

# Package ‘CohortMethod’

August 14, 2022

**Type** Package

**Title** New-User Cohort Method with Large Scale Propensity and Outcome Models

**Version** 4.2.3

**Date** 2022-08-12

**Maintainer** Martijn Schuemie <schuemie@ohdsi.org>

**Description** Functions for performing new-user cohort studies in an observational database in the OMOP Common Data Model. Can extract the necessary data from a database and use a large set of covariates for both the propensity and outcome model, including for example all drugs, diagnoses, procedures, as well as age, comorbidity indexes, etc. Large scale regularized regression is used to fit the propensity and outcome models. Functions are included for trimming, stratifying, (variable and fixed ratio) matching and weighting by propensity scores, as well as diagnostic functions, such as propensity score distribution plots and plots showing covariate balance before and after matching and/or trimming. Supported outcome models are (conditional) logistic regression, (conditional) Poisson regression, and (stratified) Cox regression. Also included are Kaplan-Meier plots that can adjust for the stratification or matching.

**License** Apache License 2.0

**VignetteBuilder** knitr

**URL** <https://ohdsi.github.io/CohortMethod>, <https://github.com/OHDSI/CohortMethod>

**BugReports** <https://github.com/OHDSI/CohortMethod/issues>

**Depends** R (>= 3.5.0),  
DatabaseConnector (>= 4.0.0),  
Cyclops (>= 3.1.2),  
FeatureExtraction (>= 3.0.0),  
Andromeda (>= 0.5.0)

**Imports** methods,  
ggplot2,  
gridExtra,  
grid,  
readr,  
plyr,  
dplyr,  
rlang,  
cli,  
pillar,

Rcpp ( $\geq 0.11.2$ ),  
 SqlRender ( $\geq 1.7.0$ ),  
 survival,  
 ParallelLogger ( $\geq 2.0.0$ ),  
 bit64

**Suggests** testthat,  
 pROC,  
 knitr,  
 rmarkdown,  
 EmpiricalCalibration,  
 Eunomia,  
 withr,  
 R.utils

**Remotes** ohdsi/FeatureExtraction,  
 ohdsi/Eunomia

**LinkingTo** Rcpp

**NeedsCompilation** yes

**RoxygenNote** 7.1.2

**Roxygen** list(markdown = TRUE)

**Encoding** UTF-8

## R topics documented:

adjustedKm . . . . .	3
checkCmInstallation . . . . .	4
CohortMethodData-class . . . . .	4
cohortMethodDataSimulationProfile . . . . .	5
computeCovariateBalance . . . . .	5
computeMdr . . . . .	6
computePsAuc . . . . .	7
createCmAnalysis . . . . .	8
createCmTable1 . . . . .	10
createCohortMethodDataSimulationProfile . . . . .	11
createCreatePsArgs . . . . .	11
createCreateStudyPopulationArgs . . . . .	12
createFitOutcomeModelArgs . . . . .	14
createGetDbCohortMethodDataArgs . . . . .	15
createMatchOnPsAndCovariatesArgs . . . . .	16
createMatchOnPsArgs . . . . .	17
createPs . . . . .	18
createStratifyByPsAndCovariatesArgs . . . . .	19
createStratifyByPsArgs . . . . .	19
createStudyPopulation . . . . .	20
createTargetComparatorOutcomes . . . . .	22
createTrimByIptwArgs . . . . .	23
createTrimByPsArgs . . . . .	23
createTrimByPsToEquipoiseArgs . . . . .	24
drawAttritionDiagram . . . . .	24
fitOutcomeModel . . . . .	25
getAttritionTable . . . . .	26

getDbCohortMethodData . . . . .	27
getDefaultCmTable1Specifications . . . . .	29
getFollowUpDistribution . . . . .	30
getOutcomeModel . . . . .	30
getPsModel . . . . .	31
isCohortMethodData . . . . .	31
loadCmAnalysisList . . . . .	32
loadCohortMethodData . . . . .	32
loadTargetComparatorOutcomesList . . . . .	33
matchOnPs . . . . .	33
matchOnPsAndCovariates . . . . .	35
plotCovariateBalanceOfTopVariables . . . . .	36
plotCovariateBalanceScatterPlot . . . . .	37
plotCovariatePrevalence . . . . .	38
plotFollowUpDistribution . . . . .	39
plotKaplanMeier . . . . .	40
plotPs . . . . .	41
plotTimeToEvent . . . . .	43
runCmAnalyses . . . . .	45
saveCmAnalysisList . . . . .	48
saveCohortMethodData . . . . .	48
saveTargetComparatorOutcomesList . . . . .	49
simulateCohortMethodData . . . . .	49
stratifyByPs . . . . .	50
stratifyByPsAndCovariates . . . . .	51
summarizeAnalyses . . . . .	52
trimByIptw . . . . .	52
trimByPs . . . . .	53
trimByPsToEquipoise . . . . .	54
<b>Index</b>	<b>55</b>

adjustedKm

*Compute a weight-adjusted Kaplan-Meier curve***Description**

Compute a weight-adjusted Kaplan-Meier curve

**Usage**

```
adjustedKm(weight, time, y)
```

**Arguments**

weight	Vector of observation weights
time	Vector of event times
y	Vector outcomes (0 indicates censoring, 1 indicates event-of-interest)



---

cohortMethodDataSimulationProfile

*A simulation profile*


---

### Description

A simulation profile

### Usage

```
data(cohortMethodDataSimulationProfile)
```

---

computeCovariateBalance

*Compute covariate balance before and after matching and trimming*


---

### Description

For every covariate, prevalence in treatment and comparator groups before and after matching/trimming are computed. When variable ratio matching was used the balance score will be corrected according the method described in Austin et al (2008).

### Usage

```
computeCovariateBalance(
  population,
  cohortMethodData,
  subgroupCovariateId = NULL,
  maxCohortSize = 250000
)
```

### Arguments

- |                     |  |
|---------------------|--|
| population          | A data frame containing the people that are remaining after matching and/or trimming.  |
| cohortMethodData    | An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .  |
| subgroupCovariateId | Optional: a covariate ID of a binary covariate that indicates a subgroup of interest. Both the before and after populations will be restricted to this subgroup before computing covariate balance.        |
| maxCohortSize       | If the target or comparator cohort are larger than this number, they will be down-sampled before computing covariate balance to save time. Setting this number to 0 means no downsampling will be applied. |

## Details

The population data frame should have the following three columns:

- `rowId` (numeric): A unique identifier for each row (e.g. the person ID).
- `treatment` (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- `propensityScore` (numeric): Propensity score.

## Value

Returns a tibble describing the covariate balance before and after matching/trimming.

## References

Austin, P.C. (2008) Assessing balance in measured baseline covariates when using many-to-one matching on the propensity-score. *Pharmacoepidemiology and Drug Safety*, 17: 1218-1225.

---

<code>computeMdr</code>	<i>Compute the minimum detectable relative risk</i>
-------------------------	---

---

## Description

Compute the minimum detectable relative risk

## Usage

```
computeMdr(
  population,
  alpha = 0.05,
  power = 0.8,
  twoSided = TRUE,
  modelType = "cox"
)
```

## Arguments

<code>population</code>	A data frame describing the study population as created using the <a href="#">createStudyPopulation</a> function. This should at least have these columns: <code>personSeqId</code> , <code>treatment</code> , <code>outcomeCount</code> , <code>timeAtRisk</code> .
<code>alpha</code>	Type I error.
<code>power</code>	1 - beta, where beta is the type II error.
<code>twoSided</code>	Consider a two-sided test?
<code>modelType</code>	The type of outcome model that will be used. Possible values are "logistic", "poisson", or "cox". Currently only "cox" is supported.

## Details

Compute the minimum detectable relative risk (MDRR) and expected standard error (SE) for a given study population, using the actual observed sample size and number of outcomes. Currently, only computations for Cox models are implemented. For Cox model, the computations by Schoenfeld (1983) is used.

**Value**

A data frame with the MDRR and some counts.

**References**

Schoenfeld DA (1983) Sample-size formula for the proportional-hazards regression model, *Biometrics*, 39(3), 499-503

---

computePsAuc	<i>Compute the area under the ROC curve</i>
--------------	---

---

**Description**

Compute the area under the ROC curve of the propensity score.

**Usage**

```
computePsAuc(data, confidenceIntervals = FALSE)
```

**Arguments**

**data** A data frame with at least the two columns described below

**confidenceIntervals** Compute 95 percent confidence intervals (computationally expensive for large data sets)

**Details**

The data frame should have at least the following two columns:

- **treatment** (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- **propensityScore** (numeric): Propensity score.

