

Package: PAmeasures (via r-universe)

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Type Package

Title Performance Assessment for Continuous and Survival Outcomes

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Description Evaluates prediction accuracy for continuous outcomes and the predictive performance of survival models in a unified framework. Implements the R-squared, L-squared, pseudo R-squared and related explained variation measures of Li and Wang (2019) <doi:10.1080/01621459.2018.1515079>, the concordance indices of Harrell et al. (1982) <doi:10.1001/jama.1982.03320430047030> and Uno et al. (2011) <doi:10.1002/sim.4154>, the time-dependent area under the receiver operating characteristic curve (AUC) of Heagerty, Lumley and Pepe (2000) <doi:10.1111/j.0006-341X.2000.00337.x>, and the Brier score of Graf et al. (1999) <doi:10.1002/(SICI)1097-0258(19990915/30)18:17/18%3C2529::AID-SIM274%3E3.0.CO;2-5>. Supports right-censored data; competing risks, following Wolbers et al. (2014) <doi:10.1093/biostatistics/kxt059>, Wu and Li (2018) <doi:10.1002/sim.7806> and Zhuang et al. (2025) <doi:10.48550/arXiv.2507.15040>; and two-phase sampling designs such as case-cohort and nested case-control studies, following Lee, Zeleniuch-Jacquotte and Liu (2021) <doi:10.1080/02664763.2020.1861225>. Revives and expands the original 'PAmeasures' package with a unified interface for survival prediction assessment.

Depends R (>= 4.1.0)

Imports Hmisc, survival, stats, utils, tdROC, survivalROC, yardstick, ggplot2, patchwork, dplyr, magrittr, purrr, tibble, tidyr, pec, pmcalibration (>= 0.2.0), prodlim

URL <https://github.com/Larryzza/PAmeasures>

BugReports <https://github.com/Larryzza/PAmeasures/issues>

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moore

*Moore's Law Data***Description**

Transistor counts for commercial microprocessors together with their year of introduction. Moore's Law observes that the number of transistors in a dense integrated circuit roughly doubles every two years.

Usage

moore

Format

A data frame with 48 rows and 3 variables:

year Year of introduction, from 1971 to 2012.

time Years elapsed since 1970.

count Number of transistors.

Source

Compiled from published microprocessor transistor counts.

pa_calibration

*Calibration of Predicted Event Probabilities at One Time Point***Description**

Assess calibration from externally supplied probabilities and observed outcomes, without requiring the original prediction model.

Usage

```
pa_calibration(p, event_time, status, t_star, nk = 5, case_weights = NULL)
```

```
pa_calibration_cr(
  p,
  event_time,
  status,
  t_star,
  event_type = 1,
  bandwidth = NULL
)
```

Arguments

<code>p</code>	Numeric vector of predicted event probabilities at <code>t_star</code> , in the same subject order as <code>event_time</code> and <code>status</code> . For ordinary survival supply $1 - S(t_star)$; for competing risks supply the CIF of <code>event_type</code> at <code>t_star</code> .
<code>event_time</code>	Numeric vector of finite, non-negative observed times.
<code>status</code>	Event indicators: 0 for censoring and 1 for an event in <code>pa_calibration()</code> ; non-negative integer event codes in <code>pa_calibration_cr()</code> (0 for censoring).
<code>t_star</code>	One positive finite evaluation time, not exceeding the maximum observed follow-up. Use the same time units as <code>event_time</code> .
<code>nk</code>	Number of restricted cubic spline knots (at least 3; default 5).
<code>case_weights</code>	Optional positive finite inverse-inclusion sampling weights, aligned with the subjects. For <code>pa_calibration()</code> only. Used in both the calibration model and the metric summaries.
<code>event_type</code>	Positive integer code for the cause of interest.
<code>bandwidth</code>	Nearest-neighborhood smoothing bandwidth passed to <code>prodlim::prodlim()</code> . NULL uses its automatic bandwidth.

Details

`pa_calibration()` uses `pmcalibration::pmcalibration()` with a restricted cubic spline and no confidence interval computation. The knot count is capped at the number of distinct fitting probabilities; with fewer than five distinct probabilities or fewer than six subjects it uses the backend's single-predictor Cox calibration instead. Constant predictions are compared with the marginal Kaplan–Meier risk.

The right-censored backend requires probabilities strictly inside (0, 1). Probabilities within `sqrt(.Machine$double.eps)` of 0 or 1 are clipped to that distance from the boundary for fitting, with a warning (except for the constant-prediction Kaplan–Meier branch). Metrics compare calibrated values with the original probabilities and retain every subject.

`pa_calibration_cr()` uses local Aalen–Johansen estimates from `prodlim`, conditional on the predicted CIF. All nonzero causes other than `event_type` are pooled as competing events, not censored. Constant or few-valued predictions use the backend's discrete risk groups. This is a different curve estimator from Fine–Gray spline calibration.

For unweighted evaluations, both functions use `pmcalibration::cal_metrics()` to summarize absolute differences between predicted and curve-implied probabilities at the original subjects. Only `ici` (the backend's `Eavg`), `e50`, and `e90` are returned. They are probability-scale errors, with 0 ideal. ICI averages over subjects, not over follow-up time or a uniform risk grid.

With `case_weights`, `pa_calibration()` fits a sampling-weighted Cox model directly, using the same complementary log-log transformation and restricted cubic spline knot-placement rule as the ordinary backend. Both coefficients and baseline cumulative hazard come from the weighted fit. Constant predictions use weighted Kaplan–Meier risk; with fewer than five distinct fitting probabilities or six subjects a single linear predictor is used. Knot locations are recorded in `knots`. ICI is the normalized sampling-weighted mean absolute error. `E50` and `E90` use weighted order-statistic interpolation, with weights normalized to the sample size (`Hmisc::wtd.quantile(..., normwt = TRUE)`). Multiplying every sampling weight by the same constant has no effect.

Supply subject inclusion weights for the actual case-cohort/NCC design, rather than weights already combined with censoring adjustments. Only sampled subjects with positive inclusion probabilities should be supplied. The calibration Cox model assumes conditionally independent censoring given its predictor and an appropriate proportional-hazards spline relationship. The competing-risks function remains an unweighted cohort evaluation. No optimism correction or design-based confidence interval is applied; preferably supply held-out or out-of-fold predictions.

Value

A list with `metrics`, a named numeric vector `c(ici, e50, e90)`; `curve`, a data frame with predicted and calibrated probabilities in input subject order; `t_star`; `event_type`; and `method`, the curve estimator. Weighted results also include the original `case_weights` and spline knots on the complementary log-log scale (empty for simpler fits).

References

Austin PC, Steyerberg EW (2019). The Integrated Calibration Index (ICI) and related metrics for quantifying the calibration of logistic regression models. *Statistics in Medicine* 38, 4051–4065. doi:10.1002/sim.8281. Austin PC, Harrell FE Jr, van Klaveren D (2020). Graphical calibration curves and the integrated calibration index (ICI) for survival models. *Statistics in Medicine* 39, 2714–2742. doi:10.1002/sim.8570. Breslow NE, Wellner JA (2007). Weighted likelihood for semi-parametric models and two-phase stratified samples, with application to Cox regression. *Scandinavian Journal of Statistics* 34, 86–102. doi:10.1111/j.14679469.2006.00523.x. Samuelsen SO (1997). A pseudolikelihood approach to analysis of nested case-control studies. *Biometrika* 84, 379–394. doi:10.1093/biomet/84.2.379.

