of the model. Row is step and column is variable.
`beta_path`	Matrix representing the solution path of the coefficients in the latency portion of the model. Row is step and column is variable.
`b0_path`	Vector representing the solution path of the intercept in the incidence portion of the model.
`logLik_inc`	Vector representing the expected penalized complete-data log-likelihood for the incidence portion of the model for each step in the solution path.
`logLik_lat`	Vector representing the expected penalized complete-data log-likelihood for the latency portion of the model for each step in the solution path.
`x_incidence`	Matrix representing the design matrix of the incidence predictors.
`x_latency`	Matrix representing the design matrix of the latency predictors.
`y`	Vector representing the survival object response as returned by the `Surv` function
`model`	Character string indicating the type of regression model used for the latency portion of mixture cure model ("weibull" or "exponential").
`scale`	Logical value indicating whether the predictors were centered and scaled.
`method`	Character string indicating the EM algorithm was used in fitting the mixture cure model.
`rate_path`	Vector representing the solution path of the rate parameter for the Weibull or exponential density in the latency portion of the model.
`alpha_path`	Vector representing the solution path of the shape parameter for the Weibull density in the latency portion of the model.
`call`	the matched call.

References

Archer, K. J., Fu, H., Mrozek, K., Nicolet, D., Mims, A. S., Uy, G. L., Stock, W., Byrd, J. C., Hiddemann, W., Braess, J., Spiekermann, K., Metzeler, K. H., Herold, T., Eisfeld, A.-K. (2024) Identifying long-term survivors and those at higher or lower risk of relapse among patients with cytogenetically normal acute myeloid leukemia using a high-dimensional mixture cure model. *Journal of Hematology & Oncology*, **17**:28.

See Also

`cv_cureem`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 80, j = 100, n_true = 10, a = 1.8)
training <- temp$training
fit <- cureem(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "cox", penalty = "lasso", lambda_inc = 0.1,
  lambda_lat = 0.1, gamma_inc = 6, gamma_lat = 10
)
```

`curegmifs` *Fit penalized parametric mixture cure model using the GMIFS algorithm*

Description

Fits a penalized Weibull or exponential mixture cure model using the generalized monotone incremental forward stagewise (GMIFS) algorithm (Hastie et al 2007) and yields solution paths for parameters in the incidence and latency portions of the model.

Usage

```
curegmifs(
  formula,
  data,
  subset,
  x_latency = NULL,
  model = c("weibull", "exponential"),
  penalty_factor_inc = NULL,
  penalty_factor_lat = NULL,
  epsilon = 0.001,
  thresh = 1e-05,
  scale = TRUE,
  maxit = 10000,
  inits = NULL,
  verbose = TRUE,
  suppress_warning = FALSE,
  na.action = na.omit,
  ...
)
```

Arguments

`formula`	an object of class "`formula`" (or one that can be coerced to that class): a symbolic description of the model to be fitted. The response must be a survival object as returned by the `Surv` function while the variables on the right side of

the formula are the covariates that are included in the incidence portion of the model.

`data`	a data.frame in which to interpret the variables named in the `formula` or in the `subset` argument. Rows with missing data are omitted (only `na.action = na.omit` is operational) therefore users may want to impute missing data prior to calling this function.
`subset`	an optional expression indicating which subset of observations to be used in the fitting process, either a numeric or factor variable should be used in subset, not a character variable. All observations are included by default.
`x_latency`	specifies the variables to be included in the latency portion of the model and can be either a matrix of predictors, a model formula with the right hand side specifying the latency variables, or the same data.frame passed to the `data` parameter. Note that when using the model formula syntax for `x_latency` it cannot handle `x_latency = ~ .`.
`model`	type of regression model to use for the latency portion of mixture cure model. Can be "weibull" or "exponential"; default is "weibull".
`penalty_factor_inc`	vector of binary indicators representing the penalty to apply to each incidence coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all incidence variables.
`penalty_factor_lat`	vector of binary indicators representing the penalty to apply to each latency coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all latency variables.
`epsilon`	small numeric value reflecting the incremental value used to update a coefficient at a given step (default is 0.001).
`thresh`	small numeric value. The iterative process stops when the differences between successive expected penalized log-likelihoods for both incidence and latency components are less than this specified level of tolerance (default is 10^-5).
`scale`	logical, if TRUE the predictors are centered and scaled.
`maxit`	integer specifying the maximum number of steps to run in the iterative algorithm (default is 10^4).
`inits`	an optional list specifying the initial values as follows: • `itct` a numeric value for the unpenalized incidence intercept. • `b_u` a numeric vector for the unpenalized incidence coefficients. • `beta_u` a numeric vector for unpenalized latency coefficients. • `lambda` a numeric value for the rate parameter. • `alpha` a numeric value for the shape parameter when `model = "weibull"`. If not supplied or improperly supplied, initialization is automatically provided by the function.
`verbose`	logical, if TRUE running information is printed to the console (default is FALSE).
`suppress_warning`	logical, if TRUE, suppresses echoing the warning that the maximum number of iterations was reached so that the algorithm may not have converged. Instead, warning is returned as part of the output with this message.

`na.action`	this function requires complete data so `"na.omit"` is invoked. Users can impute missing data as an alternative prior to model fitting.
`...`	additional arguments.

Value

`b_path`	Matrix representing the solution path of the coefficients in the incidence portion of the model. Row is step and column is variable.
`beta_path`	Matrix representing the solution path of the coefficients in the latency portion of the model. Row is step and column is variable.
`b0_path`	Vector representing the solution path of the intercept in the incidence portion of the model.
`rate_path`	Vector representing the solution path of the rate parameter for the Weibull or exponential density in the latency portion of the model.
`logLik`	Vector representing the log-likelihood for each step in the solution path.
`x_incidence`	Matrix representing the design matrix of the incidence predictors.
`x_latency`	Matrix representing the design matrix of the latency predictors.
`y`	Vector representing the survival object response as returned by the `Surv` function
`model`	Character string indicating the type of regression model used for the latency portion of mixture cure model ("weibull" or "exponential").
`scale`	Logical value indicating whether the predictors were centered and scaled.
`alpha_path`	Vector representing the solution path of the shape parameter for the Weibull density in the latency portion of the model.
`call`	the matched call.
`warning`	message indicating whether the maximum number of iterations was achieved which may indicate the model did not converge.

References

Fu, H., Nicolet, D., Mrozek, K., Stone, R. M., Eisfeld, A. K., Byrd, J. C., Archer, K. J. (2022) Controlled variable selection in Weibull mixture cure models for high-dimensional data. *Statistics in Medicine*, **41**(22), 4340–4366.

Hastie, T., Taylor J., Tibshirani R., Walther G. (2007) Forward stagewise regression and the monotone lasso. *Electron J Stat*, **1**:1–29.

See Also

`cv_curegmifs`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
```

```
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000, epsilon = 0.01,
  verbose = FALSE
)
```

---

cv_cureem *Fit penalized mixture cure model using the E-M algorithm with cross-validation for parameter tuning*

Description

Fits penalized parametric and semi-parametric mixture cure models (MCM) using the E-M algorithm with with k-fold cross-validation for parameter tuning. The lasso (L1), MCP and SCAD penalty are supported for the Cox MCM while only lasso is currently supported for parametric MCMs. When FDR controlled variable selection is used, the model-X knockoffs method is applied and indices of selected variables are returned.