**Value**

A tibble holding the AUC and its 95 percent confidence interval

**Examples**

```
treatment <- rep(0:1, each = 100)
propensityScore <- c(rnorm(100, mean = 0.4, sd = 0.25), rnorm(100, mean = 0.6, sd = 0.25))
data <- data.frame(treatment = treatment, propensityScore = propensityScore)
data <- data[data$propensityScore > 0 & data$propensityScore < 1, ]
computePsAuc(data)
```

---

createCmAnalysis	Create a CohortMethod analysis specification
------------------	--

---

## Description

Create a CohortMethod analysis specification

## Usage

```
createCmAnalysis(
  analysisId = 1,
  description = "",
  targetType = NULL,
  comparatorType = NULL,
  getDbCohortMethodDataArgs,
  createStudyPopArgs,
  createPs = FALSE,
  createPsArgs = NULL,
  trimByPs = FALSE,
  trimByPsArgs = NULL,
  trimByPsToEquipoise = FALSE,
  trimByPsToEquipoiseArgs = NULL,
  trimByIptw = FALSE,
  trimByIptwArgs = NULL,
  matchOnPs = FALSE,
  matchOnPsArgs = NULL,
  matchOnPsAndCovariates = FALSE,
  matchOnPsAndCovariatesArgs = NULL,
  stratifyByPs = FALSE,
  stratifyByPsArgs = NULL,
  stratifyByPsAndCovariates = FALSE,
  stratifyByPsAndCovariatesArgs = NULL,
  fitOutcomeModel = FALSE,
  fitOutcomeModelArgs = NULL
)
```

## Arguments

analysisId	An integer that will be used later to refer to this specific set of analysis choices.
description	A short description of the analysis.
targetType	If more than one target is provided for each drugComparatorOutcome, this field should be used to select the specific target to use in this analysis.
comparatorType	If more than one comparator is provided for each drugComparatorOutcome, this field should be used to select the specific comparator to use in this analysis.
getDbCohortMethodDataArgs	An object representing the arguments to be used when calling the <a href="#">getDbCohortMethodData()</a> function.
createStudyPopArgs	An object representing the arguments to be used when calling the <a href="#">createStudyPopulation()</a> function.



createPs	Should the <a href="#">createPs()</a> function be used in this analysis?
createPsArgs	An object representing the arguments to be used when calling the <a href="#">createPs()</a> function.
trimByPs	Should the <a href="#">trimByPs()</a> function be used in this analysis?
trimByPsArgs	An object representing the arguments to be used when calling the <a href="#">trimByPs()</a> function.
trimByPsToEquipoise	Should the <a href="#">trimByPsToEquipoise()</a> function be used in this analysis?
trimByPsToEquipoiseArgs	An object representing the arguments to be used when calling the <a href="#">trimByPsToEquipoise()</a> function.
trimByIptw	Should the <a href="#">trimByPsToEquipoise()</a> function be used in this analysis?
trimByIptwArgs	An object representing the arguments to be used when calling the <a href="#">trimByIptw()</a> function.
matchOnPs	Should the <a href="#">matchOnPs()</a> function be used in this analysis?
matchOnPsArgs	An object representing the arguments to be used when calling the <a href="#">matchOnPs()</a> function.
matchOnPsAndCovariates	Should the <a href="#">matchOnPsAndCovariates()</a> function be used in this analysis?
matchOnPsAndCovariatesArgs	An object representing the arguments to be used when calling the <a href="#">matchOnPsAndCovariates()</a> function.
stratifyByPs	Should the <a href="#">stratifyByPs()</a> function be used in this analysis?
stratifyByPsArgs	An object representing the arguments to be used when calling the <a href="#">stratifyByPs()</a> function.
stratifyByPsAndCovariates	Should the <a href="#">stratifyByPsAndCovariates()</a> function be used in this analysis?
stratifyByPsAndCovariatesArgs	An object representing the arguments to be used when calling the <a href="#">stratifyByPsAndCovariates()</a> function.
fitOutcomeModel	Should the <a href="#">fitOutcomeModel()</a> function be used in this analysis?
fitOutcomeModelArgs	An object representing the arguments to be used when calling the <a href="#">fitOutcomeModel()</a> function.

## Details

Create a set of analysis choices, to be used with the [runCmAnalyses\(\)](#) function.

createCmTable1

*Create a table 1***Description**

Creates a formatted table of cohort characteristics, to be included in publications or reports.

**Usage**

```
createCmTable1(
  balance,
  specifications = getDefaultCmTable1Specifications(),
  beforeTargetPopSize,
  beforeComparatorPopSize,
  afterTargetPopSize,
  afterComparatorPopSize,
  beforeLabel = "Before matching",
  afterLabel = "After matching",
  targetLabel = "Target",
  comparatorLabel = "Comparator",
  percentDigits = 1,
  stdDiffDigits = 2
)
```

**Arguments**

balance	A data frame created by the computeCovariateBalance function.
specifications	Specifications of which covariates to display, and how.
beforeTargetPopSize	The number of people in the target cohort before matching/stratification/trimming, to mention in the table header. If not provide, no number will be included in the header.
beforeComparatorPopSize	The number of people in the comparator cohort before matching/stratification/trimming, to mention in the table header. If not provide, no number will be included in the header.
afterTargetPopSize	The number of people in the target cohort after matching/stratification/trimming, to mention in the table header. If not provide, no number will be included in the header.
afterComparatorPopSize	The number of people in the comparator cohort after matching/stratification/trimming, to mention in the table header. If not provide, no number will be included in the header.
beforeLabel	Label for identifying columns before matching / stratification / trimming.
afterLabel	Label for identifying columns after matching / stratification / trimming.
targetLabel	Label for identifying columns of the target cohort.
comparatorLabel	Label for identifying columns of the comparator cohort.
percentDigits	Number of digits to be used for percentages.
stdDiffDigits	Number of digits to be used for the standardized differences.

**Value**

A data frame with the formatted table 1.

---

```
createCohortMethodDataSimulationProfile
```

*Create simulation profile*

---

**Description**

Creates a profile based on the provided [CohortMethodData](#) object, which can be used to generate simulated data that has similar characteristics.

**Usage**

```
createCohortMethodDataSimulationProfile(cohortMethodData)
```

**Arguments**

cohortMethodData

An object of type [CohortMethodData](#) as generated using [getDbCohortMethodData\(\)](#).

**Details**

The output of this function is an object that can be used by the [simulateCohortMethodData\(\)](#) function to generate a cohortMethodData object.

**Value**

An object of type CohortDataSimulationProfile.

---

```
createCreatePsArgs
```

*Create a parameter object for the function createPs*

---

**Description**

Create a parameter object for the function createPs

**Usage**

```
createCreatePsArgs(
  excludeCovariateIds = c(),
  includeCovariateIds = c(),
  maxCohortSizeForFitting = 250000,
  errorOnHighCorrelation = TRUE,
  stopOnError = TRUE,
  prior = createPrior("laplace", exclude = c(0), useCrossValidation = TRUE),
  control = createControl(noiseLevel = "silent", cvType = "auto", seed = 1, tolerance =
    2e-07, cvRepetitions = 10, startingVariance = 0.01)
)
```

**Arguments**

<code>excludeCovariateIds</code>	Exclude these covariates from the propensity model.
<code>includeCovariateIds</code>	Include only these covariates in the propensity model.
<code>maxCohortSizeForFitting</code>	If the target or comparator cohort are larger than this number, they will be down-sampled before fitting the propensity model. The model will be used to compute propensity scores for all subjects. The purpose of the sampling is to gain speed. Setting this number to 0 means no downsampling will be applied.
<code>errorOnHighCorrelation</code>	If true, the function will test each covariate for correlation with the treatment assignment. If any covariate has an unusually high correlation (either positive or negative), this will throw an error.
<code>stopOnError</code>	If an error occurs, should the function stop? Else, the two cohorts will be assumed to be perfectly separable.
<code>prior</code>	The prior used to fit the model. See <code>Cyclops::createPrior()</code> for details.
<code>control</code>	The control object used to control the cross-validation used to determine the hyperparameters of the prior (if applicable). See <code>Cyclops::createControl()</code> for details.

**Details**

Create an object defining the parameter values.

---

`createCreateStudyPopulationArgs`

*Create a parameter object for the function `createStudyPopulation`*

---

**Description**

Create a parameter object for the function `createStudyPopulation`

**Usage**

```
createCreateStudyPopulationArgs(
  firstExposureOnly = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  removeDuplicateSubjects = FALSE,
  removeSubjectsWithPriorOutcome = TRUE,
  priorOutcomeLookback = 99999,
  minDaysAtRisk = 1,
  maxDaysAtRisk = 99999,
  riskWindowStart = 0,
  addExposureDaysToStart = NULL,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  addExposureDaysToEnd = NULL,
```

```

    endAnchor = "cohort end",
    censorAtNewRiskWindow = FALSE
)

```

## Arguments

firstExposureOnly	Should only the first exposure per subject be included?
restrictToCommonPeriod	Restrict the analysis to the period when both exposures are observed?
washoutPeriod	The minimum required continuous observation time prior to index date for a person to be included in the cohort.
removeDuplicateSubjects	Remove subjects that are in both the target and comparator cohort? See details for allowed values.
removeSubjectsWithPriorOutcome	Remove subjects that have the outcome prior to the risk window start?
priorOutcomeLookback	How many days should we look back when identifying prior outcomes?
minDaysAtRisk	The minimum required number of days at risk. Risk windows with fewer days than this number are removed from the analysis.
maxDaysAtRisk	The maximum allowed number of days at risk. Risk windows that are longer will be truncated to this number of days.
riskWindowStart	The start of the risk window (in days) relative to the startAnchor.
addExposureDaysToStart	DEPRECATED: Add the length of exposure the start of the risk window? Use startAnchor instead.
startAnchor	The anchor point for the start of the risk window. Can be "cohort start" or "cohort end".
riskWindowEnd	The end of the risk window (in days) relative to the endAnchor.
addExposureDaysToEnd	DEPRECATED: Add the length of exposure the risk window? Use endAnchor instead.
endAnchor	The anchor point for the end of the risk window. Can be "cohort start" or "cohort end".
censorAtNewRiskWindow	If a subject is in multiple cohorts, should time-at-risk be censored when the new time-at-risk starts to prevent overlap?

## Details

Create an object defining the parameter values.