Examples

```
set.seed(19)
x <- stats::rnorm(100)
event <- stats::rexp(100, rate = exp(x))
censor <- stats::rexp(100, rate = 0.3)
time <- pmin(event, censor)
status <- as.integer(event <= censor)
p <- 1 - exp(-exp(x) * 0.5)
pa_calibration(p, time, status, t_star = 0.5)$metrics

# Externally supplied CIFs; no fitted model object is needed.
competing <- stats::rexp(100, rate = 0.4)
cr_time <- pmin(event, competing, censor)
cr_status <- ifelse(censor < pmin(event, competing), 0L,
                  ifelse(event <= competing, 1L, 2L))
cif <- exp(x) / (exp(x) + 0.4) * (1 - exp(-(exp(x) + 0.4) * 0.5))
pa_calibration_cr(cif, cr_time, cr_status, t_star = 0.5)$metrics
```

pa_case_cohort_weights

Case-cohort sampling weights

Description

Computes static sampling weights for case-cohort evaluation. If strata is provided, non-case weights are calibrated within each stratum.

Usage

```
pa_case_cohort_weights(time, status, subcohort = NULL, strata = NULL)
```

Arguments

time	Numeric vector of follow-up times.
status	Numeric {0,1}, 1 = event, 0 = censored.
subcohort	Logical vector; TRUE if in subcohort.
strata	Optional factor/character for strata (stratified cc).

Details

Supply the full phase-one cohort, not only sampled subjects. Cases receive weight 1. Non-cases receive the number of full-cohort non-cases divided by the number of subcohort non-cases, within each stratum if supplied. These are calibrated inverse-inclusion weights, not the time-dependent risk-set construction of the Prentice Cox estimator. Passing these as ordinary Cox case weights does not implement that estimator.

Retain only the random subcohort and all cases for evaluation, and align weights with the evaluation subjects. At least one sampled non-case is needed in every stratum containing non-cases to obtain finite weights.

Value

Numeric vector of weights in full-cohort input order. Weights for non-sampled non-cases are also returned and must be excluded from the sampled analysis dataset.

Examples

```
# Compute weights using the full cohort, then retain the random subcohort
# and all cases for analysis. Sampled non-cases represent cohort non-cases.
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
                        beta = c(0.5, -0.5), seed = 2025)
set.seed(2025)
w <- pa_case_cohort_weights(time = d$time, status = d$status,
                            subcohort = stats::rbinom(nrow(d), 1, 0.4))
summary(w)
```

Description

This function constructs a standardized evaluation dataset (dat1) from the analysis dataset, predicted survival probabilities, and sampling weights, and computes key performance metrics in a tidy table format. Works for common two-phase sampling designs: - Case-cohort (unstratified or stratified) - Nested case-control (matched or unmatched)

Usage

```
pa_evaluate_two_phase(
  pred_results,
  t_star = NULL,
  tau = 1e+11,
  km_cens_fit,
  case_weights,
  metrics = c("pseudo_r2", "harrell_c", "uno_c", "brier_score", "td_auc", "ici"),
  calibration_args = list()
)
```

Arguments

pred_results	A prediction list, such as the output of <code>pa_predict_coxph()</code> , with times (or legacy time), status (0 = censored, 1 = event), pred (predicted mean survival times), and surv_prob (subjects by sorted observed times).
t_star	(Optional) Evaluation time for Brier score, time-dependent AUC, and calibration. If NULL, it defaults to the median observed follow-up time.
tau	(Optional) A numeric value specifying the truncation time for calculating explained variation metrics. If NULL, it defaults to the maximum observed survival time.
km_cens_fit	Kaplan-Meier estimate of the censoring distribution, a <code>survival::survfit</code> object. Not required when only calibration is requested.
case_weights	Numeric sampling weights in the same subject order as <code>pred_results</code> . Prediction helpers sort subjects by observed time; reorder weights using <code>order(new_data\$time)</code> as well.
metrics	Character vector of metrics. NULL uses the default metrics, including ICI; "all" additionally returns <code>r_square</code> , <code>l_square</code> , <code>e50</code> , and <code>e90</code> .
calibration_args	Optional named list of calibration settings, e.g. <code>list(nk = 3)</code> . Sampling weights are supplied by <code>case_weights</code> , not through this list.

Details

Calibration uses the event risk from the last prediction-grid column at or before `t_star`, with the exact requested outcome horizon. Both the Cox spline calibration model and absolute-error summaries use sampling weights; see `pa_calibration()`. Calibration-only inputs need times, status, and `surv_prob`, without fitted models or mean time predictions.

Value

A tibble with columns: - Metric: name of metric - Value: numeric value

Examples

```
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
  beta = c(0.5, -0.5), seed = 2025)
d <- d[, c("time", "status", "x1", "x2")]
fit <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
  data = d, x = TRUE, y = TRUE)
pred <- pa_predict_coxph(model = fit, covs = c("x1", "x2"), new_data = d)

# case-cohort sampling weights, and the censoring distribution they need
set.seed(2025)
in_subcohort <- stats::rbinom(nrow(d), 1, 0.4) == 1
w <- pa_case_cohort_weights(d$time, d$status, subcohort = in_subcohort)
keep <- which(in_subcohort | d$status == 1)
keep <- keep[order(d$time[keep])]
pred <- pa_predict_coxph(fit, c("x1", "x2"), new_data = d[keep, ])
w <- w[keep]
km <- survival::survfit(survival::Surv(d$time, 1 - d$status) ~ 1)

pa_evaluate_two_phase(pred_results = pred, km_cens_fit = km,
  case_weights = w)
```

pa_fit_and_eval

Performance Metrics for Survival Analysis Models

Description

This function computes a comprehensive set of performance metrics for survival analysis models. It provides metrics such as `R_square`, `L_square`, `Pseudo_R`, Harrell's C, Uno's C, Brier Score, and Time-dependent AUC. Users can specify particular metrics and model types, enabling tailored performance evaluation for various survival models.

Usage

```
pa_fit_and_eval(
  train_data,
  covariates,
  models = "coxph",
```

```

    metrics = NULL,
    predicted_data = NULL,
    t_star = NULL,
    tau = NULL,
    calibration_args = list()
  )

```

Arguments

<code>train_data</code>	A data frame containing the survival data.
<code>covariates</code>	A character vector of covariate names to include in the model.
<code>models</code>	Character vector of model types: "coxph", "exp", "lognormal", or "weibull". The default fits only "coxph"; use "all" to fit all four.
<code>metrics</code>	A character string or vector specifying the metrics to compute. The default NULL includes ICI and the existing performance metrics; use "all" to additionally include E50 and E90. Options include: <ul style="list-style-type: none"> • "r_square": R-squared metric. • "l_square": L-squared metric. • "pseudo_r2": Pseudo-R-squared metric. • "harrell_c": Harrell's Concordance Index. • "uno_c": Uno's Concordance Index. • "brier_score": Brier Score. • "td_auc": Time-dependent AUC. • "ici", "e50", "e90": Absolute calibration error summaries.
<code>predicted_data</code>	(Optional) A data frame containing validation data. If NULL, the function uses <code>train_data</code> for model evaluation.
<code>t_star</code>	(Optional) Positive evaluation time for Brier score, AUC, and calibration; the median evaluation follow-up if omitted.
<code>tau</code>	(Optional) A time point for truncating the survival time. If provided, the function evaluates predictions up to this time point.
<code>calibration_args</code>	Optional named list passed to <code>pa_calibration()</code> . For sampling-adjusted calibration use <code>pa_sample_design()</code> or <code>pa_calibration()</code> directly; weights cannot be supplied through this list.

Details

Calibration evaluates model-predicted event probabilities exactly at `t_star` (the median evaluation follow-up if omitted).