Usage

```
cv_cureem(
  formula,
  data,
  subset,
  x_latency = NULL,
  model = c("cox", "weibull", "exponential"),
  penalty = c("lasso", "MCP", "SCAD"),
  penalty_factor_inc = NULL,
  penalty_factor_lat = NULL,
  fdr_control = FALSE,
  fdr = 0.2,
  grid_tuning = FALSE,
  thresh = 0.001,
  scale = TRUE,
  maxit = NULL,
  inits = NULL,
  lambda_inc_list = NULL,
  lambda_lat_list = NULL,
  nlambda_inc = NULL,
  nlambda_lat = NULL,
  gamma_inc = 3,
  gamma_lat = 3,
  lambda_min_ratio_inc = 0.1,
  lambda_min_ratio_lat = 0.1,
  n_folds = 5,
  measure_inc = c("c", "auc"),
```

```
  one_se = FALSE,
  cure_cutoff = 5,
  parallel = FALSE,
  seed = NULL,
  verbose = TRUE,
  na.action = na.omit,
  ...
)
```

Arguments

`formula`	an object of class "`formula`" (or one that can be coerced to that class): a symbolic description of the model to be fitted. The response must be a survival object as returned by the `Surv` function while the variables on the right side of the formula are the covariates that are included in the incidence portion of the model.
`data`	a data.frame in which to interpret the variables named in the `formula` or in the `subset` argument. Rows with missing data are omitted (only `na.action = na.omit` is operational) therefore users may want to impute missing data prior to calling this function.
`subset`	an optional expression indicating which subset of observations to be used in the fitting process, either a numeric or factor variable should be used in subset, not a character variable. All observations are included by default.
`x_latency`	specifies the variables to be included in the latency portion of the model and can be either a matrix of predictors, a model formula with the right hand side specifying the latency variables, or the same data.frame passed to the `data` parameter. Note that when using the model formula syntax for `x_latency` it cannot handle `x_latency = ~ .`.
`model`	type of regression model to use for the latency portion of mixture cure model. Can be "cox", "weibull", or "exponential" (default is "cox").
`penalty`	type of penalty function. Can be "lasso", "MCP", or "SCAD" (default is "lasso").
`penalty_factor_inc`	vector of binary indicators representing the penalty to apply to each incidence coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all incidence variables.
`penalty_factor_lat`	vector of binary indicators representing the penalty to apply to each latency coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all latency variables.
`fdr_control`	logical, if TRUE, model-X knockoffs are used for FDR-controlled variable selection and indices of selected variables are returned (default is FALSE).
`fdr`	numeric value in (0, 1) range specifying the target FDR level to use for variable selection when `fdr_control = TRUE` (default is 0.2).
`grid_tuning`	logical, if TRUE a 2-D grid tuning approach is used to select the optimal pair of λ_b and λ_β penalty parameters for the incidence and latency portions of the model, respectively. Otherwise the λ_b and λ_β are selected from a 1-D sequence and are equal to one another (default is FALSE).

`thresh`	small numeric value. The iterative process stops when the differences between successive expected penalized complete-data log-likelihoods for both incidence and latency components are less than this specified level of tolerance (default is 10^-3).
`scale`	logical, if TRUE the predictors are centered and scaled.
`maxit`	maximum number of passes over the data for each lambda. If not specified, 100 is applied when `penalty = "lasso"` and 1000 is applied when `penalty = "MCP"` or `penalty = "SCAD"`.
`inits`	an optional list specifiying the initial values to be used for model fitting as follows: • `itct` the incidence intercept. • `b_u` a numeric vector for the unpenalized. incidence coefficients for the incidence portion of the model. • `beta_u` a numeric vector for unpenalized latency coefficients in the incidence portion of the model. • `lambda` a numeric value for the rate parameter when fitting either a Weibull or exponential MCM using `model = "weibull"` or `model = "exponential"`. • `alpha` a numeric value for the shape parameter when fitting a Weibull MCM using `model = "weibull"`. • `survprob` a numeric vector for the latency survival probabilities $S_u(t_i\|w_i)$ for i=1,...,N when fitting a Cox MCM `model = "cox"`. Penalized coefficients are initialized to zero. If `inits` is not specified or improperly specified, initialization is automatically provided by the function.
`lambda_inc_list`	a numeric vector used to search for the optimal λ_b tuning parameter. If not supplied, the function computes a λ_b sequence based on `nlambda_inc` and `lambda_min_ratio_inc`. If `grid_tuning = FALSE`, the same sequence should be used for both λ_b and λ_β.
`lambda_lat_list`	a numeric vector used to search for the optimal λ_β tuning parameter. If not supplied, the function computes a λ_β sequence based on `nlambda_lat` and `lambda_min_ratio_lat`. If `grid_tuning = FALSE`, the same sequence should be used for both λ_b and λ_β.
`nlambda_inc`	an integer specifying the number of values to search for the optimal λ_b tuning parameter; default is 10 if `grid_tuning = TRUE` and 50 otherwise.
`nlambda_lat`	an integer specifying the number of values to search for the optimal λ_β tuning parameter; default is 10 if `grid_tuning = TRUE` and 50 otherwise.
`gamma_inc`	numeric value for the penalization parameter γ for variables in the incidence portion of the model when `penalty = "MCP"` or `penalty = "SCAD"` (default is 3).
`gamma_lat`	numeric value for the penalization parameter γ for variables in the latency portion of the model when `penalty = "MCP"` or `penalty = "SCAD"` (default is 3).
`lambda_min_ratio_inc`	numeric value in (0,1) representing the smallest value for λ_b as a fraction of `lambda.max_inc`, the data-derived entry value at which essentially all penalized variables in the incidence portion of the model have a coefficient estimate of 0 (default is 0.1).