---

```
createFitOutcomeModelArgs
```

*Create a parameter object for the function fitOutcomeModel*

---

## Description

Create a parameter object for the function fitOutcomeModel

## Usage

```
createFitOutcomeModelArgs(
  modelType = "logistic",
  stratified = FALSE,
  useCovariates = FALSE,
  inversePtWeighting = FALSE,
  estimator = "ate",
  maxWeight = 0,
  interactionCovariateIds = c(),
  excludeCovariateIds = c(),
  includeCovariateIds = c(),
  profileGrid = NULL,
  profileBounds = c(log(0.1), log(10)),
  prior = createPrior("laplace", useCrossValidation = TRUE),
  control = createControl(cvType = "auto", seed = 1, startingVariance = 0.01, tolerance
    = 2e-07, cvRepetitions = 10, noiseLevel = "quiet")
)
```

## Arguments

modelType	The type of outcome model that will be used. Possible values are "logistic", "poisson", or "cox".
stratified	Should the regression be conditioned on the strata defined in the population object (e.g. by matching or stratifying on propensity scores)?
useCovariates	Whether to use the covariates in the cohortMethodData object in the outcome model.
inversePtWeighting	Use inverse probability of treatment weighting (IPTW)? See details.
estimator	for IPTW: the type of estimator. Options are estimator = "ate" for the average treatment effect, and estimator = "att" for the average treatment effect in the treated.
maxWeight	for IPTW: the maximum weight. Larger values will be truncated to this value. maxWeight = 0 means no truncation takes place.
interactionCovariateIds	An optional vector of covariate IDs to use to estimate interactions with the main treatment effect.
excludeCovariateIds	Exclude these covariates from the outcome model.
includeCovariateIds	Include only these covariates in the outcome model.

profileGrid	A one-dimensional grid of points on the log(relative risk) scale where the likelihood for coefficient of variables is sampled. See details.
profileBounds	The bounds (on the log relative risk scale) for the adaptive sampling of the likelihood function. See details.
prior	The prior used to fit the model. See Cyclops::createPrior() for details.
control	The control object used to control the cross-validation used to determine the hyperparameters of the prior (if applicable). See Cyclops::createControl() for details.

## Details

Create an object defining the parameter values.

---

```
createGetDbCohortMethodDataArgs
```

*Create a parameter object for the function getDbCohortMethodData*

---

## Description

Create a parameter object for the function getDbCohortMethodData

## Usage

```
createGetDbCohortMethodDataArgs(
  studyStartDate = "",
  studyEndDate = "",
  excludeDrugsFromCovariates = NULL,
  firstExposureOnly = FALSE,
  removeDuplicateSubjects = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  maxCohortSize = 0,
  covariateSettings
)
```

## Arguments

studyStartDate	A calendar date specifying the minimum date that a cohort index date can appear. Date format is 'yyyymmdd'.
studyEndDate	A calendar date specifying the maximum date that a cohort index date can appear. Date format is 'yyyymmdd'. Important: the study end data is also used to truncate risk windows, meaning no outcomes beyond the study end date will be considered.
excludeDrugsFromCovariates	DEPRECATED: Should the target and comparator drugs (and their descendant concepts) be excluded from the covariates? Note that this will work if the drugs are actually drug concept IDs (and not cohort IDs).
firstExposureOnly	Should only the first exposure per subject be included? Note that this is typically done in the createStudyPopulation() function, but can already be done here for efficiency reasons.

removeDuplicateSubjects	Remove subjects that are in both the target and comparator cohort? See details for allowed values. Note that this is typically done in the createStudyPopulation function, but can already be done here for efficiency reasons.
restrictToCommonPeriod	Restrict the analysis to the period when both treatments are observed?
washoutPeriod	The minimum required continuous observation time prior to index date for a person to be included in the cohort. Note that this is typically done in the createStudyPopulation function, but can already be done here for efficiency reasons.
maxCohortSize	If either the target or the comparator cohort is larger than this number it will be sampled to this size. maxCohortSize = 0 indicates no maximum size.
covariateSettings	An object of type covariateSettings as created using the FeatureExtraction::createCovariateSettings() function.

## Details

Create an object defining the parameter values.

---

```
createMatchOnPsAndCovariatesArgs
```

*Create a parameter object for the function matchOnPsAndCovariates*

---

## Description

Create a parameter object for the function matchOnPsAndCovariates

## Usage

```
createMatchOnPsAndCovariatesArgs(
  caliper = 0.2,
  caliperScale = "standardized logit",
  maxRatio = 1,
  allowReverseMatch = FALSE,
  covariateIds
)
```

## Arguments

caliper	The caliper for matching. A caliper is the distance which is acceptable for any match. Observations which are outside of the caliper are dropped. A caliper of 0 means no caliper is used.
caliperScale	The scale on which the caliper is defined. Three scales are supported: caliperScale = 'propensity score', caliperScale = 'standardized', or caliperScale = 'standardized logit'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution. 'standardized logit' is similar, except that the propensity score is transformed to the logit scale because the PS is more likely to be normally distributed on that scale (Austin, 2011).
maxRatio	The maximum number of persons in the comparator arm to be matched to each person in the treatment arm. A maxRatio of 0 means no maximum: all comparators will be assigned to a target person.



allowReverseMatch	Allows n-to-1 matching if target arm is larger
covariateIds	One or more covariate IDs in the cohortMethodData object on which subjects should be also matched.

## Details

Create an object defining the parameter values.

---

createMatchOnPsArgs	<i>Create a parameter object for the function matchOnPs</i>
---------------------	---

---

## Description

Create a parameter object for the function matchOnPs

## Usage

```
createMatchOnPsArgs(
  caliper = 0.2,
  caliperScale = "standardized logit",
  maxRatio = 1,
  allowReverseMatch = FALSE,
  stratificationColumns = c()
)
```

## Arguments

caliper	The caliper for matching. A caliper is the distance which is acceptable for any match. Observations which are outside of the caliper are dropped. A caliper of 0 means no caliper is used.
caliperScale	The scale on which the caliper is defined. Three scales are supported: caliperScale = 'propensity score', caliperScale = 'standardized', or caliperScale = 'standardized logit'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution. 'standardized logit' is similar, except that the propensity score is transformed to the logit scale because the PS is more likely to be normally distributed on that scale (Austin, 2011).
maxRatio	The maximum number of persons in the comparator arm to be matched to each person in the treatment arm. A maxRatio of 0 means no maximum: all comparators will be assigned to a target person.
allowReverseMatch	Allows n-to-1 matching if target arm is larger
stratificationColumns	Names or numbers of one or more columns in the data data.frame on which subjects should be stratified prior to matching. No persons will be matched with persons outside of the strata identified by the values in these columns.

## Details

Create an object defining the parameter values.

createPs

*Create propensity scores***Description**

Creates propensity scores using a regularized logistic regression.

**Usage**

```
createPs(
  cohortMethodData,
  population = NULL,
  excludeCovariateIds = c(),
  includeCovariateIds = c(),
  maxCohortSizeForFitting = 250000,
  errorOnHighCorrelation = TRUE,
  stopOnError = TRUE,
  prior = createPrior("laplace", exclude = c(), useCrossValidation = TRUE),
  control = createControl(noiseLevel = "silent", cvType = "auto", seed = 1, tolerance =
    2e-07, cvRepetitions = 10, startingVariance = 0.01)
)
```

**Arguments**

cohortMethodData	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .
population	A data frame describing the population. This should at least have a rowId column corresponding to the rowId column in the <a href="#">CohortMethodData</a> covariates object and a treatment column. If population is not specified, the full population in the <a href="#">CohortMethodData</a> will be used.
excludeCovariateIds	Exclude these covariates from the propensity model.
includeCovariateIds	Include only these covariates in the propensity model.
maxCohortSizeForFitting	If the target or comparator cohort are larger than this number, they will be down-sampled before fitting the propensity model. The model will be used to compute propensity scores for all subjects. The purpose of the sampling is to gain speed. Setting this number to 0 means no downsampling will be applied.
errorOnHighCorrelation	If true, the function will test each covariate for correlation with the treatment assignment. If any covariate has an unusually high correlation (either positive or negative), this will throw an error.
stopOnError	If an error occurs, should the function stop? Else, the two cohorts will be assumed to be perfectly separable.
prior	The prior used to fit the model. See <a href="#">Cyclops::createPrior()</a> for details.
control	The control object used to control the cross-validation used to determine the hyperparameters of the prior (if applicable). See <a href="#">Cyclops::createControl()</a> for details.

**Examples**

```
data(cohortMethodDataSimulationProfile)
cohortMethodData <- simulateCohortMethodData(cohortMethodDataSimulationProfile, n = 1000)
ps <- createPs(cohortMethodData)
```

---

```
createStratifyByPsAndCovariatesArgs
```

*Create a parameter object for the function stratifyByPsAndCovariates*

---

**Description**

Create a parameter object for the function stratifyByPsAndCovariates

**Usage**

```
createStratifyByPsAndCovariatesArgs(
  numberOfStrata = 5,
  baseSelection = "all",
  covariateIds
)
```

**Arguments**

numberOfStrata	Into how many strata should the propensity score be divided? The boundaries of the strata are automatically defined to contain equal numbers of target persons.
baseSelection	What is the base selection of subjects where the strata bounds are to be determined? Strata are defined as equally-sized strata inside this selection. Possible values are "all", "target", and "comparator".
covariateIds	One or more covariate IDs in the cohortMethodData object on which subjects should also be stratified.

**Details**

Create an object defining the parameter values.

---

```
createStratifyByPsArgs
```

*Create a parameter object for the function stratifyByPs*

---

**Description**

Create a parameter object for the function stratifyByPs

**Usage**

```
createStratifyByPsArgs(
  numberOfStrata = 5,
  stratificationColumns = c(),
  baseSelection = "all"
)
```

**Arguments**

- numberOfStrata** How many strata? The boundaries of the strata are automatically defined to contain equal numbers of target persons.
- stratificationColumns** Names of one or more columns in the data data.frame on which subjects should also be stratified in addition to stratification on propensity score.
- baseSelection** What is the base selection of subjects where the strata bounds are to be determined? Strata are defined as equally-sized strata inside this selection. Possible values are "all", "target", and "comparator".