Rows with missing values in time, status, or a requested covariate are excluded from training and evaluation with a warning. All requested metrics use the same complete evaluation rows. The warning can be silenced with `suppressWarnings()`.

Value

A data frame containing the selected model's performance metrics.

Examples

```
data(pbc, package = "survival")
pbc <- pbc[!is.na(pbc$trt), ]
pbc$log_albumin <- log(pbc$albumin)
pbc$log_bili    <- log(pbc$bili)
pbc$log_protime <- log(pbc$protime)
pbc$status <- ifelse(pbc$status == 2, 1, 0)
covariates <- c("age", "log_albumin", "log_bili", "log_protime", "edema")
dat <- pbc[, c("time", "status", covariates)]
dat <- dat[stats::complete.cases(dat), ]

# Default Cox model and metrics, including ICI
results <- pa_fit_and_eval(train_data = dat, covariates = covariates)
results

# A specific subset of models and metrics
results2 <- pa_fit_and_eval(
  train_data = dat,
  covariates = covariates,
  models     = c("lognormal", "weibull"),
  metrics    = c("r_square", "l_square", "brier_score")
)
results2
```

pa_metric_names	<i>Canonical metric names</i>
-----------------	-------------------------------

Description

The metric identifiers accepted by the metrics argument of the evaluation functions, and emitted in the Metric column of their results.

Usage

```
pa_metric_names()
```

Value

A character vector of canonical metric names.

Examples

```
pa_metric_names()
```

pa_nested_case_control_weights

Nested case-control (NCC) sampling weights

Description

Computes inverse-selection weights for NCC designs. If strata is provided, matched NCC weights are computed.

Usage

```
pa_nested_case_control_weights(time, status, strata = NULL, m = NULL)
```

Arguments

time	Numeric vector of follow-up times.
status	Numeric {0,1}, 1 = event, 0 = censored.
strata	Optional factor/character defining matching strata in the full phase-one cohort, not IDs of sampled case-control sets.
m	Positive integer, number of controls per case (required).

Details

Supply full-cohort follow-up and event information, not only sampled rows. Controls are sampled without replacement within each case's risk set and independently across cases. At tied failure times, all simultaneous cases are excluded from the eligible controls. Events precede censoring at ties. Matching restricts both cases and controls to the same stratum. If fewer than *m* controls are eligible, all eligible controls are sampled. A non-case with no sampling opportunity has infinite inverse probability; such a subject cannot belong to the sampled analysis dataset. Subset the returned weights to the subjects actually sampled and align their order with the prediction objects before evaluation.

Value

Numeric vector of weights, in full-cohort input order.

Examples

```
# Unmatched nested case-control design with m = 2 controls per case.
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
                       beta = c(0.5, -0.5), seed = 2025)

set.seed(2025)
w <- pa_nested_case_control_weights(time = d$time, status = d$status, m = 2)
summary(w)
```

pa_nlm

*Prediction Accuracy for Continuous Outcomes***Description**

Evaluate observed and predicted continuous responses using the R-squared and L-squared measures of Li and Wang (2019). Predictions may come from a linear or nonlinear model; the original fitted model is not required.

Usage

```
pa_nlm(y, y.predict)
```

```
pam.nlm(y, y.predict)
```

Arguments

<code>y</code>	A finite numeric vector of observed continuous responses.
<code>y.predict</code>	A finite numeric vector of predictions, paired with <code>y</code> in the same order and of the same length. At least two pairs are required. Missing and infinite values are rejected.

Details

An ordinary least-squares regression of `y` on `y.predict`, with an intercept, gives the corrected predictions $\hat{y}_c$. The measures are

$$R^2 = \frac{\sum_i (\hat{y}_{c,i} - \bar{y})^2}{\sum_i (y_i - \bar{y})^2}$$

and

$$L^2 = \frac{\sum_i (y_i - \hat{y}_{c,i})^2}{\sum_i (y_i - y.predict_i)^2}.$$

R-squared describes explained response variation after affine correction; L-squared compares the corrected and original prediction errors. These are the original PAmesures definitions, not $1 - L^2$.

Constant predictions use the intercept-only correction `mean(y)`. A constant observed response makes R-squared undefined. Predictions exactly equal to the observations make L-squared undefined because its denominator is zero. Undefined measures are `NA_real_`, with a warning; any other defined measure is retained. Inputs whose squared deviations overflow or whose nonzero deviations underflow to zero are rejected with an error and should be rescaled to common units.

`pam.nlm()` is the retained PAmesures 0.1.0 compatibility entry point. It uses the same calculation, then rounds and formats each result to four decimal places as in the historical package. Its ordinary, defined results retain the historical field names and character format; input validation and undefined-result handling follow `pa_nlm()`. No deprecation warning is issued for this supported compatibility entry.

Value

A list with R.squared and L.squared. pa_nlm() returns unrounded numeric scalars. pam.nlm() returns the historical formatted character scalars.

References

Li, G. and Wang, X. (2019). Prediction Accuracy Measures for a Nonlinear Model and for Right-Censored Time-to-Event Data. *Journal of the American Statistical Association*, 114, 1815–1825. doi:10.1080/01621459.2018.1515079.

Examples

```
y <- c(1, 2, 4, 5)
prediction <- c(1, 2, 3, 4)
pa_nlm(y, prediction)
pam.nlm(y, prediction)

data(moore)
fit <- stats::lm(log2(count) ~ time, data = moore)
pa_nlm(moore$count, 2^stats::predict(fit))
```

pa_plot_pred

Descriptive Plot of Predicted and Observed Survival Times

Description

Plots a model-derived survival-time summary and observed follow-up times against the model's linear predictor. Event and censoring status are shown with different point shapes. This is a descriptive display, not a probability-calibration curve.

Usage

```
pa_plot_pred(
  data,
  title = NULL,
  xlab = "Risk Score",
  ylab = "Days",
  label_name = c("censored", "event"),
  levels = NULL,
  shape_style = NULL,
  legend_name = "status",
  invert_linear = TRUE,
  sample_index = NULL,
  restrict_time = NULL
)
```



```
d <- d[, c("time", "status", "x1", "x2")]
fit <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
                      data = d, x = TRUE, y = TRUE)
pred <- pa_predict_coxph(model = fit, covs = c("x1", "x2"), new_data = d)

# observed time against predicted risk score, shaped by event status
pa_plot_pred(pred, title = "Cox model", invert_linear = FALSE)
```

pa_plot_summary	<i>Arrange multiple prediction plots with letter tags</i>
-----------------	---

Description

Builds one panel per data frame using `plot_fun` (default: `pa_plot_pred`) and arranges them with letter tags. The function collects a single shared legend and applies global styling (titles, axis labels, legend options) across panels.