`lambda_min_ratio_lat`	numeric value in (0.1) representing the smallest value for λ_β as a fraction of `lambda.max_lat`, the data-derived entry value at essentially all penalized variables in the latency portion of the model have a coefficient estimate of 0 (default is 0.1).
`n_folds`	an integer specifying the number of folds for the k-fold cross-valiation procedure (default is 5).
`measure_inc`	character string specifying the evaluation criterion used in selecting the optimal λ_b which can be either - `"c"` specifying to use the C-statistic for cure status weighting (CSW) method proposed by Asano and Hirakawa (2017) to select both λ_b and λ_β - `"auc"` specifying to use the AUC for cure prediction using the mean score imputation (MSI) method proposed by Asano et al. (2014) to select λ_b while the C-statistic with CSW is used for λ_β.
`one_se`	logical, if TRUE then the one standard error rule is applied for selecting the optimal parameters. The one standard error rule selects the most parsimonious model having evaluation criterion no more than one standard error worse than that of the best evaluation criterion (default is FALSE).
`cure_cutoff`	numeric value representing the cutoff time value that represents subjects not experiencing the event by this time are cured. This value is used to produce a proxy for the unobserved cure status when calculating C-statistic and AUC (default is 5 representing 5 years). Users should be careful to note the time scale of their data and adjust this according to the time scale and clinical application.
`parallel`	logical. If TRUE, parallel processing is performed for K-fold CV using `foreach` and the **doParallel** package is required.
`seed`	optional integer representing the random seed. Setting the random seed fosters reproducibility of the results.
`verbose`	logical, if TRUE running information is printed to the console (default is FALSE).
`na.action`	this function requires complete data so `"na.omit"` is invoked. Users can impute missing data as an alternative prior to model fitting.
`...`	additional arguments.

Value

`b0`	Estimated intercept for the incidence portion of the model.
`b`	Estimated coefficients for the incidence portion of the model.
`beta`	Estimated coefficients for the latency portion of the model.
`alpha`	Estimated shape parameter if the Weibull model is fit.
`rate`	Estimated rate parameter if the Weibull or exponential model is fit.
`logLik_inc`	Expected penalized complete-data log-likelihood for the incidence portion of the model.
`logLik_lat`	Expected penalized complete-data log-likelihood for the latency portion of the model.

`selected_lambda_inc`	Value of λ_b selected using cross-validation. NULL when fdr_control is TRUE.
`selected_lambda_lat`	Value of λ_β selected using cross-validation. NULL when fdr_control is TRUE.
`max_c`	Maximum C-statistic achieved.
`max_auc`	Maximum AUC for cure prediction achieved; only output when `measure_inc="auc"`.
`selected_index_inc`	Indices of selected variables for the incidence portion of the model when `fdr_control=TRUE`. If no variables are selected, `int(0)` will be returned.
`selected_index_lat`	Indices of selected variables for the latency portion of the model when `fdr_control=TRUE`. If no variables are selected, `int(0)` will be returned.
`call`	the matched call.

References

Archer, K. J., Fu, H., Mrozek, K., Nicolet, D., Mims, A. S., Uy, G. L., Stock, W., Byrd, J. C., Hiddemann, W., Braess, J., Spiekermann, K., Metzeler, K. H., Herold, T., Eisfeld, A.-K. (2024) Identifying long-term survivors and those at higher or lower risk of relapse among patients with cytogenetically normal acute myeloid leukemia using a high-dimensional mixture cure model. *Journal of Hematology & Oncology*, **17**:28.

See Also

`cureem`

Examples

```
## Not run:
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 200, j = 25, n_true = 5, a = 1.8, rho = 0.2)
training <- temp$training
# Fit a penalized Cox MCM selecting parameters using 2-fold CV
fit.cv <- cv_cureem(Surv(Time, Censor) ~ .,
  data = training,
  x_latency = training, fdr_control = FALSE,
  grid_tuning = FALSE, nlambda_inc = 10, nlambda_lat = 10,
  n_folds = 2, seed = 23, verbose = FALSE
)
fit.cv.fdr <- cv_cureem(Surv(Time, Censor) ~ .,
   data = training,
   x_latency = training, model = "weibull", penalty = "lasso",
   fdr_control = TRUE, grid_tuning = FALSE, nlambda_inc = 10,
   nlambda_lat = 10, n_folds = 2, seed = 23, verbose = TRUE
   )

## End(Not run)
```

`cv_curegmifs` *Fit a penalized parametric mixture cure model using the generalized monotone incremental forward stagewise (GMIFS) algorithm (Hastie et al 2007) with cross-validation for model selection*

Description

Fits a penalized Weibull or exponential mixture cure model using the generalized monotone incremental forward stagewise (GMIFS) algorithm with k-fold cross-validation to select the optimal iteration step along the solution path. When FDR controlled variable selection is used, the model-X knockoffs method is applied and indices of selected variables are returned.

Usage

```
cv_curegmifs(
  formula,
  data,
  subset,
  x_latency = NULL,
  model = c("weibull", "exponential"),
  penalty_factor_inc = NULL,
  penalty_factor_lat = NULL,
  fdr_control = FALSE,
  fdr = 0.2,
  epsilon = 0.001,
  thresh = 1e-05,
  scale = TRUE,
  maxit = 10000,
  inits = NULL,
  n_folds = 5,
  measure_inc = c("c", "auc"),
  one_se = FALSE,
  cure_cutoff = 5,
  parallel = FALSE,
  seed = NULL,
  verbose = TRUE,
  na.action = na.omit,
  ...
)
```

Arguments

`formula`	an object of class "`formula`" (or one that can be coerced to that class): a symbolic description of the model to be fitted. The response must be a survival object as returned by the `Surv` function while the variables on the right side of the formula are the covariates that are included in the incidence portion of the model.