**Details**

Create an object defining the parameter values.

---

`createStudyPopulation` *Create a study population*

---

**Description**

Create a study population

**Usage**

```
createStudyPopulation(
  cohortMethodData,
  population = NULL,
  outcomeId,
  firstExposureOnly = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  removeDuplicateSubjects = FALSE,
  removeSubjectsWithPriorOutcome = TRUE,
  priorOutcomeLookback = 99999,
  minDaysAtRisk = 1,
  maxDaysAtRisk = 99999,
  riskWindowStart = 0,
  addExposureDaysToStart = NULL,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  addExposureDaysToEnd = NULL,
  endAnchor = "cohort end",
  censorAtNewRiskWindow = FALSE
)
```

**Arguments**

- cohortMethodData** An object of type [CohortMethodData](#) as generated using [getDbCohortMethodData\(\)](#).
- population** If specified, this population will be used as the starting point instead of the cohorts in the cohortMethodData object.

outcomeId	The ID of the outcome. If not specified, no outcome-specific transformations will be performed.
firstExposureOnly	Should only the first exposure per subject be included?
restrictToCommonPeriod	Restrict the analysis to the period when both exposures are observed?
washoutPeriod	The minimum required continuous observation time prior to index date for a person to be included in the cohort.
removeDuplicateSubjects	Remove subjects that are in both the target and comparator cohort? See details for allowed values.
removeSubjectsWithPriorOutcome	Remove subjects that have the outcome prior to the risk window start?
priorOutcomeLookback	How many days should we look back when identifying prior outcomes?
minDaysAtRisk	The minimum required number of days at risk. Risk windows with fewer days than this number are removed from the analysis.
maxDaysAtRisk	The maximum allowed number of days at risk. Risk windows that are longer will be truncated to this number of days.
riskWindowStart	The start of the risk window (in days) relative to the startAnchor.
addExposureDaysToStart	DEPRECATED: Add the length of exposure the start of the risk window? Use startAnchor instead.
startAnchor	The anchor point for the start of the risk window. Can be "cohort start" or "cohort end".
riskWindowEnd	The end of the risk window (in days) relative to the endAnchor.
addExposureDaysToEnd	DEPRECATED: Add the length of exposure the risk window? Use endAnchor instead.
endAnchor	The anchor point for the end of the risk window. Can be "cohort start" or "cohort end".
censorAtNewRiskWindow	If a subject is in multiple cohorts, should time-at-risk be censored when the new time-at-risk starts to prevent overlap?

## Details

Create a study population by enforcing certain inclusion and exclusion criteria, defining a risk window, and determining which outcomes fall inside the risk window.

The removeduplicateSubjects argument can have one of the following values:

- "keep all": Do not remove subjects that appear in both target and comparator cohort
- "keep first": When a subjects appear in both target and comparator cohort, only keep whichever cohort is first in time. If both cohorts start simultaneous, the person is removed from the analysis.
- "remove all": Remove subjects that appear in both target and comparator cohort completely from the analysis."

**Value**

A tibble specifying the study population. This tibble will have the following columns:

- `rowId`: A unique identifier for an exposure.
- `personSeqId`: The person sequence ID of the subject.
- `cohortStartDate`: The index date.
- `outcomeCount`: The number of outcomes observed during the risk window.
- `timeAtRisk`: The number of days in the risk window.
- `survivalTime`: The number of days until either the outcome or the end of the risk window.

---

```
createTargetComparatorOutcomes
```

*Create target-comparator-outcomes combinations.*

---

**Description**

Create target-comparator-outcomes combinations.

**Usage**

```
createTargetComparatorOutcomes(
  targetId,
  comparatorId,
  outcomeIds,
  excludedCovariateConceptIds = c(),
  includedCovariateConceptIds = c()
)
```

**Arguments**

- |  |   |
|--|---|
| <code>targetId</code>                    | A concept ID identifying the target drug in the exposure table. If multiple strategies for picking the target will be tested in the analysis, a named list of numbers can be provided instead. In the analysis, the name of the number to be used can be specified using the <code>#'</code> <code>targetType</code> parameter in the <code>createCmAnalysis()</code> function.             |
| <code>comparatorId</code>                | A concept ID identifying the comparator drug in the exposure table. If multiple strategies for picking the comparator will be tested in the analysis, a named list of numbers can be provided instead. In the analysis, the name of the number to be used can be specified using the <code>#'</code> <code>comparatorType</code> parameter in the <code>createCmAnalysis()</code> function. |
| <code>outcomeIds</code>                  | A vector of concept IDs identifying the outcome(s) in the outcome table.  |
| <code>excludedCovariateConceptIds</code> | A list of concept IDs that cannot be used to construct covariates. This argument is to be used only for exclusion concepts that are specific to the drug-comparator combination.  |
| <code>includedCovariateConceptIds</code> | A list of concept IDs that must be used to construct covariates. This argument is to be used only for inclusion concepts that are specific to the drug-comparator combination.  |

**Details**

Create a set of hypotheses of interest, to be used with the `runCmAnalyses()` function.

---

`createTrimByIptwArgs`    *Create a parameter object for the function `trimByIptw`*

---

**Description**

Create a parameter object for the function `trimByIptw`

**Usage**

```
createTrimByIptwArgs(maxWeight = 10, estimator = "ate")
```

**Arguments**

<code>maxWeight</code>	The maximum allowed IPTW.
<code>estimator</code>	The type of estimator. Options are <code>estimator = "ate"</code> for the average treatment effect, and <code>estimator = "att"</code> for the average treatment effect in the treated.

**Details**

Create an object defining the parameter values.

---

`createTrimByPsArgs`    *Create a parameter object for the function `trimByPs`*

---

**Description**

Create a parameter object for the function `trimByPs`

**Usage**

```
createTrimByPsArgs(trimFraction = 0.05)
```

**Arguments**

<code>trimFraction</code>	This fraction will be removed from each treatment group. In the target group, persons with the highest propensity scores will be removed, in the comparator group person with the lowest scores will be removed.
---------------------------	--

**Details**

Create an object defining the parameter values.

---

```
createTrimByPsToEquipoiseArgs
```

*Create a parameter object for the function trimByPsToEquipoise*

---

### Description

Create a parameter object for the function trimByPsToEquipoise

### Usage

```
createTrimByPsToEquipoiseArgs(bounds = c(0.3, 0.7))
```

### Arguments

bounds	The upper and lower bound on the preference score for keeping persons.
--------	--

### Details

Create an object defining the parameter values.

---

```
drawAttritionDiagram
```

*Draw the attrition diagram*

---

### Description

drawAttritionDiagram draws the attrition diagram, showing how many people were excluded from the study population, and for what reasons.

### Usage

```
drawAttritionDiagram(
  object,
  targetLabel = "Target",
  comparatorLabel = "Comparator",
  fileName = NULL
)
```

### Arguments

object	Either an object of type cohortMethodData, a population object generated by functions like createStudyPopulation, or an object of type outcomeModel.
targetLabel	A label to us for the target cohort.
comparatorLabel	A label to us for the comparator cohort.
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function ggsave in the ggplot2 package for supported file formats.

### Value

A ggplot object. Use the [ggsave](#) function to save to file in a different format.



---

fitOutcomeModel	Create an outcome model, and compute the relative risk
-----------------	--

---

## Description

Create an outcome model, and computes the relative risk

## Usage

```
fitOutcomeModel(
  population,
  cohortMethodData = NULL,
  modelType = "logistic",
  stratified = FALSE,
  useCovariates = FALSE,
  inversePtWeighting = FALSE,
  estimator = "ate",
  maxWeight = 0,
  interactionCovariateIds = c(),
  excludeCovariateIds = c(),
  includeCovariateIds = c(),
  profileGrid = NULL,
  profileBounds = c(log(0.1), log(10)),
  prior = createPrior("laplace", useCrossValidation = TRUE),
  control = createControl(cvType = "auto", seed = 1, startingVariance = 0.01, tolerance
    = 2e-07, cvRepetitions = 10, noiseLevel = "quiet")
)
```

## Arguments

population	A population object generated by <a href="#">createStudyPopulation()</a> , potentially filtered by other functions.
cohortMethodData	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> . Can be omitted if not using covariates and not using interaction terms.
modelType	The type of outcome model that will be used. Possible values are "logistic", "poisson", or "cox".
stratified	Should the regression be conditioned on the strata defined in the population object (e.g. by matching or stratifying on propensity scores)?
useCovariates	Whether to use the covariates in the cohortMethodData object in the outcome model.
inversePtWeighting	Use inverse probability of treatment weighting (IPTW)? See details.
estimator	for IPTW: the type of estimator. Options are estimator = "ate" for the average treatment effect, and estimator = "att" for the average treatment effect in the treated.
maxWeight	for IPTW: the maximum weight. Larger values will be truncated to this value. maxWeight = 0 means no truncation takes place.

interactionCovariateIds	An optional vector of covariate IDs to use to estimate interactions with the main treatment effect.
excludeCovariateIds	Exclude these covariates from the outcome model.
includeCovariateIds	Include only these covariates in the outcome model.
profileGrid	A one-dimensional grid of points on the log(relative risk) scale where the likelihood for coefficient of variables is sampled. See details.
profileBounds	The bounds (on the log relative risk scale) for the adaptive sampling of the likelihood function. See details.
prior	The prior used to fit the model. See <a href="#">Cyclops::createPrior()</a> for details.
control	The control object used to control the cross-validation used to determine the hyperparameters of the prior (if applicable). See <a href="#">Cyclops::createControl()</a> for details.