Usage

```
pa_plot_summary(
  data_list,
  titles = NULL,
  plot_fun = pa_plot_pred,
  ncol = 2,
  tag_levels = "a",
  invert_linear = TRUE,
  xlab = "Risk Score",
  ylab = "Days",
  label_name = c("Censored", "Event"),
  levels = NULL,
  shape_style = NULL,
  sample_size = NULL,
  restrict_time = NULL,
  legend_name = "Status",
  seed = NULL
)
```

Arguments

<code>data_list</code>	List of data frames, each containing columns: <code>linear.pred</code> , <code>pred</code> , <code>times</code> , and <code>status</code> .
<code>titles</code>	Optional character vector, same length as <code>data_list</code> ; if <code>NULL</code> , defaults to "Panel 1", "Panel 2", etc.
<code>plot_fun</code>	Function that creates a single panel (default = <code>pa_plot_pred</code>).
<code>ncol</code>	Number of columns in the panel layout (default = 2).
<code>tag_levels</code>	Letter style for tags: "a", "A", "1", "i", or "I".

invert_linear	Logical or logical vector; if length 1, recycled to all panels. Use FALSE for Cox log-hazard scores and TRUE for AFT location scores when larger x values should indicate higher risk.
xlab, ylab	Axis labels applied to all panels (defaults are "Risk Score" and "Days").
label_name	Character vector of labels for status legend (e.g., c("censored", "event")).
levels	Numeric vector for status level ordering.
shape_style	Numeric vector of plotting symbols for censoring/event status.
sample_size	Optional integer. If supplied, that many subjects are sampled at random for each panel using seed; otherwise every subject is plotted.
restrict_time	Optional numeric. If supplied, observed times above this value are capped at it and a dashed horizontal reference line is drawn, which keeps a few long follow-up times from compressing the plot.
legend_name	Character string for the legend title (default = "status").
seed	Optional integer. If supplied, the random number generator is seeded with it before sample_size subjects are drawn, so the sampled points are reproducible. The default NULL uses the current random number stream without resetting it; random draws advance its state.

Value

A patchwork ggplot object combining all panels.

Examples

```
# One panel per model, with larger x values indicating higher risk.
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
  beta = c(0.5, -0.5), seed = 2025)
d <- d[, c("time", "status", "x1", "x2")]
cox <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
  data = d, x = TRUE, y = TRUE)
wei <- survival::survreg(survival::Surv(time, status) ~ x1 + x2,
  data = d, dist = "weibull", x = TRUE, y = TRUE)
p_cox <- pa_predict_coxph(model = cox, covs = c("x1", "x2"), new_data = d)
p_wei <- pa_predict_survreg(model = wei, covs = c("x1", "x2"), new_data = d)

pa_plot_summary(
  list(p_cox, p_wei),
  titles = c("Cox PH", "Weibull AFT"),
  invert_linear = c(FALSE, TRUE),
  ncol = 2,
  legend_name = "Status"
)
```

pa_predict_cif

*Predict Subject-Specific Cumulative Incidence Function (CIF)***Description**

This function calculates cumulative incidence function (CIF) predictions for individual subjects across multiple types of competing risks models. It supports cause-specific Cox models, accelerated failure time (AFT) models (Weibull, Log-normal, Log-logistic), Fine-Gray models, and competing risks random survival forests.

Usage

```
pa_predict_cif(
  model1 = NULL,
  model2 = NULL,
  fg_model = NULL,
  cr_model = NULL,
  tau = NULL,
  newdata,
  event.type = 1,
  covs
)
```

Arguments

model1	The fitted model for the target cause (either a coxph or survreg object).
model2	The fitted model for competing causes (either a coxph or survreg object).
fg_model	Optional Fine-Gray model object (crr from cmprsk package).
cr_model	Optional competing risks random forest model object.
tau	The maximum time horizon for prediction.
newdata	A data frame containing observed event times (time), event indicators (status).
event.type	The event type of interest for which CIF is to be predicted.
covs	A character vector specifying the names of the covariates used in the model.

Details

For cause-specific AFT models, CIFs are evaluated by adaptive quadrature on the target cause's probability scale. This handles Weibull densities that are unbounded at time zero, including fixed scales above 1. The time summary integrates one minus the CIF on the returned time grid.

Value

A list with the following components:

times Numeric vector of observed times from newdata\$time.

status Integer vector of event indicators from `newdata$status`.

cif_pred A matrix where the first column is time, and subsequent columns are predicted CIFs for each subject.

pred Numeric vector of the integral of one minus the CIF from 0 to τ .

linear.pred Numeric vector of subject-specific linear predictors.

Examples

```
data(pbc, package = "survival")
pbc <- pbc[!is.na(pbc$strt), ]
pbc$log_albumin <- log(pbc$albumin)
pbc$log_bili <- log(pbc$bili)
pbc$log_protime <- log(pbc$protime)
covariates <- c("age", "log_albumin", "log_bili", "log_protime", "edema")
dat <- pbc[, c("time", "status", covariates)]
dat <- dat[stats::complete.cases(dat), ]

# Cause-specific Cox models: event of interest (1) and competing event (2)
m1 <- survival::coxph(survival::Surv(time, status == 1) ~ age + log_albumin +
  log_bili + log_protime + edema,
  data = dat, x = TRUE, y = TRUE)
m2 <- survival::coxph(survival::Surv(time, status == 2) ~ age + log_albumin +
  log_bili + log_protime + edema,
  data = dat, x = TRUE, y = TRUE)

pred <- pa_predict_cif(model1 = m1, model2 = m2, newdata = dat,
  covs = covariates, event.type = 1,
  tau = max(dat$time))

str(pred$pred)

# Fine-Gray backend, available when cmprsk is installed
if (requireNamespace("cmprsk", quietly = TRUE)) {
  fg <- cmprsk::crr(ftime = dat$time, fstatus = dat$status,
    cov1 = as.matrix(dat[, covariates]),
    failcode = 1, cencode = 0)
  pred_fg <- pa_predict_cif(fg_model = fg, newdata = dat,
    covs = covariates, event.type = 1,
    tau = max(dat$time))

  str(pred_fg$pred)
}
```

Description

Returns subject-specific survival probabilities and a time summary from a fitted model when `predict = TRUE`. With `predict = FALSE`, returns R-squared, L-squared and pseudo-R-squared instead. R-squared measures explained variation after affine correction. L-squared is the corrected prediction

error divided by the original prediction error; it is not the fraction of error removed by correction. Pseudo-R-squared is their product.

Usage

```
pa_predict_coxph(model, covs, tau = 1e+11, new_data = NULL, predict = TRUE)
```

Arguments

model	An object of class <code>coxph</code> representing a fitted Cox proportional hazards regression model. The model must be fitted with <code>x = TRUE</code> and <code>y = TRUE</code> to include the design matrix and response vector.
covs	A character vector specifying the names of the covariates used in the model.
tau	(Optional) A restriction time for computing restricted survival time. If provided, the function evaluates predictions up to this time point. Default is <code>10e10</code> .
new_data	Optional evaluation data. Its outcomes are returned for metric computation and define the evaluation time grid; they never estimate the baseline hazard. All prediction parameters come from the training fit.
predict	A logical value indicating whether to return individual predictions. Default is <code>TRUE</code> .

Details

Probabilities use the fitted training baseline, including its case weights, strata, tie method and offsets. Offset centering follows the weighted training mean used by `survfit()`. With `predict = FALSE`, R-squared metrics return missing values with a warning if censoring weights cannot be normalized, for example when the training censoring survival is zero at an observed evaluation event.

Value

With `predict = TRUE`, a single prediction list containing:

- `model` – the fitted survival model object.
- `times` – numeric vector of observed follow-up times for the evaluation dataset.
- `status` – event indicators (1 = event, 0 = censored).
- `surv_prob` – an $n \times K$ matrix of predicted subject-specific survival probabilities on a common time grid.
- `pred` – numerical integral of the training-model survival curve on the evaluation grid, up to its maximum time or `tau`, whichever is smaller. This preserves the legacy integration rule; it is not an identified unrestricted mean. Fix the grid and `tau` for comparisons.
- `linear.pred` – the model's linear predictor, aligned with the returned subjects.
- `covs` – character vector of covariate names used for prediction.
- `new_data` – dataset on which the predictions were computed.

Subjects are sorted by observed follow-up time. The columns of `surv_prob` correspond to the sorted follow-up times, including ties. With `predict = FALSE`, the result is a list of formatted character scalars named `R.squared`, `L.squared`, and `Psuedo.R`. The last spelling is retained for compatibility.