`data`	a data.frame in which to interpret the variables named in the `formula` or in the `subset` argument. Rows with missing data are omitted (only `na.action = na.omit` is operational) therefore users may want to impute missing data prior to calling this function.
`subset`	an optional expression indicating which subset of observations to be used in the fitting process, either a numeric or factor variable should be used in subset, not a character variable. All observations are included by default.
`x_latency`	specifies the variables to be included in the latency portion of the model and can be either a matrix of predictors, a model formula with the right hand side specifying the latency variables, or the same data.frame passed to the `data` parameter. Note that when using the model formula syntax for `x_latency` it cannot handle `x_latency = ~ .`.
`model`	type of regression model to use for the latency portion of mixture cure model. Can be "weibull" or "exponential"; default is "weibull".
`penalty_factor_inc`	vector of binary indicators representing the penalty to apply to each incidence coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all incidence variables.
`penalty_factor_lat`	vector of binary indicators representing the penalty to apply to each latency coefficient: 0 implies no shrinkage and 1 implies shrinkage. If not supplied, 1 is applied to all latency variables.
`fdr_control`	logical, if TRUE, model-X knockoffs are used for FDR-controlled variable selection and indices of selected variables are returned (default is FALSE).
`fdr`	numeric value in (0, 1) range specifying the target FDR level to use for variable selection when `fdr_control = TRUE` (default is 0.2).
`epsilon`	small numeric value reflecting incremental value used to update a coefficient at a given step (default is 0.001).
`thresh`	small numeric value. The iterative process stops when the differences between successive expected penalized complete-data log-likelihoods for both incidence and latency components are less than this specified level of tolerance (default is 10^-5).
`scale`	logical, if TRUE the predictors are centered and scaled.
`maxit`	integer specifying the maximum number of steps to run in the iterative algorithm (default is 10^4).
`inits`	an optional list specifying the initial values as follows: • `itct` a numeric value for the unpenalized incidence intercept. • `b_u` a numeric vector for the unpenalized incidence coefficients. • `beta_u` a numeric vector for unpenalized latency coefficients. • `lambda` a numeric value for the rate parameter. • `alpha` a numeric value for the shape parameter when `model = "weibull"`. If `inits` is not specified or improperly supplied, initialization is automatically provided by the function.

`n_folds`	an integer specifying the number of folds for the k-fold cross-validation procedure (default is 5).
`measure_inc`	character string specifying the evaluation criterion used in selecting the optimal λ_b which can be either • `"c"` specifying to use the C-statistic for cure status weighting (CSW) method proposed by Asano and Hirakawa (2017) to select both λ_b and λ_β • `"auc"` specifying to use the AUC for cure prediction using the mean score imputation (MSI) method proposed by Asano et al. (2014) to select λ_b while the C-statistic with CSW is used for λ_β.
`one_se`	logical, if TRUE then the one standard error rule is applied for selecting the optimal parameters. The one standard error rule selects the most parsimonious model having evaluation criterion no more than one standard error worse than that of the best evaluation criterion (default is FALSE).
`cure_cutoff`	numeric value representing the cutoff time value that represents subjects not experiencing the event by this time are cured. This value is used to produce a proxy for the unobserved cure status when calculating C-statistic and AUC (default is 5 representing 5 years). Users should be careful to note the time scale of their data and adjust this according to the time scale and clinical application.
`parallel`	logical. If TRUE, parallel processing is performed for K-fold CV using `foreach` and the **doParallel** package is required.
`seed`	optional integer representing the random seed. Setting the random seed fosters reproducibility of the results.
`verbose`	logical, if TRUE running information is printed to the console (default is FALSE).
`na.action`	this function requires complete data so `"na.omit"` is invoked. Users can impute missing data as an alternative prior to model fitting.
`...`	additional arguments.

Value

`b0`	Estimated intercept for the incidence portion of the model.
`b`	Estimated coefficients for the incidence portion of the model.
`beta`	Estimated coefficients for the latency portion of the model.
`alpha`	Estimated shape parameter if the Weibull model is fit.
`rate`	Estimated rate parameter if the Weibull or exponential model is fit.
`logLik`	Log-likelihood value.
`selected_step_inc`	Iteration step selected for the incidence portion of the model using cross-validation. NULL when fdr_control is TRUE.
`selected_step_lat`	Iteration step selected for the latency portion of the model using cross-validation. NULL when fdr_control is TRUE.
`max_c`	Maximum C-statistic achieved
`max_auc`	Maximum AUC for cure prediction achieved; only output when `measure_inc = "auc"`.

`selected_index_inc`	Indices of selected variables for the incidence portion of the model when `fdr_control = TRUE`. If none selected, `int(0)` will be returned.
`selected_index_lat`	Indices of selected variables for the latency portion of the model when `fdr_control = TRUE`. If none selected, `int(0)` will be returned.
`call`	the matched call.

References

Fu, H., Nicolet, D., Mrozek, K., Stone, R. M., Eisfeld, A. K., Byrd, J. C., Archer, K. J. (2022) Controlled variable selection in Weibull mixture cure models for high-dimensional data. *Statistics in Medicine*, **41**(22), 4340–4366.

Hastie, T., Taylor J., Tibshirani R., Walther G. (2007) Forward stagewise regression and the monotone lasso. *Electron J Stat*, **1**:1–29.

See Also

`curegmifs`

Examples

```
library(survival)
withr::local_seed(123)
temp <- generate_cure_data(n = 100, j = 15, n_true = 3, a = 1.8, rho = 0.2)
training <- temp$training

fit.cv <- cv_curegmifs(Surv(Time, Censor) ~ .,
  data = training,
  x_latency = training, fdr_control = FALSE,
  maxit = 450, epsilon = 0.01, n_folds = 2,
  seed = 23, verbose = TRUE
)
```

`dim.mixturecure` *Dimension method for mixturecure objects*

Description

Dimension method for `mixturecure` objects.

Usage

```
## S3 method for class 'mixturecure'
dim(x)
```

Arguments

`x`	An object of class `mixturecure`.

Value

`nobs`	number of subjects in the dataset.
`p_incidence`	number of variables in the incidence portion of the model.
`p_latency`	number of variables in the latency portion of the model.

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
dim(fit)
```

`family.mixturecure` *Return model family and fitting algorithm for mixturecure model fits*

Description

Return model family and fitting algorithm for`mixturecure` model fits.

Usage

```
## S3 method for class 'mixturecure'
family(object, ...)
```

Arguments

`object`	an object of class `mixturecure`
`...`	other arguments.

Value

the parametric or semi-parametric model fit and the fitting algorithm.

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
```

```
  epsilon = 0.01, verbose = FALSE
)
family(fit)
```

---

formula.mixturecure *Extract model formula for mixturecure object*

Description

Extract the model formula for `mixturecure` object

Usage

```
## S3 method for class 'mixturecure'
formula(x, ...)
```

Arguments

`x`	an object from class `mixturecure`.
`...`	other arguments.