### Details

IPTW estimates either the average treatment effect (ate) or average treatment effect in the treated (att) using stabilized inverse propensity scores (Xu et al. 2010).

For likelihood profiling, either specify the profileGrid for a completely user- defined grid, or profileBounds for an adaptive grid. Both should be defined on the log effect size scale. When both profileGrid and profileGrid are NULL likelihood profiling is disabled.

### Value

An object of class OutcomeModel. Generic function print, coef, and confint are available.

### References

Xu S, Ross C, Raebel MA, Shetterly S, Blanchette C, Smith D. Use of stabilized inverse propensity scores as weights to directly estimate relative risk and its confidence intervals. Value Health. 2010;13(2):273-277. doi:10.1111/j.1524-4733.2009.00671.x

---

getAttritionTable	<i>Get the attrition table for a population</i>
-------------------	---

---

### Description

Get the attrition table for a population

### Usage

```
getAttritionTable(object)
```

### Arguments

object	Either an object of type <a href="#">CohortMethodData</a> , a population object generated by functions like <a href="#">createStudyPopulation()</a> , or an object of type outcomeModel.
--------	--

**Value**

A tibble specifying the number of people and exposures in the population after specific steps of filtering.

---

getDbCohortMethodData *Get the cohort data from the server*

---

**Description**

This function executes a large set of SQL statements against the database in OMOP CDM format to extract the data needed to perform the analysis.

**Usage**

```
getDbCohortMethodData(
  connectionDetails,
  cdmDatabaseSchema,
  oracleTempSchema = NULL,
  tempEmulationSchema = getOption("sqlRenderTempEmulationSchema"),
  targetId,
  comparatorId,
  outcomeIds,
  studyStartDate = "",
  studyEndDate = "",
  exposureDatabaseSchema = cdmDatabaseSchema,
  exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence",
  cdmVersion = "5",
  excludeDrugsFromCovariates = NULL,
  firstExposureOnly = FALSE,
  removeDuplicateSubjects = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  maxCohortSize = 0,
  covariateSettings
)
```

**Arguments**

connectionDetails

An R object of type connectionDetails created using the [DatabaseConnector::createConnection](#) function.

cdmDatabaseSchema

The name of the database schema that contains the OMOP CDM instance. Requires read permissions to this database. On SQL Server, this should specify both the database and the schema, so for example 'cdm\_instance.dbo'.

oracleTempSchema

DEPRECATED: use tempEmulationSchema instead.

tempEmulationSchema	Some database platforms like Oracle and Impala do not truly support temp tables. To emulate temp tables, provide a schema with write privileges where temp tables can be created.
targetId	A unique identifier to define the target cohort. If exposureTable = DRUG_ERA, targetId is a concept ID and all descendant concepts within that concept ID will be used to define the cohort. If exposureTable <> DRUG_ERA, targetId is used to select the COHORT_DEFINITION_ID in the cohort-like table.
comparatorId	A unique identifier to define the comparator cohort. If exposureTable = DRUG_ERA, comparatorId is a concept ID and all descendant concepts within that concept ID will be used to define the cohort. If exposureTable <> DRUG_ERA, comparatorId is used to select the COHORT_DEFINITION_ID in the cohort-like table.
outcomeIds	A list of cohort IDs used to define outcomes.
studyStartDate	A calendar date specifying the minimum date that a cohort index date can appear. Date format is 'yyyymmdd'.
studyEndDate	A calendar date specifying the maximum date that a cohort index date can appear. Date format is 'yyyymmdd'. Important: the study end data is also used to truncate risk windows, meaning no outcomes beyond the study end date will be considered.
exposureDatabaseSchema	The name of the database schema that is the location where the exposure data used to define the exposure cohorts is available.
exposureTable	The tablename that contains the exposure cohorts. If exposureTable <> DRUG_ERA, then expectation is exposureTable has format of COHORT table: COHORT_DEFINITION_ID, SUBJECT_ID, COHORT_START_DATE, COHORT_END_DATE.
outcomeDatabaseSchema	The name of the database schema that is the location where the data used to define the outcome cohorts is available.
outcomeTable	The tablename that contains the outcome cohorts. If outcomeTable <> CONDITION_OCCURRENCE, then expectation is outcomeTable has format of COHORT table: COHORT_DEFINITION_ID, SUBJECT_ID, COHORT_START_DATE, COHORT_END_DATE.
cdmVersion	Define the OMOP CDM version used: currently supports "5".
excludeDrugsFromCovariates	DEPRECATED: Should the target and comparator drugs (and their descendant concepts) be excluded from the covariates? Note that this will work if the drugs are actually drug concept IDs (and not cohort IDs).
firstExposureOnly	Should only the first exposure per subject be included? Note that this is typically done in the <a href="#">createStudyPopulation()</a> function, but can already be done here for efficiency reasons.
removeDuplicateSubjects	Remove subjects that are in both the target and comparator cohort? See details for allowed values. Note that this is typically done in the <a href="#">createStudyPopulation</a> function, but can already be done here for efficiency reasons.
restrictToCommonPeriod	Restrict the analysis to the period when both treatments are observed?
washoutPeriod	The minimum required continuous observation time prior to index date for a person to be included in the cohort. Note that this is typically done in the

	createStudyPopulation function, but can already be done here for efficiency reasons.
maxCohortSize	If either the target or the comparator cohort is larger than this number it will be sampled to this size. maxCohortSize = 0 indicates no maximum size.
covariateSettings	An object of type covariateSettings as created using the <a href="#">FeatureExtraction::createCovariate</a> function.

## Details

Based on the arguments, the treatment and comparator cohorts are retrieved, as well as outcomes occurring in exposed subjects. The treatment and comparator cohorts can be identified using the DRUG\_ERA table, or through user-defined cohorts in a cohort table either inside the CDM schema or in a separate schema. Similarly, outcomes are identified using the CONDITION\_ERA table or through user-defined cohorts in a cohort table either inside the CDM schema or in a separate schema. Covariates are automatically extracted from the appropriate tables within the CDM.

**Important:** The target and comparator drug must not be included in the covariates, including any descendant concepts. You will need to manually add the drugs and descendants to the excludedCovariateConceptIds of the covariateSettings argument.

The removeduplicateSubjects argument can have one of the following values:

- "keep all": Do not remove subjects that appear in both target and comparator cohort
- "keep first": When a subjects appear in both target and comparator cohort, only keep whichever cohort is first in time.
- "remove all": Remove subjects that appear in both target and comparator cohort completely from the analysis."

## Value

A [CohortMethodData](#) object.

---

```
getDefaultCmTable1Specifications
```

*Get the default table 1 specifications*

---

## Description

Loads the default specifications for a table 1, to be used with the [createTable1](#) function.

Important: currently only works for binary covariates.

## Usage

```
getDefaultCmTable1Specifications()
```

## Value

A specifications objects.

---

```
getFollowUpDistribution
```

*Get the distribution of follow-up time*

---

### Description

Get the distribution of follow-up time

### Usage

```
getFollowUpDistribution(population, quantiles = c(0, 0.25, 0.5, 0.75, 1))
```

### Arguments

population	A data frame describing the study population as created using the <a href="#">createStudyPopulation</a> function. This should at least have these columns: treatment, timeAtRisk.
quantiles	The quantiles of the population to compute minimum follow-up time for.

### Details

Get the distribution of follow-up time as quantiles. Follow-up time is defined as time-at-risk, so not censored at the outcome.

### Value

A data frame with per treatment group at each quantile the amount of follow-up time available.

---

```
getOutcomeModel
```

*Get the outcome model*

---

### Description

Get the full outcome model, so showing the betas of all variables included in the outcome model, not just the treatment variable.

### Usage

```
getOutcomeModel(outcomeModel, cohortMethodData)
```

### Arguments

outcomeModel	An object of type OutcomeModel as generated using the <a href="#">fitOutcomeModel()</a> function.
cohortMethodData	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .

### Value

A tibble.

---

getPsModel	<i>Get the propensity model</i>
------------	---------------------------------

---

**Description**

Returns the coefficients and names of the covariates with non-zero coefficients.

**Usage**

```
getPsModel(propensityScore, cohortMethodData)
```

**Arguments**

propensityScore

The propensity scores as generated using the [createPs\(\)](#) function.

cohortMethodData

An object of type [CohortMethodData](#) as generated using [getDbCohortMethodData\(\)](#).

**Value**

A tibble.

---

isCohortMethodData	<i>Check whether an object is a CohortMethodData object</i>
--------------------	---

---

**Description**

Check whether an object is a CohortMethodData object

**Usage**

```
isCohortMethodData(x)
```

**Arguments**

x

The object to check.

**Value**

A logical value.

---

loadCmAnalysisList	<i>Load a list of cmAnalysis from file</i>
--------------------	--

---

**Description**

Load a list of objects of type cmAnalysis from file. The file is in JSON format.

**Usage**

```
loadCmAnalysisList(file)
```

**Arguments**

file	The name of the file
------	----------------------

**Value**

A list of objects of type cmAnalysis.

---

loadCohortMethodData	<i>Load the cohort method data from a file</i>
----------------------	--

---

**Description**

Loads an object of type [CohortMethodData](#) from a file in the file system.

**Usage**

```
loadCohortMethodData(file)
```

**Arguments**

file	The name of the file containing the data.
------	---

**Value**

An object of class [CohortMethodData](#).



---

loadTargetComparatorOutcomesList

*Load a list of targetComparatorOutcomes from file*


---

**Description**

Load a list of objects of type targetComparatorOutcomes from file. The file is in JSON format.

**Usage**

```
loadTargetComparatorOutcomesList(file)
```

**Arguments**

file	The name of the file
------	----------------------

**Value**

A list of objects of type targetComparatorOutcomes.