References

Li, G., & Wang, X. (2016). Prediction Accuracy Measures for a Nonlinear Model and for Right-Censored Time-to-Event Data. arXiv preprint arXiv:1611.03063. Available at <https://arxiv.org/abs/1611.03063>.

Examples

```
data(pbc, package = "survival")
pbc <- pbc[!is.na(pbc$trt), ]
pbc$log_albumin <- log(pbc$albumin)
pbc$log_bili <- log(pbc$bili)
pbc$log_protime <- log(pbc$protime)
pbc$status <- ifelse(pbc$status == 2, 1, 0)
covs <- c("age", "log_albumin", "log_bili", "log_protime", "edema")
df <- pbc[, c("time", "status", covs)]
df <- df[stats::complete.cases(df), ]

set.seed(2025)
idx <- sample(seq_len(nrow(df)), round(2 / 3 * nrow(df)))
train_data <- df[idx, ]
test_data <- df[-idx, ]

m.cox <- survival::coxph(survival::Surv(time, status) ~ .,
                        data = train_data, x = TRUE, y = TRUE)
cox_pred <- pa_predict_coxph(model = m.cox, covs = covs,
                           new_data = test_data)

str(cox_pred$pred)
```

pa_predict_survreg	<i>Prediction Accuracy Measures for Parametric Survival Regression Models</i>
--------------------	---

Description

Returns subject-specific survival probabilities and a time summary from a fitted model when predict = TRUE. With predict = FALSE, returns R-squared, L-squared and pseudo-R-squared instead. R-squared measures explained variation after affine correction. L-squared is the corrected prediction error divided by the original prediction error; it is not the fraction of error removed by correction. Pseudo-R-squared is their product.

Usage

```
pa_predict_survreg(model, covs, tau = 1e+11, new_data = NULL, predict = TRUE)
```

Arguments

model	An object of class survreg fitted with distribution "exponential", "weibull", or "lognormal". Fit with <code>x = TRUE</code> and <code>y = TRUE</code> to store the design and response.
covs	A vector of covariate names to be used for prediction when <code>new_data</code> is provided.
tau	(Optional) A restriction time for computing restricted survival time. If provided, the function evaluates predictions up to this time point. Default is 10e10.
new_data	(Optional) A new dataset for evaluating the model. If <code>NULL</code> , the function uses the training data stored in model.
predict	(Optional) A logical value indicating whether to return individual predictions. Default is <code>TRUE</code> .

Details

With `predict = FALSE`, R-squared metrics return missing values with a warning if censoring weights cannot be normalized, for example when the training censoring survival is zero at an observed evaluation event. Individual predictions remain available with `predict = TRUE`.

Value

With `predict = TRUE`, a single prediction list containing:

- `model` – the fitted survival model object.
- `times` – numeric vector of observed follow-up times for the evaluation dataset.
- `status` – event indicators (1 = event, 0 = censored).
- `surv_prob` – an $n \times K$ matrix of predicted subject-specific survival probabilities on a common time grid.
- `pred` – model-based restricted mean survival time $E[\min(T, \tau) \mid X]$, calculated analytically up to `tau`; it is not the median survival time.
- `linear.pred` – the model's linear predictor, aligned with the returned subjects.
- `covs` – character vector of covariate names used for prediction.
- `new_data` – dataset on which the predictions were computed.

Subjects are sorted by observed follow-up time. The columns of `surv_prob` correspond to the sorted follow-up times, including ties. With `predict = FALSE`, the result is a list of formatted character scalars named `R.squared`, `L.squared`, and `Psuedo.R`. The last spelling is retained for compatibility.

Examples

```
data(pbc, package = "survival")
pbc <- pbc[!is.na(pbc$trt), ]
pbc$log_albumin <- log(pbc$albumin)
pbc$log_bili <- log(pbc$bili)
pbc$log_protime <- log(pbc$protime)
```

```

pbc$status <- ifelse(pbc$status == 2, 1, 0)
covs <- c("age", "log_albumin", "log_bili", "log_protime", "edema")
df <- pbc[, c("time", "status", covs)]
df <- df[stats::complete.cases(df), ]

set.seed(2025)
idx <- sample(seq_len(nrow(df)), round(2 / 3 * nrow(df)))
train_data <- df[idx, ]
test_data <- df[-idx, ]

m.wei <- survival::survreg(survival::Surv(time, status) ~ .,
                          data = train_data, dist = "weibull",
                          x = TRUE, y = TRUE)
wei_pred <- pa_predict_survreg(model = m.wei, covs = covs,
                              new_data = test_data)
str(wei_pred$pred)

```

pa_sample_design	<i>Summarize multiple two-phase (case-cohort) models into a wide table</i>
------------------	--

Description

For a list of models (each providing times, status, surv_prob, pred), calls `pa_evaluate_two_phase()` for each and returns a wide comparison table with metrics as rows and model names as columns.

Usage

```

pa_sample_design(
  models,
  case_weights,
  km_cens,
  metrics = c("pseudo_r2", "harrell_c", "uno_c", "brier_score", "td_auc", "ici"),
  t_star = NULL,
  tau = NULL,
  digits = 2,
  calibration_args = list()
)

```

Arguments

<code>models</code>	A named list of prediction objects, such as those returned by <code>pa_predict_coxph()</code> . Each entry contains <code>times</code> (observed follow-up), <code>status</code> (0 = censored, 1 = event), and <code>surv_prob</code> (subjects by sorted follow-up times). Include <code>pred</code> , the predicted survival-time summary, for the standard metric workflow. Calibration-only evaluation does not require <code>pred</code> or a fitted model.
<code>case_weights</code>	Numeric vector of inverse-inclusion sampling weights, aligned with each model's evaluation subjects.

km_cens	A Kaplan-Meier fit for censoring (based on training data) used to compute IPCW. Not required for calibration-only evaluation.
metrics	Character vector of metrics to compute (default shown below).
t_star	Optional evaluation time for Brier score, AUC and calibration. Defaults to the median observed follow-up time.
tau	Optional truncation time for explained-variation metrics. If NULL, the single-model evaluator uses maximum observed follow-up.
digits	Integer number of decimal places to round values (default 2).
calibration_args	Optional named list passed to <code>pa_calibration()</code> , e.g. <code>list(nk = 3)</code> . Weights come from <code>case_weights</code> .

Value

A data.frame: rows = metrics, columns = model names.

Examples

```
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
  beta = c(0.5, -0.5), seed = 2025)
d <- d[, c("time", "status", "x1", "x2")]
fit <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
  data = d, x = TRUE, y = TRUE)
pred <- pa_predict_coxph(model = fit, covs = c("x1", "x2"), new_data = d)

# case-cohort sampling weights, and the censoring distribution they need
set.seed(2025)
in_subcohort <- stats::rbinom(nrow(d), 1, 0.4) == 1
w <- pa_case_cohort_weights(d$time, d$status, subcohort = in_subcohort)
keep <- which(in_subcohort | d$status == 1)
keep <- keep[order(d$time[keep])]
pred <- pa_predict_coxph(fit, c("x1", "x2"), new_data = d[keep, ])
w <- w[keep]
km <- survival::survfit(survival::Surv(d$time, 1 - d$status) ~ 1)

# the wide summary of the same two-phase evaluation
pa_sample_design(models = list(cc = pred), case_weights = w, km_cens = km)
```

pa_sim_cox_weibull	<i>Simulate Cox survival data with optional interaction terms and targeted censoring</i>
--------------------	--

Description

Generates survival data under a Cox proportional hazards model with a Weibull baseline cumulative hazard $H_0(t) = (0.5t)^v$. Event times are sampled via inverse transform:

$$Y = H_0^{-1}(-\log U \cdot \exp(-\beta^\top Z)), \quad U \sim \text{Unif}(0, 1),$$

where $H_0^{-1}(s) = 2s^{1/v}$.