Value

a formula representing the incidence and variables for the latency portion of the model

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
formula(fit)
```

`generate_cure_data` *Simulate data under a mixture cure model*

Description

Simulate data under a mixture cure model.

Usage

```
generate_cure_data(
  n = 400,
  j = 500,
  nonp = 2,
  train_prop = 0.75,
  n_true = 10,
  a = 1,
  rho = 0.5,
  itct_mean = 0.5,
  cens_ub = 20,
  alpha = 1,
  lambda = 2,
  same_signs = FALSE,
  model = "weibull"
)
```

Arguments

`n`	an integer denoting the total sample size.
`j`	an integer denoting the number of penalized predictors which is the same for both the incidence and latency portions of the model.
`nonp`	an integer denoting the number of unpenalized predictors (which is the same for both the incidence and latency portions of the model).
`train_prop`	a numeric value in [0, 1) representing the fraction of `n` to be used in forming the training dataset.
`n_true`	an integer less than `j` denoting the number of variables truly associated with the outcome (i.e., the number of covariates with nonzero parameter values) among the penalized predictors.
`a`	a numeric value denoting the effect size (signal amplitude) which is the same for both the incidence and latency portions of the model.
`rho`	a numeric value in [0, 1) representing the correlation between adjacent covariates in the same block.
`itct_mean`	a numeric value representing the expectation of the incidence intercept which controls the cure rate.
`cens_ub`	a numeric value representing the upper bound on the censoring time distribution which follows a uniform distribution on (0, `cens_ub`].

`alpha`	a numeric value representing the shape parameter in the Weibull density.
`lambda`	a numeric value representing the rate parameter in the Weibull density.
`same_signs`	logical, if TRUE the incidence and latency coefficients have the same signs.
`model`	type of regression model to use for the latency portion of mixture cure model. Can be one of the following: • `"weibull"` to generate times from a Weibull distribution. • `"GG"` to generate times from a generalized gamma distribution. • `"Gompertz"` to generate times from a Gomertz distribution. • `"nonparametric"` to generate times non-parametrically. • `"GG_baseline"` to generate times from a generalized gamma baseline distribution.

Value

`training`	training data.frame which includes Time, Censor, and covariates. Variables prefixed with `"U"` indicates unpenalized covariates and is equal to the value passed to `nonp` (default is 2). Variables prefixed with `"X"` indicates penalized covariates and is equal to the value passed to `j`.
`training_y`	the true status for the training set: uncured = 1; cured = 0
`testing`	testing data.frame which includes Time, Censor, Y (the true uncured = 1; cured = 0 status), and covariates. Variables prefixed with `"U"` indicates unpenalized covariates and is equal to the value passed to `nonp` (default is 2). Variables prefixed with `"X"` indicates penalized covariates and is equal to the value passed to `j`.
`testing_y`	the true status for the testing set: uncured = 1; cured = 0
`parameters`	a list including: the indices of true incidence signals (`nonzero_b`), indices of true latency signals (`nonzero_beta`), unpenalized incidence parameter values (`b_u`), unpenalized latency parameter values (`beta_u`), parameter values for the true incidence signals among penalized covariates (`b_p_nz`), parameter values for the true latency signals among penalized covariates (`beta_p_nz`), parameter value for the incidence intercept (`itct`)

Examples

```
library(survival)
withr::local_seed(1234)
# This dataset has 2 penalized features associated with the outcome,
# 3 penalized features not associated with the outcome (noise features), and 1
# unpenalized noise feature.
data <- generate_cure_data(n = 1000, j = 5, n_true = 2, nonp = 1, a = 2)
# Extract the training data
training <- data$training
# Extract the testing data
testing <- data$testing
# To identify the features truly associated with incidence
names(training)[grep("^X", names(training))][data$parameters$nonzero_b]
# To identify the features truly associated with latency
```

```
names(training)[grep("^X", names(training))][data$parameters$nonzero_beta]
# Fit the model to the training data
fitem <- cureem(Surv(Time, Censor) ~ ., data = training,
  x_latency = training)
# Examine the estimated coefficients at the (default) minimum AIC
coef(fitem)
# As the penalty increases, the coefficients for the noise variables shrink
# to or remain at zero, while the truly associated features have coefficient
# paths that depart from zero. This shows the model's ability to distinguish
# signal from noise.
plot(fitem, label = TRUE)
```

`logLik.mixturecure` *Log-likelihood for fitted mixture cure model*

Description

This function returns the log-likelihood for a user-specified model criterion or step for a `curegmifs`, `cureem`, `cv_curegmifs` or `cv_cureem` fitted object.

Usage

```
## S3 method for class 'mixturecure'
logLik(object, model_select = "AIC", ...)
```

Arguments

`object` a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, `cv_cureem`.

`model_select` either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are:

- `"AIC"` for the minimum AIC (default).
- `"mAIC"` for the minimum modified AIC.
- `"cAIC"` for the minimum corrected AIC.
- `"BIC"`, for the minimum BIC.
- `"mBIC"` for the minimum modified BIC.
- `"EBIC"` for the minimum extended BIC.
- `"logLik"` for the step that maximizes the log-likelihood.
- `n` where `n` is any numeric value from the solution path.

This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.

`...` other arguments.

Value

log-likelihood of the fitted mixture cure model using the specified criteria.

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
logLik(fit, model_select = "AIC")
```

`nobs.mixturecure` *Number of observations in mixturecure object*

Description

Number of observations in fitted `mixturecure` object.

Usage

```
## S3 method for class 'mixturecure'
nobs(object, ...)
```

Arguments

`object`	An object of class `mixturecure`.
`...`	other arguments.

Value

number of subjects in the dataset.