---

matchOnPs

*Match persons by propensity score*


---

**Description**

Use the provided propensity scores to match target to comparator persons.

**Usage**

```
matchOnPs(
  population,
  caliper = 0.2,
  caliperScale = "standardized logit",
  maxRatio = 1,
  allowReverseMatch = FALSE,
  stratificationColumns = c()
)
```

**Arguments**

population	A data frame with the three columns described below.
caliper	The caliper for matching. A caliper is the distance which is acceptable for any match. Observations which are outside of the caliper are dropped. A caliper of 0 means no caliper is used.
caliperScale	The scale on which the caliper is defined. Three scales are supported: caliperScale = 'propensity score', caliperScale = 'standardized', or caliperScale = 'standardized logit'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution. 'standardized logit' is similar, except that the propensity score is transformed to the logit scale because the PS is more likely to be normally distributed on that scale (Austin, 2011).

<code>maxRatio</code>	The maximum number of persons in the comparator arm to be matched to each person in the treatment arm. A <code>maxRatio</code> of 0 means no maximum: all comparators will be assigned to a target person.
<code>allowReverseMatch</code>	Allows n-to-1 matching if target arm is larger
<code>stratificationColumns</code>	Names or numbers of one or more columns in the data <code>data.frame</code> on which subjects should be stratified prior to matching. No persons will be matched with persons outside of the strata identified by the values in these columns.

## Details

The data frame should have the following three columns:

- `rowId` (numeric): A unique identifier for each row (e.g. the person ID).
- `treatment` (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- `propensityScore` (numeric): Propensity score.

The default caliper (0.2 on the standardized logit scale) is the one recommended by Austin (2011).

## Value

Returns a data frame with the same columns as the input data plus one extra column: `stratumId`. Any rows that could not be matched are removed

## References

Rassen JA, Shelat AA, Myers J, Glynn RJ, Rothman KJ, Schneeweiss S. (2012) One-to-many propensity score matching in cohort studies, *Pharmacoepidemiology and Drug Safety*, May, 21 Suppl 2:69-80.

Austin, PC. (2011) Optimal caliper widths for propensity-score matching when estimating differences in means and differences in proportions in observational studies, *Pharmaceutical statistics*, March, 10(2):150-161.

## Examples

```
rowId <- 1:5
treatment <- c(1, 0, 1, 0, 1)
propensityScore <- c(0, 0.1, 0.3, 0.4, 1)
age_group <- c(1, 1, 1, 1, 1)
data <- data.frame(rowId = rowId,
                   treatment = treatment,
                   propensityScore = propensityScore,
                   age_group = age_group)
result <- matchOnPs(data, caliper = 0, maxRatio = 1, stratificationColumns = "age_group")
```

---

matchOnPsAndCovariates

*Match by propensity score as well as other covariates*


---

## Description

Use the provided propensity scores and a set of covariates to match target to comparator persons.

## Usage

```
matchOnPsAndCovariates(
  population,
  caliper = 0.2,
  caliperScale = "standardized logit",
  maxRatio = 1,
  allowReverseMatch = FALSE,
  cohortMethodData,
  covariateIds
)
```

## Arguments

population	A data frame with the three columns described below.
caliper	The caliper for matching. A caliper is the distance which is acceptable for any match. Observations which are outside of the caliper are dropped. A caliper of 0 means no caliper is used.
caliperScale	The scale on which the caliper is defined. Three scales are supported: caliperScale = 'propensity score', caliperScale = 'standardized', or caliperScale = 'standardized logit'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution. 'standardized logit' is similar, except that the propensity score is transformed to the logit scale because the PS is more likely to be normally distributed on that scale (Austin, 2011).
maxRatio	The maximum number of persons in the comparator arm to be matched to each person in the treatment arm. A maxRatio of 0 means no maximum: all comparators will be assigned to a target person.
allowReverseMatch	Allows n-to-1 matching if target arm is larger
cohortMethodData	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .
covariateIds	One or more covariate IDs in the cohortMethodData object on which subjects should be also matched.

## Details

The data frame should have the following three columns:

- rowId (numeric): A unique identifier for each row (e.g. the person ID).
- treatment (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- propensityScore (numeric): Propensity score.

The default caliper (0.2 on the standardized logit scale) is the one recommended by Austin (2011).

**Value**

Returns a tibble with the same columns as the input data plus one extra column: stratumId. Any rows that could not be matched are removed

**References**

Rassen JA, Shelat AA, Myers J, Glynn RJ, Rothman KJ, Schneeweiss S. (2012) One-to-many propensity score matching in cohort studies, *Pharmacoepidemiology and Drug Safety*, May, 21 Suppl 2:69-80.

Austin, PC. (2011) Optimal caliper widths for propensity-score matching when estimating differences in means and differences in proportions in observational studies, *Pharmaceutical statistics*, March, 10(2):150-161.

---

plotCovariateBalanceOfTopVariables

*Plot variables with largest imbalance*

---

**Description**

Create a plot showing those variables having the largest imbalance before matching, and those variables having the largest imbalance after matching. Requires running computeCovariateBalance first.

**Usage**

```
plotCovariateBalanceOfTopVariables(
  balance,
  n = 20,
  maxNameWidth = 100,
  title = NULL,
  fileName = NULL,
  beforeLabel = "before matching",
  afterLabel = "after matching"
)
```

**Arguments**

balance	A data frame created by the computeCovariateBalance function.
n	(Maximum) count of covariates to plot.
maxNameWidth	Covariate names longer than this number of characters are truncated to create a nicer plot.
title	Optional: the main title for the plot.
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function ggsave in the ggplot2 package for supported file formats.
beforeLabel	Label for identifying data before matching / stratification / trimming.
afterLabel	Label for identifying data after matching / stratification / trimming.

**Value**

A ggplot object. Use the [ggplot2::ggsave](#) function to save to file in a different format.

---

plotCovariateBalanceScatterPlot

*Create a scatterplot of the covariate balance*


---

## Description

Create a scatterplot of the covariate balance, showing all variables with balance before and after matching on the x and y axis respectively. Requires running `computeCovariateBalance` first.

## Usage

```
plotCovariateBalanceScatterPlot(
  balance,
  absolute = TRUE,
  threshold = 0,
  title = "Standardized difference of mean",
  fileName = NULL,
  beforeLabel = "Before matching",
  afterLabel = "After matching",
  showCovariateCountLabel = FALSE,
  showMaxLabel = FALSE
)
```

## Arguments

<code>balance</code>	A data frame created by the <code>computeCovariateBalance</code> function.
<code>absolute</code>	Should the absolute value of the difference be used?
<code>threshold</code>	Show a threshold value for after matching standardized difference.
<code>title</code>	The main title for the plot.
<code>fileName</code>	Name of the file where the plot should be saved, for example 'plot.png'. See the function <code>ggsave</code> in the <code>ggplot2</code> package for supported file formats.
<code>beforeLabel</code>	Label for the x-axis.
<code>afterLabel</code>	Label for the y-axis.
<code>showCovariateCountLabel</code>	Show a label with the number of covariates included in the plot?
<code>showMaxLabel</code>	Show a label with the maximum absolute standardized difference after matching/stratification?

## Value

A ggplot object. Use the [ggplot2::ggsave](#) function to save to file in a different format.

---

`plotCovariatePrevalence`*Plot covariate prevalence*

---

### Description

Plot prevalence of binary covariates in the target and comparator cohorts, before and after matching. Requires running `computeCovariateBalance` first.

### Usage

```
plotCovariatePrevalence(  
  balance,  
  threshold = 0,  
  title = "Covariate prevalence",  
  fileName = NULL,  
  beforeLabel = "Before matching",  
  afterLabel = "After matching",  
  targetLabel = "Target",  
  comparatorLabel = "Comparator"  
)
```

### Arguments

<code>balance</code>	A data frame created by the <code>computeCovariateBalance</code> function.
<code>threshold</code>	A threshold value for standardized difference. When exceeding the threshold, covariates will be marked in a different color. If <code>threshold = 0</code> , no color coding will be used.
<code>title</code>	The main title for the plot.
<code>fileName</code>	Name of the file where the plot should be saved, for example 'plot.png'. See the function <code>ggsave</code> in the <code>ggplot2</code> package for supported file formats.
<code>beforeLabel</code>	Label for the before matching / stratification panel.
<code>afterLabel</code>	Label for the after matching / stratification panel.
<code>targetLabel</code>	Label for the x-axis.
<code>comparatorLabel</code>	Label for the y-axis.

### Value

A ggplot object. Use the [ggplot2::ggsave](#) function to save to file in a different format.

---

`plotFollowUpDistribution`*Plot the distribution of follow-up time*

---

## Description

Plot the distribution of follow-up time

## Usage

```
plotFollowUpDistribution(  
  population,  
  targetLabel = "Target",  
  comparatorLabel = "Comparator",  
  yScale = "percent",  
  logYScale = FALSE,  
  dataCutoff = 0.95,  
  title = NULL,  
  fileName = NULL  
)
```

## Arguments

population	A data frame describing the study population as created using the <a href="#">createStudyPopulation</a> function. This should at least have these columns: treatment, timeAtRisk.
targetLabel	A label to us for the target cohort.
comparatorLabel	A label to us for the comparator cohort.
yScale	Should be either 'percent' or 'count'.
logYScale	Should the Y axis be on the log scale?
dataCutoff	Fraction of the data (number censored) after which the graph will not be shown.
title	The main title of the plot.
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function <code>ggsave</code> in the <code>ggplot2</code> package for supported file formats.

## Details

Plot the distribution of follow-up time, stratified by treatment group. Follow-up time is defined as time-at-risk, so not censored at the outcome.