Covariates are simulated from a standard normal distribution. With both `interact = FALSE` and `nonlinear = FALSE`, the design matrix Z contains only main effects. When `interact = TRUE`, the function automatically detects whether `length(beta)` equals 3 or 7:

- If `length(beta) = 3`, two covariates (x_1, x_2) and their two-way interaction (x_1x_2) are included.
- If `length(beta) = 7`, three covariates (x_1, x_2, x_3) , their three two-way interactions (x_1x_2, x_1x_3, x_2x_3) , and their three-way interaction $(x_1x_2x_3)$ are included.

Right-censoring times are generated as $C_i = \exp(\mu + r_i)$ with $r_i \sim \mathcal{N}(0, \text{sd}^2)$. By default, `sd` is set to `sd(log Y)`, and `mu` is calibrated so that the expected censoring proportion equals `pi_c`. If `mu` and/or `sd` are supplied, calibration is skipped accordingly:

- **mu and sd provided:** use them directly (no calibration).
- **sd only:** solve for `mu` using the given `sd`.
- **mu only:** use the given `mu` and estimate `sd(log Y)`.
- **neither:** estimate both `sd(log Y)` and `mu`.

Usage

```
pa_sim_cox_weibull(
  n,
  pi_c,
  v,
  beta,
  mu = NULL,
  sd = NULL,
  seed = NULL,
  interact = FALSE,
  nonlinear = FALSE
)
```

Arguments

<code>n</code>	Integer. Sample size.
<code>pi_c</code>	Numeric in $[0, 1)$. Zero returns uncensored data. For a positive value, this is the target censoring proportion when <code>mu</code> is omitted; supplying <code>mu</code> overrides target calibration.
<code>v</code>	Positive numeric. Weibull shape in $H_0(t) = (0.5t)^v$.
<code>beta</code>	Numeric vector of regression coefficients. If <code>interact = FALSE</code> , its length defines the number of main effects. If <code>interact = TRUE</code> , its length must be either 3 (two main + one interaction) or 7 (three main + four interactions).
<code>mu</code>	Optional numeric. Mean of $\log C$. If supplied with <code>sd</code> , no calibration is performed.
<code>sd</code>	Optional positive numeric. Standard deviation used in $\log C = \mu + r_i$. If <code>NULL</code> , defaults to <code>sd(log Y)</code> .

seed	Optional integer seed for reproducibility.
interact	Logical (default FALSE). If TRUE, include pre-defined interaction terms as described above.
nonlinear	Logical. If TRUE, the latent event-time model uses x_1, x_2, x_1x_2, x_1^2 , in that coefficient order, and beta must have length 4. Takes precedence over interact.

Value

A data.frame with columns:

- time: observed time $T_i = \min(Y_i, C_i)$.
- status: event indicator $1\{Y_i \leq C_i\}$.
- x1, x2, ...: simulated covariates.
- y_true: latent event time Y_i , only when $\text{pi_c} > 0$.
- cens_time: censoring time C_i , only when $\text{pi_c} > 0$.

Interaction and quadratic terms are used internally to generate event times, but only the original covariates are returned as columns. With $\text{pi_c} = 0$, time is the latent event time and all status values are 1; no model or censoring attributes are attached. With $\text{pi_c} > 0$, additional attributes are:

- mu: censoring shift used.
- sd_log: standard deviation used in $\log C$.
- sd_logY: $\text{sd}(\log Y)$ from the generated events.
- pi_c_target: requested censoring rate.
- pi_c_observed: realized censoring rate.
- v, beta: model parameters.

Examples

```
# (1) Main effects only
set.seed(1)
d0 <- pa_sim_cox_weibull(500, 0.3, v = 1.2, beta = c(0.4, -0.2))

# (2) Two covariates + their interaction
d1 <- pa_sim_cox_weibull(500, 0.3, v = 1.2,
  beta = c(0.5, -0.3, 0.2),
  interact = TRUE)

# (3) Three covariates + all two-way and three-way interactions
b <- c(0.4, -0.2, 0.1, 0.15, -0.05, 0.08, 0.02)
d2 <- pa_sim_cox_weibull(500, 0.3, v = 1.0, beta = b, interact = TRUE)
```

pa_simulate_fine_gray *Simulate data from a two-cause Fine-Gray model*

Description

This function generates a simulated dataset for a competing risks model with two causes, following a Fine-Gray-like model specification.

Usage

```
pa_simulate_fine_gray(
  n,
  v,
  beta1,
  beta2,
  lambda1 = 1,
  X = NULL,
  mu = 0,
  p = 0.7,
  c_scale = 1,
  censor = 0,
  sd.time = NULL,
  mu.c.time = NULL,
  independent_c = TRUE,
  report.mu_and_sd = FALSE,
  seed = NULL
)
```

Arguments

n	An integer specifying the number of subjects to simulate.
v	A numeric value for the Weibull shape parameter.
beta1	A numeric vector of coefficients for cause 1.
beta2	A numeric vector of coefficients for cause 2.
lambda1	Positive coefficient in the baseline term $1 - \exp(-\lambda_1 t^v)$ for cause 1.
X	A matrix of covariates, if not provided, covariates are generated from a standard normal distribution.
mu	A numeric value for the mean of the generated covariates.
p	Baseline probability of a cause 1 event when the cause-1 linear predictor is zero. Conditional on covariates, this probability is $1 - (1 - p)^{\exp(X\beta_1)}$; it is not generally the marginal probability across the generated covariates.
c_scale	Positive multiplier for the default standard deviation of log censoring times.

sensor	Target censoring proportion. Zero produces uncensored data. For a positive value, the log-censoring shift is calibrated to this target unless <code>mu.c.time</code> is supplied. Because calibration uses the generated sample, the attained fraction can differ by approximately one subject.
sd.time	Standard deviation of log censoring times. If <code>NULL</code> , it is calculated from the log cause-1 event times and <code>c_scale</code> .
mu.c.time	Optional additive shift of log censoring times. When <code>sensor > 0</code> , a supplied shift overrides target calibration; otherwise it is found using <code>uniroot()</code> . Ignored when <code>sensor = 0</code> .
independent_c	A logical value. If <code>TRUE</code> , censoring times are independent of covariates.
report.mu_and_sd	A logical value. If <code>TRUE</code> , a summary of censoring parameters and event time quantiles is returned instead of the full dataset.
seed	Optional integer. If supplied, the random number generator is seeded with it for reproducibility. The default <code>NULL</code> uses the current random number stream without resetting it; random draws advance its state.

Value

A data frame containing the simulated data with columns `obs.times`, `obs.event`, and the covariates `X`. Alternatively, if `report.mu_and_sd` is `TRUE`, it returns a data frame with censoring parameters and event time quantiles.

Examples

```
# Simulate a dataset with 100 subjects
sim_data <- pa_simulate_fine_gray(
  n = 100, v = 1, beta1 = c(0.5, -0.2), beta2 = c(-0.3, 0.4)
)
head(sim_data)

# Simulate with a specific censoring proportion and report parameters
params_summary <- pa_simulate_fine_gray(
  n = 1000, v = 1, beta1 = c(0.5, -0.2), beta2 = c(-0.3, 0.4),
  sensor = 0.2, report.mu_and_sd = TRUE
)
params_summary
```

pa_summarize

Summarize multiple survival models into a wide comparison table

Description

Calls `pa_survival_eval()` for each model in a named list and outputs a wide table: each row corresponds to one metric and each column corresponds to a model.