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
nobs(fit)
```

`nonzerocure_test` *Non-parametric test for a non-zero cured fraction*

Description

Tests the null hypothesis that the proportion of observations susceptible to the event = 1 against the alternative that the proportion of observations susceptible to the event is < 1. If the null hypothesis is rejected, there is a significant cured fraction.

Usage

```
nonzerocure_test(object, reps = 1000, seed = NULL, plot = FALSE, b = NULL)
```

Arguments

`object`	a `survfit` object.
`reps`	number of simulations on which to base the p-value (default = 1000).
`seed`	optional random seed.
`plot`	logical. If TRUE a histogram of the estimated susceptible proportions over all simulations is produced.
`b`	optional. If specified the maximum observed time for the uniform distribution for generating the censoring times. If not specified, an exponential model is used for generating the censoring times (default).

Value

`proportion_susceptible`	estimated proportion of susceptibles
`proportion_cured`	estimated proportion of those cured
`p_value`	p-value testing the null hypothesis that the proportion of susceptibles = 1 (cured fraction = 0) against the alternative that the proportion of susceptibles < 1 (non-zero cured fraction)
`time_95_percent_of_events`	estimated time at which 95% of events should have occurred

References

Maller, R. A. and Zhou, X. (1996) *Survival Analysis with Long-Term Survivors*. John Wiley & Sons.

See Also

`survfit`, `cure_estimate`, `sufficient_fu_test`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
km_fit <- survfit(Surv(Time, Censor) ~ 1, data = training)
nonzerocure_test(km_fit)
```

`npar_mixturecure` *Number of parameters in fitted mixture cure model*

Description

This function returns the number of parameters in a user-specified model criterion or step for a `curegmifs`, `cureem`, `cv_curegmifs` or `cv_cureem` fitted object.

Usage

```
npar_mixturecure(object, model_select = "AIC")
```

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, `cv_cureem`.
`model_select`	either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are: • `"AIC"` for the minimum AIC (default). • `"mAIC"` for the minimum modified AIC. • `"cAIC"` for the minimum corrected AIC. • `"BIC"`, for the minimum BIC. • `"mBIC"` for the minimum modified BIC. • `"EBIC"` for the minimum extended BIC. • `"logLik"` for the step that maximizes the log-likelihood. • `n` where `n` is any numeric value from the solution path. This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.

Value

number of paramaters of the fitted mixture cure model using the specified criteria.

`pediatric_flt3` *Pediatric acute myeloid leukemia patients with FLT3-ITD rearrangement data*

Description

Event-free survival for 246 pediatric acute myeloid leukemia patients having a FLT3-ITD rearrangement along with expression of 200 transcripts from RNA-sequencing.

Usage

```
pediatric_flt3
```

Format

A data frame with 246 rows (subjects) and 202 columns:

efs event-free survival

status censoring indicator: 1 = event; 0 = censored

ENSG00000229807.12 normalized expression for indicated transcript

ENSG00000211459.2 normalized expression for indicated transcript

ENSG00000264063.1 normalized expression for indicated transcript

ENSG00000264462.1 normalized expression for indicated transcript

ENSG00000227195.11 normalized expression for indicated transcript

ENSG00000198795.11 normalized expression for indicated transcript

ENSG00000177133.11 normalized expression for indicated transcript

ENSG00000257883.1 normalized expression for indicated transcript

ENSG00000189223.15 normalized expression for indicated transcript

ENSG00000210082.2 normalized expression for indicated transcript

ENSG00000142611.17 normalized expression for indicated transcript

ENSG00000054598.9 normalized expression for indicated transcript

ENSG00000147180.17 normalized expression for indicated transcript

ENSG00000275708.1 normalized expression for indicated transcript

ENSG00000227063.5 normalized expression for indicated transcript

ENSG00000179455.10 normalized expression for indicated transcript

ENSG00000243478.9 normalized expression for indicated transcript

ENSG00000239839.7 normalized expression for indicated transcript

ENSG00000274253.5 normalized expression for indicated transcript

ENSG00000198840.2 normalized expression for indicated transcript

ENSG00000186407.7 normalized expression for indicated transcript

ENSG00000198763.3 normalized expression for indicated transcript

ENSG00000107562.16 normalized expression for indicated transcript

ENSG00000188536.13 normalized expression for indicated transcript

ENSG00000164821.5 normalized expression for indicated transcript

ENSG00000255833.2 normalized expression for indicated transcript

ENSG00000107447.8 normalized expression for indicated transcript

ENSG00000237541.4 normalized expression for indicated transcript

ENSG00000274618.2 normalized expression for indicated transcript

ENSG00000206172.8 normalized expression for indicated transcript

ENSG00000198886.2 normalized expression for indicated transcript

ENSG00000122592.8 normalized expression for indicated transcript

ENSG00000038427.16 normalized expression for indicated transcript

ENSG00000134755.17 normalized expression for indicated transcript

ENSG00000248527.1 normalized expression for indicated transcript

ENSG00000273018.7 normalized expression for indicated transcript

ENSG00000261377.5 normalized expression for indicated transcript

ENSG00000074964.17 normalized expression for indicated transcript

ENSG00000118113.12 normalized expression for indicated transcript

ENSG00000086548.9 normalized expression for indicated transcript

ENSG00000172037.14 normalized expression for indicated transcript

ENSG00000198744.5 normalized expression for indicated transcript

ENSG00000198899.2 normalized expression for indicated transcript

ENSG00000186818.12 normalized expression for indicated transcript

ENSG00000154864.13 normalized expression for indicated transcript

ENSG00000227081.5 normalized expression for indicated transcript

ENSG00000101425.14 normalized expression for indicated transcript

ENSG00000158473.8 normalized expression for indicated transcript

ENSG00000102362.15 normalized expression for indicated transcript

ENSG00000212907.2 normalized expression for indicated transcript

ENSG00000121807.6 normalized expression for indicated transcript

ENSG00000138829.12 normalized expression for indicated transcript

ENSG00000185052.12 normalized expression for indicated transcript

ENSG00000111057.11 normalized expression for indicated transcript

ENSG00000228253.1 normalized expression for indicated transcript

ENSG00000188153.14 normalized expression for indicated transcript

ENSG00000166681.14 normalized expression for indicated transcript

ENSG00000198888.2 normalized expression for indicated transcript

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ENSG00000230076.1	normalized expression for indicated transcript
ENSG00000260314.3	normalized expression for indicated transcript
ENSG00000187164.20	normalized expression for indicated transcript
ENSG00000212443.1	normalized expression for indicated transcript
ENSG00000251692.8	normalized expression for indicated transcript
ENSG00000198727.2	normalized expression for indicated transcript
ENSG00000131401.11	normalized expression for indicated transcript
ENSG00000225345.3	normalized expression for indicated transcript
ENSG00000197253.13	normalized expression for indicated transcript
ENSG00000163737.4	normalized expression for indicated transcript
ENSG00000204010.3	normalized expression for indicated transcript
ENSG00000174403.16	normalized expression for indicated transcript
ENSG00000249790.3	normalized expression for indicated transcript
ENSG00000239559.2	normalized expression for indicated transcript
ENSG00000142192.21	normalized expression for indicated transcript
ENSG00000151650.8	normalized expression for indicated transcript
ENSG00000198712.1	normalized expression for indicated transcript
ENSG00000124469.12	normalized expression for indicated transcript
ENSG00000105854.13	normalized expression for indicated transcript
ENSG00000197629.6	normalized expression for indicated transcript
ENSG00000278588.2	normalized expression for indicated transcript
ENSG00000196565.15	normalized expression for indicated transcript
ENSG00000241743.4	normalized expression for indicated transcript
ENSG00000174059.17	normalized expression for indicated transcript
ENSG00000150760.13	normalized expression for indicated transcript
ENSG00000133742.14	normalized expression for indicated transcript
ENSG00000210196.2	normalized expression for indicated transcript
ENSG00000204860.5	normalized expression for indicated transcript
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ENSG00000106080.11	normalized expression for indicated transcript
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ENSG00000248334.6	normalized expression for indicated transcript
ENSG00000169908.12	normalized expression for indicated transcript
ENSG00000171509.16	normalized expression for indicated transcript
ENSG00000008438.5	normalized expression for indicated transcript
ENSG00000130147.16	normalized expression for indicated transcript