## Value

A ggplot object. Use the [ggsave](#) function to save to file in a different format.

---

plotKaplanMeier

*Plot the Kaplan-Meier curve*


---

## Description

plotKaplanMeier creates the Kaplan-Meier (KM) survival plot. Based (partially) on recommendations in Pocock et al (2002).

When variable-sized strata are detected, an adjusted KM plot is computed to account for stratified data, as described in Galimberti et al (2002), using the closed form variance estimator described in Xie et al (2005).

## Usage

```
plotKaplanMeier(
  population,
  censorMarks = FALSE,
  confidenceIntervals = TRUE,
  includeZero = FALSE,
  dataTable = TRUE,
  dataCutoff = 0.9,
  targetLabel = "Treated",
  comparatorLabel = "Comparator",
  title,
  fileName = NULL
)
```

## Arguments

population	A population object generated by createStudyPopulation, potentially filtered by other functions.
censorMarks	Whether or not to include censor marks in the plot.
confidenceIntervals	Plot 95 percent confidence intervals? Default is TRUE, as recommended by Pocock et al.
includeZero	Should the y axis include zero, or only go down to the lowest observed survival? The default is FALSE, as recommended by Pocock et al.
dataTable	Should the numbers at risk be shown in a table? Default is TRUE, as recommended by Pocock et al.
dataCutoff	Fraction of the data (number censored) after which the graph will not be shown. The default is 90 percent as recommended by Pocock et al.
targetLabel	A label to us for the target cohort.
comparatorLabel	A label to us for the comparator cohort.
title	The main title of the plot.
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function ggsave in the ggplot2 package for supported file formats.



**Value**

A ggplot object. Use the [ggsave](#) function to save to file in a different format.

**References**

Pocock SJ, Clayton TC, Altman DG. (2002) Survival plots of time-to-event outcomes in clinical trials: good practice and pitfalls, *Lancet*, 359:1686-89.

Galimberti S, Sasieni P, Valsecchi MG (2002) A weighted Kaplan-Meier estimator for matched data with application to the comparison of chemotherapy and bone-marrow transplant in leukaemia. *Statistics in Medicine*, 21(24):3847-64.

Xie J, Liu C. (2005) Adjusted Kaplan-Meier estimator and log-rank test with inverse probability of treatment weighting for survival data. *Statistics in Medicine*, 26(10):2276.

---

plotPs	<i>Plot the propensity score distribution</i>
--------	---

---

**Description**

Plots the propensity (or preference) score distribution.

**Usage**

```
plotPs(
  data,
  unfilteredData = NULL,
  scale = "preference",
  type = "density",
  binWidth = 0.05,
  targetLabel = "Target",
  comparatorLabel = "Comparator",
  showCountsLabel = FALSE,
  showAucLabel = FALSE,
  showEquiposeLabel = FALSE,
  equipoiseBounds = c(0.3, 0.7),
  unitOfAnalysis = "subjects",
  title = NULL,
  fileName = NULL
)
```

**Arguments**

data	A data frame with at least the two columns described below
unfilteredData	To be used when computing preference scores on data from which subjects have already been removed, e.g. through trimming and/or matching. This data frame should have the same structure as data.
scale	The scale of the graph. Two scales are supported: scale = 'propensity' or scale = 'preference'. The preference score scale is defined by Walker et al (2013).

type	Type of plot. Four possible values: type = 'density' type = 'histogram', type = 'histogramCount', or type = 'histogramProportion'. 'histogram' defaults to 'histogramCount'.
binWidth	For histograms, the width of the bins
targetLabel	A label to us for the target cohort.
comparatorLabel	A label to us for the comparator cohort.
showCountsLabel	Show subject counts?
showAucLabel	Show the AUC?
showEquiposeLabel	Show the percentage of the population in equipoise?
equipoiseBounds	The bounds on the preference score to determine whether a subject is in equipoise.
unitOfAnalysis	The unit of analysis in the input data. Defaults to 'subjects'.
title	Optional: the main title for the plot.
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function <a href="#">ggplot2::ggsave()</a> for supported file formats.

## Details

The data frame should have a least the following two columns:

- treatment (integer): Column indicating whether the person is in the target (1) or comparator (0) group
- propensityScore (numeric): Propensity score

## Value

A ggplot object. Use the [ggplot2::ggsave\(\)](#) function to save to file in a different format.

## References

Walker AM, Patrick AR, Lauer MS, Hornbrook MC, Marin MG, Platt R, Roger VL, Stang P, and Schneeweiss S. (2013) A tool for assessing the feasibility of comparative effectiveness research, Comparative Effective Research, 3, 11-20

## Examples

```
treatment <- rep(0:1, each = 100)
propensityScore <- c(rnorm(100, mean = 0.4, sd = 0.25), rnorm(100, mean = 0.6, sd = 0.25))
data <- data.frame(treatment = treatment, propensityScore = propensityScore)
data <- data[data$propensityScore > 0 & data$propensityScore < 1, ]
plotPs(data)
```

---

plotTimeToEvent	<i>Plot time-to-event</i>
-----------------	---------------------------

---

## Description

Plot time-to-event

## Usage

```
plotTimeToEvent(
  cohortMethodData,
  population = NULL,
  outcomeId,
  firstExposureOnly = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  removeDuplicateSubjects = FALSE,
  minDaysAtRisk = 1,
  riskWindowStart = 0,
  addExposureDaysToStart = NULL,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  addExposureDaysToEnd = NULL,
  endAnchor = "cohort end",
  censorAtNewRiskWindow = FALSE,
  periodLength = 7,
  numberOfPeriods = 52,
  highlightExposedEvents = TRUE,
  includePostIndexTime = TRUE,
  showFittedLines = TRUE,
  targetLabel = "Target",
  comparatorLabel = "Comparator",
  title = NULL,
  fileName = NULL
)
```

## Arguments

cohortMethodData	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .
population	If specified, this population will be used as the starting point instead of the cohorts in the cohortMethodData object.
outcomeId	The ID of the outcome. If not specified, no outcome-specific transformations will be performed.
firstExposureOnly	(logical) Should only the first exposure per subject be included?
restrictToCommonPeriod	(logical) Restrict the analysis to the period when both exposures are observed?
washoutPeriod	The minimum required continuous observation time prior to index date for a person to be included in the cohort.

<code>removeDuplicateSubjects</code>	Remove subjects that are in both the target and comparator cohort? See details for allowed values.
<code>minDaysAtRisk</code>	The minimum required number of days at risk.
<code>riskWindowStart</code>	The start of the risk window (in days) relative to the <code>startAnchor</code> .
<code>addExposureDaysToStart</code>	DEPRECATED: Add the length of exposure the start of the risk window? Use <code>startAnchor</code> instead.
<code>startAnchor</code>	The anchor point for the start of the risk window. Can be "cohort start" or "cohort end".
<code>riskWindowEnd</code>	The end of the risk window (in days) relative to the <code>endAnchor</code> .
<code>addExposureDaysToEnd</code>	DEPRECATED: Add the length of exposure the risk window? Use <code>endAnchor</code> instead.
<code>endAnchor</code>	The anchor point for the end of the risk window. Can be "cohort start" or "cohort end".
<code>censorAtNewRiskWindow</code>	If a subject is in multiple cohorts, should time-at-risk be censored when the new time-at-risk starts to prevent overlap?
<code>periodLength</code>	The length in days of each period shown in the plot.
<code>numberOfPeriods</code>	Number of periods to show in the plot. The periods are equally divided before and after the index date.
<code>highlightExposedEvents</code>	(logical) Highlight event counts during exposure in a different color?
<code>includePostIndexTime</code>	(logical) Show time after the index date?
<code>showFittedLines</code>	(logical) Fit lines to the proportions and show them in the plot?
<code>targetLabel</code>	A label to us for the target cohort.
<code>comparatorLabel</code>	A label to us for the comparator cohort.
<code>title</code>	Optional: the main title for the plot.
<code>fileName</code>	Name of the file where the plot should be saved, for example 'plot.png'. See <a href="#">ggplot2::ggsave()</a> for supported file formats.

## Details

Creates a plot showing the number of events over time in the target and comparator cohorts, both before and after index date. The plot also distinguishes between events inside and outside the time-at-risk period. This requires the user to (re)specify the time-at-risk using the same arguments as the [createStudyPopulation\(\)](#) function. Note that it is not possible to specify that people with the outcome prior should be removed, since the plot will show these prior events.

## Value

A ggplot object. Use the [ggplot2::ggsave\(\)](#) function to save to file in a different format.