Usage

```
pa_summarize(
  models,
  metrics = NULL,
  t_star = NULL,
  tau = 1e+11,
  digits = 2,
  calibration_args = list()
)
```

Arguments

models	<p>A named list in which each element corresponds to a fitted model. Each model entry must itself be a list containing:</p> <ul style="list-style-type: none"> • times – numeric vector of observed follow-up times. • surv_prob – an $n \times K$ matrix (or data frame) of subject-specific predicted survival probabilities on a common time grid. • status – event indicator (1 = event, 0 = censored). • pred – (optional) predicted mean survival time (restricted or unrestricted). • new_data – (optional) dataset used for prediction. • covs – character vector of covariate names used for prediction. • model – (optional) the underlying fitted survival model object. <p>Note: If pred is provided, it will be used to calculate R2 and concordance measure.</p>
metrics	Optional character vector of metrics to compute (passed through).
t_star	Optional evaluation time for point metrics and calibration. Defaults to the median observed follow-up.
tau	Optional numeric scalar, specify the max time horizon for R2 measure and concordance measure (default = 10e10).
digits	Number of decimal places to round the metric values (default = 2).
calibration_args	Optional named list passed to pa_calibration() .

Value

A data frame with:

- Each row = metric
- Each column = model name
- Cell values = corresponding rounded metric values

Examples

```
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
  beta = c(0.5, -0.5), seed = 2025)
d <- d[, c("time", "status", "x1", "x2")]
fit <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
  data = d, x = TRUE, y = TRUE)
pred <- pa_predict_coxph(model = fit, covs = c("x1", "x2"), new_data = d)

# one column per model, metrics down the rows
pa_summarize(list(cox = pred))
```

pa_summarize_cr	<i>Summarize multiple competing-risks models into a wide comparison table</i>
-----------------	---

Description

Calls `pa_survival_eval_cr()` for each model in a named list and returns a wide table where rows are metrics and columns are model names.

Usage

```
pa_summarize_cr(
  models,
  metrics = NULL,
  t_star = NULL,
  tau = NULL,
  event_type = 1,
  digits = 2,
  censoring = c("cox", "km"),
  censor_covariates = NULL,
  calibration_args = list()
)
```

Arguments

models	<p>A named list of prediction objects, such as those returned by <code>pa_predict_cif()</code>. Each entry must contain:</p> <ul style="list-style-type: none"> • <code>cif_pred</code> – numeric matrix with the prediction time grid in the first column and one target-event CIF column per subject. • <code>times</code> – observed follow-up times, in subject-column order. • <code>status</code> – event codes (0 = censoring; <code>event_type</code> = target event; other positive codes = competing events).
--------	--

For external predictions, construct this format as shown in the example. The separate `pred_cif`, `time.cif`, and `event_time` arguments belong to `pa_survival_eval_cr()`, not to this wrapper.

metrics	Character vector of metrics to compute (passed through to <code>pa_survival_eval_cr()</code>); use "all" for all supported.
t_star	(Optional) Evaluation time for Brier score, AUC, point R2, and calibration. Defaults to the median observed follow-up.
tau	(Optional) The maximum prediction time horizon for R2 and C index. Default is maximum of observation time.
event_type	Integer code for the cause of interest (passed through).
digits	Number of decimal places to round the metric values (default = 2).
censoring	Censoring estimator for generalized R-squared metrics, "cox" (default) or "km"; passed to <code>pa_survival_eval_cr()</code> .
censor_covariates	Optional numeric censoring-model covariates shared across models. Rows must match each model's subject order. Passed to <code>pa_survival_eval_cr()</code> ; used only with <code>censoring = "cox"</code> .
calibration_args	Optional named list passed to <code>pa_calibration_cr()</code> .

Details

See `pa_survival_eval_cr()` for the event-observation weighting rule, original-time support requirements and probability floor.

Value

A data.frame with:

- each row = metric
- each column = model name
- cell values = metric values

Examples

```
# Competing risks: cause 1 is the event of interest, cause 2 the competing
# risk. Both cause-specific hazards are modelled.
cr <- pa_simulate_fine_gray(n = 150, v = 2, beta1 = c(0.5, -0.5),
                           beta2 = c(-0.3, 0.3), censor = 0.3, seed = 2025)

cr$time <- cr$obs.times
cr$status <- cr$obs.event
m1 <- survival::coxph(survival::Surv(time, status == 1) ~ X1 + X2,
                     data = cr, x = TRUE, y = TRUE)
m2 <- survival::coxph(survival::Surv(time, status == 2) ~ X1 + X2,
                     data = cr, x = TRUE, y = TRUE)
p <- pa_predict_cif(model1 = m1, model2 = m2, newdata = cr,
                   covs = c("X1", "X2"), event.type = 1, tau = max(cr$time))

pa_summarize_cr(list(csh = p), event_type = 1)

# External CIF matrix: rows are grid times, columns are subjects.
```

```
external_cif <- p$CIF_pred[, -1, drop = FALSE]
external_grid <- p$CIF_pred[, 1]
external <- list(cif_pred = cbind(time = external_grid, external_cif),
               times = p$times, status = p$status)
pa_summarize_cr(list(external = external), metrics = "brier_score")
```

pa_survival_eval

*Compute Survival Model Evaluation Metrics***Description**

This function calculates various predictive performance metrics for survival models based on predicted survival probabilities and observed survival data. It supports a range of evaluation measures, including explained variation, concordance indices, Brier Score, and time-dependent AUC.

Usage

```
pa_survival_eval(
  model,
  event_time,
  predicted_probability,
  pred_mean_survival = NULL,
  status,
  covariates,
  new_data = NULL,
  metrics = NULL,
  t_star = NULL,
  tau = NULL,
  calibration_args = list()
)
```

Arguments

model	A fitted survival model object. Optional for the metrics computed directly from predicted_probability.
event_time	A numeric vector of observed survival times.
predicted_probability	Numeric matrix of predicted survival probabilities, with <ul style="list-style-type: none"> rows = subjects. columns = sorted observed follow-up times, including duplicates
pred_mean_survival	Optional numeric vector of predicted mean survival times, used for explained variation and concordance. Predicted probabilities are still needed for point metrics, Brier score, AUC, and calibration.
status	A numeric vector indicating event occurrence (1 for event, 0 for censoring).
covariates	A character vector specifying the names of the covariates used in the model.

new_data	Optional data frame used by metrics that require refitting or prediction from model. If supplied, it should contain the variables needed by those procedures and columns time and status coded as above.
metrics	<p>A character vector specifying the evaluation metrics to compute. Options include:</p> <ul style="list-style-type: none"> • "pseudo_r2" - Pseudo R-squared measure • "r_square" - Explained variation R^2 • "l_square" - L-squared measure • "harrell_c" - Harrell's concordance index • "uno_c" - Uno's concordance index • "brier_score" - Overall probability prediction error • "td_auc" - Time-dependent area under the curve (AUC) • "ici", "e50", "e90" - Absolute calibration error summaries <p>The default NULL includes ICI but omits E50 and E90. Use "all" to include every supported metric.</p>
t_star	(Optional) Evaluation time for point R-squared, Brier score, time-dependent AUC, and calibration. If NULL, it defaults to the median observed follow-up time.
tau	(Optional) A numeric value specifying the truncation time for calculating explained variation metrics. If NULL, it defaults to the maximum observed survival time.
calibration_args	Optional named list passed to pa_calibration() , e.g. <code>list(nk = 3)</code> . These settings affect calibration only. For sampling weights use pa_sample_design() or pa_calibration() directly, rather than passing weights through this list.