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ENSG00000165092.13 normalized expression for indicated transcript
ENSG00000170458.14 normalized expression for indicated transcript
ENSG00000070182.21 normalized expression for indicated transcript
ENSG00000157445.15 normalized expression for indicated transcript
ENSG00000206737.1 normalized expression for indicated transcript
ENSG00000225217.1 normalized expression for indicated transcript
ENSG00000198336.9 normalized expression for indicated transcript
ENSG00000177469.13 normalized expression for indicated transcript
ENSG00000198695.2 normalized expression for indicated transcript
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ENSG00000223609.11 normalized expression for indicated transcript
ENSG00000206177.7 normalized expression for indicated transcript
ENSG00000164684.13 normalized expression for indicated transcript
ENSG00000197632.9 normalized expression for indicated transcript

ENSG00000177575.13 normalized expression for indicated transcript

ENSG00000004939.15 normalized expression for indicated transcript

ENSG00000121053.6 normalized expression for indicated transcript

ENSG00000259207.9 normalized expression for indicated transcript

ENSG00000158352.16 normalized expression for indicated transcript

ENSG00000198046.12 normalized expression for indicated transcript

ENSG00000225630.1 normalized expression for indicated transcript

ENSG00000070808.16 normalized expression for indicated transcript

ENSG00000012223.13 normalized expression for indicated transcript

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ENSG00000272398.6 normalized expression for indicated transcript

ENSG00000116667.15 normalized expression for indicated transcript

ENSG00000101916.12 normalized expression for indicated transcript

ENSG00000184293.7 normalized expression for indicated transcript

ENSG00000244040.7 normalized expression for indicated transcript

ENSG00000229344.1 normalized expression for indicated transcript

ENSG00000142405.22 normalized expression for indicated transcript

ENSG00000171051.9 normalized expression for indicated transcript

ENSG00000183023.18 normalized expression for indicated transcript

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ENSG00000244682.7 normalized expression for indicated transcript

ENSG00000166025.18 normalized expression for indicated transcript

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ENSG00000104918.8 normalized expression for indicated transcript

ENSG00000287431.1 normalized expression for indicated transcript

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ENSG00000164047.6 normalized expression for indicated transcript

ENSG00000137571.11 normalized expression for indicated transcript

ENSG00000143416.21 normalized expression for indicated transcript

ENSG00000253578.1 normalized expression for indicated transcript

ENSG00000120708.17	normalized expression for indicated transcript
ENSG00000220842.6	normalized expression for indicated transcript
ENSG00000110077.14	normalized expression for indicated transcript
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ENSG00000121316.11	normalized expression for indicated transcript
ENSG00000123689.6	normalized expression for indicated transcript
ENSG00000198938.2	normalized expression for indicated transcript
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ENSG00000233101.10	normalized expression for indicated transcript
ENSG00000163220.11	normalized expression for indicated transcript
ENSG00000100985.7	normalized expression for indicated transcript
ENSG00000166947.15	normalized expression for indicated transcript
ENSG00000275713.2	normalized expression for indicated transcript
ENSG00000143546.10	normalized expression for indicated transcript
ENSG00000136929.13	normalized expression for indicated transcript
ENSG00000103313.13	normalized expression for indicated transcript
ENSG00000186529.16	normalized expression for indicated transcript
ENSG00000109321.11	normalized expression for indicated transcript
ENSG00000104177.18	normalized expression for indicated transcript
ENSG00000257335.8	normalized expression for indicated transcript
ENSG00000197993.9	normalized expression for indicated transcript
ENSG00000134460.18	normalized expression for indicated transcript
ENSG00000211892.4	normalized expression for indicated transcript
ENSG00000180044.5	normalized expression for indicated transcript
ENSG00000268555.2	normalized expression for indicated transcript
ENSG00000266402.3	normalized expression for indicated transcript
ENSG00000163221.9	normalized expression for indicated transcript
ENSG00000225101.6	normalized expression for indicated transcript
ENSG00000197249.14	normalized expression for indicated transcript
ENSG00000093134.15	normalized expression for indicated transcript

Source

`https://www.cancer.gov/ccg/research/genome-sequencing/target/studied-cancers/acute-myeloid-leukemia`
`targets-aml-project`

`plot.mixturecure` *Plot fitted mixture cure model*

Description

This function plots either the coefficient path, the AIC, the cAIC, the BIC, or the log-likelihood for a fitted `curegmifs` or `cureem` object. This function produces a lollipop plot of the coefficient estimates for a fitted `cv_curegmifs` or `cv_cureem` object.

Usage

```
## S3 method for class 'mixturecure'
plot(
  x,
  type = c("trace", "AIC", "BIC", "logLik", "cAIC", "mAIC", "mBIC", "EBIC"),
  xlab = NULL,
  ylab = NULL,
  label = FALSE,
  main = NULL,
  ...
)
```

Arguments

`x`	a `mixturecure` object resulting from `curegmifs` or `cureem`, `cv_curegmifs` or `cv_cureem`.
`type`	a case-sensitive parameter indicating what to plot on the y-axis. The complete list of options are: • `"trace"` plots the coefficient path for the fitted object (default). • `"AIC"` plots the AIC against step of model fit. • `"mAIC"` plots the modified AIC against step of model fit. • `"cAIC"` plots the corrected AIC against step of model fit. • `"BIC"`, plots the BIC against step of model fit. • `"mBIC"` plots the modified BIC against step of model fit. • `"EBIC"` plots the extended BIC against step of model fit. • `"logLik"` plots the log-likelihood against step of model fit. This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.
`xlab`	a default x-axis label will be used which can be changed by specifying a user-defined x-axis label.
`ylab`	a default y-axis label will be used which can be changed by specifying a user-defined y-axis label.
`label`	logical. If TRUE the variable names will appear in a legend. Applicable only when `type = "trace"`. Be reminded that this works well only for small to moderate numbers of variables. For many predictors, the plot will be cluttered. The

	variables may be more easily identified using the `coef` function indicating the step of interest.
`main`	a default main title will be used which can be changed by specifying a user-defined main title. This option is not used for `cv_curegmifs` or `cv_cureem` fitted objects.
`...`	other arguments.

Value

this function has no returned value but is called for its side effects

See Also

`curegmifs`, `cureem`, `coef.mixturecure`, `summary.mixturecure`, `predict.mixturecure`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
plot(fit)
```

`predict.mixturecure` *Predicted probabilities for susceptibles, linear predictor for latency, and risk class for latency for mixture cure fit*

Description

This function returns a list that includes the predicted probabilities for susceptibles as well as the linear predictor for the latency distribution and a dichotomous risk for latency for a `curegmifs`, `cureem`, `cv_curegmifs` or `cv_cureem` fitted object.

Usage

```
## S3 method for class 'mixturecure'
predict(object, newdata, model_select = "AIC", ...)
```

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.
`newdata`	an optional data.frame that minimally includes the incidence and/or latency variables to use for predicting the response. If omitted, the training data are used.
`model_select`	either a case-sensitive parameter for models fit using `curegmifs` or `cureem` or any numeric step along the solution path can be selected. The default is `model_select = "AIC"` which calculates the predicted values using the coefficients from the model achieving the minimum AIC. The complete list of options are: • `"AIC"` for the minimum AIC (default). • `"mAIC"` for the minimum modified AIC. • `"cAIC"` for the minimum corrected AIC. • `"BIC"`, for the minimum BIC. • `"mBIC"` for the minimum modified BIC. • `"EBIC"` for the minimum extended BIC. • `"logLik"` for the step that maximizes the log-likelihood. • `n` where n is any numeric value from the solution path. This option has no effect for objects fit using `cv_curegmifs` or `cv_cureem`.
`...`	other arguments

Value

`p_uncured`	a vector of probabilities from the incidence portion of the fitted model representing the P(uncured).
`linear_latency`	a vector for the linear predictor from the latency portion of the model.
`latency_risk`	a dichotomous class representing low (below the median) versus high risk for the latency portion of the model.

See Also

`curegmifs`, `cureem`, `coef.mixturecure`, `summary.mixturecure`, `plot.mixturecure`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
predict_train <- predict(fit)
names(predict_train)
testing <- temp$testing
predict_test <- predict(fit, newdata = testing)
```

`print.mixturecure` *Print the contents of a mixture cure fitted object*

Description

This function prints the first several incidence and latency coefficients, the rate (when fitting an exponential or Weibull mixture cure model), and alpha (when fitting a Weibull mixture cure model). This function returns the fitted object invisible to the user.

Usage

```
## S3 method for class 'mixturecure'
print(x, max = 6, ...)
```

Arguments

`x`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_cureem`, or `cv_curegmifs`.
`max`	maximum number of rows in a matrix or elements in a vector to display
`...`	other arguments.

Value

prints coefficient estimates for the incidence portion of the model and if included, prints the coefficient estimates for the latency portion of the model. Also prints rate for exponential and Weibull models and scale (alpha) for the Weibull mixture cure model. Returns all objects fit using `cureem`, `curegmifs`, `cv_cureem`, or `cv_curegmifs`.

Note

The contents of a `mixturecure` fitted object differ depending upon whether the EM (`cureem`) or GMIFS (`curegmifs`) algorithm is used for model fitting or if cross-validation is used. Also, the output differs depending upon whether `x_latency` is specified in the model (i.e., variables are included in the latency portion of the model fit) or only `terms` on the right hand side of the equation are included (i.e., variables are included in the incidence portion of the model).

See Also

`curegmifs`, `cureem`, `coef.mixturecure`, `summary.mixturecure`, `plot.mixturecure`, `predict.mixturecure`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
```

```
  epsilon = 0.01, verbose = FALSE
)
print(fit)
```

`sufficient_fu_test` *Test for sufficient follow-up*

Description

Tests for sufficient follow-up using a Kaplan-Meier fitted object.

Usage

```
sufficient_fu_test(object)
```

Arguments

`object`	a `survfit` object.

Value

`p_value`	p-value from testing the null hypothesis that there was not sufficient follow-up against the alternative that there was sufficient follow-up
`n_n`	total number of events that occurred at time > pmax(0, 2*(last observed event time)-(last observed time)) and < the last observed event time
`N`	number of observations in the dataset

References

Maller, R. A. and Zhou, X. (1996) *Survival Analysis with Long-Term Survivors*. John Wiley & Sons.

See Also

`survfit`, `cure_estimate`, `nonzerocure_test`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
km_fit <- survfit(Surv(Time, Censor) ~ 1, data = training)
sufficient_fu_test(km_fit)
```

`summary.mixturecure` *Summarize a fitted mixture cure object*

Description

`summary` method for a mixturecure object fit using `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.

Usage

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

Arguments

`object`	a `mixturecure` object resulting from `curegmifs`, `cureem`, `cv_curegmifs`, or `cv_cureem`.
`...`	other arguments.

Value

prints the number of non-zero coefficients from the incidence and latency portions of the fitted mixture cure model when using the minimum AIC to select the final model. When fitting a model using `curegmifs` or `cureem` the summary function additionally prints results associated with the following model selection methods: the step and value that maximizes the log-likelihood; the step and value that minimizes the AIC, modified AIC (mAIC), corrected AIC (cAIC), BIC, modified BIC (mBIC), and extended BIC (EBIC). This information can be used to guide the user in the selection of a final model from the solution path.

See Also

`curegmifs`, `cureem`, `coef.mixturecure`, `plot.mixturecure`, `predict.mixturecure`

Examples

```
library(survival)
withr::local_seed(1234)
temp <- generate_cure_data(n = 100, j = 10, n_true = 10, a = 1.8)
training <- temp$training
fit <- curegmifs(Surv(Time, Censor) ~ .,
  data = training, x_latency = training,
  model = "weibull", thresh = 1e-4, maxit = 2000,
  epsilon = 0.01, verbose = FALSE
)
summary(fit)
```

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