Details

Point R-squared uses the event indicator $I(T \leq t_{\text{star}})$ and the predicted event probability $1 - S$ at the selected prediction-grid column. Events at the evaluation time are included. Calibration uses $1 - S(t_{\text{star}})$ from the last supplied grid column at or before t_{star} , interpreting the grid as a right-continuous step function (risk zero before its first time). Its outcome horizon is exactly t_{star} , not the nearest future grid time. See [pa_calibration\(\)](#) for the estimator and boundary-probability policy. A failed calibration fit warns and returns missing calibration metrics; other requested metrics are retained.

Value

A data frame with two columns:

Metric	The name of the computed evaluation metric.
Value	The corresponding computed value.

References

- Schemper, M. and R. Henderson (2000). Predictive accuracy and explained variation in Cox regression. *Biometrics* 56, 249–255.
- Lusa, L., R. Miceli and L. Mariani (2007). Estimation of predictive accuracy in survival analysis using R and S-PLUS. *Computer Methods and Programs in Biomedicine* 87, 132–137.
- Potapov, S., Adler, W., Schmid, M., Bertrand, F. (2024). survAUC: Estimating Time-Dependent AUC for Censored Survival Data. R package version 1.3-0. DOI: [doi:10.32614/CRAN.package.survAUC](https://doi.org/10.32614/CRAN.package.survAUC). Available at <https://CRAN.R-project.org/package=survAUC>.

Examples

```
# A Cox model evaluated on right-censored data. The simulator is seeded,
# so the reported metric values are reproducible.
d <- pa_sim_cox_weibull(n = 150, pi_c = 0.3, v = 2,
  beta = c(0.5, -0.5), seed = 2025)
d <- d[, c("time", "status", "x1", "x2")]
fit <- survival::coxph(survival::Surv(time, status) ~ x1 + x2,
  data = d, x = TRUE, y = TRUE)
pred <- pa_predict_coxph(model = fit, covs = c("x1", "x2"), new_data = d)

# the default metric set
pa_survival_eval(
  model = fit, event_time = pred$times,
  predicted_probability = pred$urv_prob, status = pred$status,
  covariates = c("x1", "x2"), new_data = d
)

# a chosen subset; see pa_metric_names() for the accepted identifiers
pa_survival_eval(
  model = fit, event_time = pred$times,
  predicted_probability = pred$urv_prob, status = pred$status,
  covariates = c("x1", "x2"), new_data = d,
  metrics = c("harrell_c", "brier_score")
)
```

pa_survival_eval_cr *Evaluate Survival Prediction Performance with Competing Risks*

Description

This function evaluates the predictive performance of a survival model in the presence of competing risks. It calculates various performance metrics including Pseudo R-squared, R-squared, L-squared, C-index, Brier Score, and Time-Dependent AUC.

Usage

```
pa_survival_eval_cr(
  pred_cif,
  event_time,
  time.cif,
  status,
  metrics = NULL,
  t_star = NULL,
  tau = NULL,
  event_type = 1,
  censoring = c("cox", "km"),
  censor_covariates = NULL,
  calibration_args = list()
)
```

Arguments

<code>pred_cif</code>	Numeric matrix of cumulative incidence functions with one row per value of <code>time.cif</code> and one column per subject.
<code>event_time</code>	A numeric vector of observed event or censoring times for subjects.
<code>time.cif</code>	A numeric vector of time points corresponding to the predicted CIFs.
<code>status</code>	A numeric vector indicating the event type for each subject: by default <ul style="list-style-type: none"> • 0 for censored cases. • 1 for the primary event of interest. • 2 (or other values) for competing events.
<code>metrics</code>	A character vector specifying the performance metrics to calculate. The default NULL computes pseudo R-squared (integrated and point), C-index, Brier score, time-dependent AUC, and ICI. Use "all" to include the components and E50/E90 as well. Valid options include: <ul style="list-style-type: none"> • "pseudo_r2" • "r_square" • "l_square" • "pseudo_r2_point" • "r2_point" • "l2_point" • "c_index" • "brier_score" • "td_auc" • "ici", "e50", "e90"
<code>t_star</code>	Optional numeric value specifying a reference time for evaluating point-version Pseudo_R_square, the Brier score, time-dependent AUC, and calibration. Defaults to the median of <code>event_time</code> .
<code>tau</code>	Optional numeric value specifying a restricted time horizon for model evaluation. Defaults to the maximum observed follow-up time.

event_type	An integer indicating the primary event type of interest (default = 1).
censoring	Censoring estimator for the generalized R-squared metrics: "cox" (default, optionally conditional on censor_covariates) or "km". Other metrics retain their own censoring adjustments.
censor_covariates	Optional numeric vector, matrix or data frame of censoring-model covariates, in the same subject order as event_time and the columns of pred_cif. Requires censoring = "cox". With NULL, the Cox censoring model has no covariates. The Cox censoring model uses Breslow ties, so without covariates it equals exp(-Nelson-Aalen).
calibration_args	Optional named list passed to pa_calibration_cr() , e.g. list(bandwidth = 0.2). These settings affect calibration only.

Details

The generalized R-squared metrics use event-observation IPCW. Censoring is estimated on the original follow-up times, treating any observed event type as an observed event. The unnormalized weight is $I(status_i > 0)/G(Y_i - | X_i)$; weights are normalized to sum to one. A censored subject has zero weight even if follow-up exceeds tau. Restricting the working outcome therefore does not limit censoring support requirements to tau: support at contributing original event times is needed. Cox weights require conditional independent censoring and a valid censoring model; marginal weights require the corresponding marginal independence. Censoring survival probabilities are floored at 10^{-4} . The interval outcome is the target-event time capped at tau, with competing events assigned to tau. The point outcome is $I(T \leq t_{\text{star}}, D = \text{event_type})$, including target events at the evaluation time, as in the CIF. Normalized ratio estimates are not asserted to be exactly unbiased in finite samples. Calibration uses the target-event CIF at or before t_{star} on the supplied grid and evaluates the outcome at t_{star} . It uses the local Aalen-Johansen estimator described in [pa_calibration_cr\(\)](#); censoring and censor_covariates do not control this estimator. A failed curve fit warns and returns missing calibration metrics without discarding others.

Value

A data frame with two columns:

- "Metric": Name of the evaluated metric.
- "Value": Computed value of the corresponding metric.

Examples

```
# Competing risks: cause 1 is the event of interest, cause 2 the competing
# risk. Both cause-specific hazards are modelled.
cr <- pa_simulate_fine_gray(n = 150, v = 2, beta1 = c(0.5, -0.5),
                           beta2 = c(-0.3, 0.3), censor = 0.3, seed = 2025)

cr$time <- cr$obs.times
cr$status <- cr$obs.event
m1 <- survival::coxph(survival::Surv(time, status == 1) ~ X1 + X2,
                      data = cr, x = TRUE, y = TRUE)
m2 <- survival::coxph(survival::Surv(time, status == 2) ~ X1 + X2,
```

```
      data = cr, x = TRUE, y = TRUE)
p <- pa_predict_cif(model1 = m1, model2 = m2, newdata = cr,
  covs = c("X1", "X2"), event.type = 1, tau = max(cr$time))

pa_survival_eval_cr(
  pred_cif = p$cif_pred[, -1], event_time = p$times,
  time.cif = p$cif_pred[, 1], status = p$status, event_type = 1
)
```

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