---

runCmAnalyses	<i>Run a list of analyses</i>
---------------	-------------------------------

---

## Description

Run a list of analyses

## Usage

```
runCmAnalyses(
  connectionDetails,
  cdmDatabaseSchema,
  oracleTempSchema = NULL,
  tempEmulationSchema = getOption("sqlRenderTempEmulationSchema"),
  exposureDatabaseSchema = cdmDatabaseSchema,
  exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence",
  cdmVersion = 5,
  outputFolder = "../CohortMethodOutput",
  cmAnalysisList,
  targetComparatorOutcomesList,
  refitPsForEveryOutcome = FALSE,
  refitPsForEveryStudyPopulation = TRUE,
  prefilterCovariates = TRUE,
  getDbCohortMethodDataThreads = 1,
  createPsThreads = 1,
  psCvThreads = 1,
  createStudyPopThreads = 1,
  trimMatchStratifyThreads = 1,
  prefilterCovariatesThreads = 1,
  fitOutcomeModelThreads = 1,
  outcomeCvThreads = 1,
  outcomeIdsOfInterest,
  analysesToExclude = NULL
)
```

## Arguments

connectionDetails

An R object of type connectionDetails created using the [DatabaseConnector::createConnection](#) function.

cdmDatabaseSchema

The name of the database schema that contains the OMOP CDM instance. Requires read permissions to this database. On SQL Server, this should specify both the database and the schema, so for example 'cdm\_instance.dbo'.

oracleTempSchema

DEPRECATED: use tempEmulationSchema instead.

tempEmulationSchema

Some database platforms like Oracle and Impala do not truly support temp tables. To emulate temp tables, provide a schema with write privileges where temp tables can be created.

exposureDatabaseSchema	The name of the database schema that is the location where the exposure data used to define the exposure cohorts is available. If exposureTable = DRUG_ERA, exposureDatabaseSchema is not used by assumed to be cdmSchema. Requires read permissions to this database.
exposureTable	The tablename that contains the exposure cohorts. If exposureTable <> DRUG_ERA, then expectation is exposureTable has format of COHORT table: COHORT_DEFINITION_ID, SUBJECT_ID, COHORT_START_DATE, COHORT_END_DATE.
outcomeDatabaseSchema	The name of the database schema that is the location where the data used to define the outcome cohorts is available. If exposureTable = CONDITION_ERA, exposureDatabaseSchema is not used by assumed to be cdmSchema. Requires read permissions to this database.
outcomeTable	The tablename that contains the outcome cohorts. If outcomeTable <> CONDITION_OCCURRENCE, then expectation is outcomeTable has format of COHORT table: COHORT_DEFINITION_ID, SUBJECT_ID, COHORT_START_DATE, COHORT_END_DATE.
cdmVersion	Define the OMOP CDM version used: currently support "4" and "5".
outputFolder	Name of the folder where all the outputs will written to.
cmAnalysisList	A list of objects of type cmAnalysis as created using the <a href="#">createCmAnalysis</a> function.
targetComparatorOutcomesList	A list of objects of type targetComparatorOutcomes as created using the <a href="#">createTargetComparatorOutcomes</a> function.
refitPsForEveryOutcome	Should the propensity model be fitted for every outcome (i.e. after people who already had the outcome are removed)? If false, a single propensity model will be fitted, and people who had the outcome previously will be removed afterwards.
refitPsForEveryStudyPopulation	Should the propensity model be fitted for every study population definition? If false, a single propensity model will be fitted, and the study population criteria will be applied afterwards.
prefilterCovariates	If TRUE, and some outcome models require filtering covariates by concept ID (e.g. because includeCovariateIds or interactionCovariateIds is specified), this filtering will be done once for all outcome models that need it. This can greatly speed up the analyses if multiple outcome models require the same filtering.
getDbCohortMethodDataThreads	The number of parallel threads to use for building the cohortMethod data objects.
createPsThreads	The number of parallel threads to use for fitting the propensity models.
psCvThreads	The number of parallel threads to use for the cross-validation when estimating the hyperparameter for the propensity model. Note that the total number of CV threads at one time could be createPsThreads * psCvThreads.
createStudyPopThreads	The number of parallel threads to use for creating the study population.

`trimMatchStratifyThreads`  
 The number of parallel threads to use for trimming, matching and stratifying.

`prefilterCovariatesThreads`  
 The number of parallel threads to use for prefiltering covariates.

`fitOutcomeModelThreads`  
 The number of parallel threads to use for fitting the outcome models.

`outcomeCvThreads`  
 The number of parallel threads to use for the cross- validation when estimating the hyperparameter for the outcome model. Note that the total number of CV threads at one time could be `fitOutcomeModelThreads * outcomeCvThreads`.

`outcomeIdsOfInterest`  
 If provided, creation of non-essential files will be skipped for all other outcome IDs. This could be helpful to speed up analyses with many controls.

`analysesToExclude`  
 Analyses to exclude. See the Analyses to Exclude section for details.

## Details

Run a list of analyses for the target-comparator-outcomes of interest. This function will run all specified analyses against all hypotheses of interest, meaning that the total number of outcome models is `length(cmAnalysisList) * length(targetComparatorOutcomesList)` (if all analyses specify an outcome model should be fitted). When you provide several analyses it will determine whether any of the analyses have anything in common, and will take advantage of this fact. For example, if we specify several analyses that only differ in the way the outcome model is fitted, then this function will extract the data and fit the propensity model only once, and re-use this in all the analysis.

### Analyses to Exclude:

Normally, `runCmAnalyses` will run all combinations of target-comparator-outcome-analyses settings. However, sometimes we may not need all those combinations. Using the `analysesToExclude` argument, we can remove certain items from the full matrix. This argument should be a data frame with at least one of the following columns:

- `targetId`
- `comparatorId`
- `outcomeId`
- `analysisId`

This data frame will be joined to the outcome model reference table before executing, and matching rows will be removed. For example, if one specifies only one target ID and analysis ID, then any analyses with that target and that analysis ID will be skipped.

## Value

A tibble describing for each target-comparator-outcome-analysisId combination where the intermediary and outcome model files can be found, relative to the `outputFolder`.

---

saveCmAnalysisList	<i>Save a list of cmAnalysis to file</i>
--------------------	--

---

**Description**

Write a list of objects of type `cmAnalysis` to file. The file is in JSON format.

**Usage**

```
saveCmAnalysisList(cmAnalysisList, file)
```

**Arguments**

<code>cmAnalysisList</code>	The <code>cmAnalysis</code> list to be written to file
<code>file</code>	The name of the file where the results will be written

---

saveCohortMethodData	<i>Save the cohort method data to file</i>
----------------------	--

---

**Description**

Saves an object of type `CohortMethodData` to a file.

**Usage**

```
saveCohortMethodData(cohortMethodData, file)
```

**Arguments**

<code>cohortMethodData</code>	An object of type <code>CohortMethodData</code> as generated using <code>getDbCohortMethodData()</code> .
<code>file</code>	The name of the file where the data will be written. If the file already exists it will be overwritten.

**Value**

Returns no output.



---

```
saveTargetComparatorOutcomesList
```

*Save a list of targetComparatorOutcomes to file*

---

### Description

Write a list of objects of type targetComparatorOutcomes to file. The file is in JSON format.

### Usage

```
saveTargetComparatorOutcomesList(targetComparatorOutcomesList, file)
```

### Arguments

targetComparatorOutcomesList

The targetComparatorOutcomes list to be written to file

file

The name of the file where the results will be written

---

```
simulateCohortMethodData
```

*Generate simulated data*

---

### Description

Creates a [CohortMethodData](#) object with simulated data.

### Usage

```
simulateCohortMethodData(profile, n = 10000)
```

### Arguments

profile

An object of type CohortMethodDataSimulationProfile as generated using the [createCohortMethodDataSimulationProfile\(\)](#) function.

n

The size of the population to be generated.

### Details

This function generates simulated data that is in many ways similar to the original data on which the simulation profile is based. The contains same outcome, comparator, and outcome concept IDs, and the covariates and their 1st order statistics should be comparable.

### Value

An object of type [CohortMethodData](#).

---

stratifyByPs	<i>Stratify persons by propensity score</i>
--------------	---

---

## Description

Use the provided propensity scores to stratify persons. Additional stratification variables for stratifications can also be used.

## Usage

```
stratifyByPs(
  population,
  numberOfStrata = 5,
  stratificationColumns = c(),
  baseSelection = "all"
)
```

## Arguments

population	A data frame with the three columns described below
numberOfStrata	How many strata? The boundaries of the strata are automatically defined to contain equal numbers of target persons.
stratificationColumns	Names of one or more columns in the data data.frame on which subjects should also be stratified in addition to stratification on propensity score.
baseSelection	What is the base selection of subjects where the strata bounds are to be determined? Strata are defined as equally-sized strata inside this selection. Possible values are "all", "target", and "comparator".

## Details

The data frame should have the following three columns:

- rowId (numeric): A unique identifier for each row (e.g. the person ID).
- treatment (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- propensityScore (numeric): Propensity score.

## Value

Returns a tibble with the same columns as the input data plus one extra column: stratumId.

## Examples

```
rowId <- 1:200
treatment <- rep(0:1, each = 100)
propensityScore <- c(runif(100, min = 0, max = 1), runif(100, min = 0, max = 1))
data <- data.frame(rowId = rowId, treatment = treatment, propensityScore = propensityScore)
result <- stratifyByPs(data, 5)
```

---

`stratifyByPsAndCovariates`*Stratify persons by propensity score and other covariates*

---

## Description

Use the provided propensity scores and covariates to stratify persons.

## Usage

```
stratifyByPsAndCovariates(  
  population,  
  numberOfStrata = 5,  
  baseSelection = "all",  
  cohortMethodData,  
  covariateIds  
)
```

## Arguments

<code>population</code>	A data frame with the three columns described below
<code>numberOfStrata</code>	Into how many strata should the propensity score be divided? The boundaries of the strata are automatically defined to contain equal numbers of target persons.
<code>baseSelection</code>	What is the base selection of subjects where the strata bounds are to be determined? Strata are defined as equally-sized strata inside this selection. Possible values are "all", "target", and "comparator".
<code>cohortMethodData</code>	An object of type <a href="#">CohortMethodData</a> as generated using <a href="#">getDbCohortMethodData()</a> .
<code>covariateIds</code>	One or more covariate IDs in the <code>cohortMethodData</code> object on which subjects should also be stratified.

## Details

The data frame should have the following three columns:

- `rowId` (numeric): A unique identifier for each row (e.g. the person ID).
- `treatment` (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- `propensityScore` (numeric): Propensity score.

## Value

Returns a data frame with the same columns as the input population plus one extra column: `stratumId`.

---

summarizeAnalyses	<i>Create a summary report of the analyses</i>
-------------------	--

---

### Description

Create a summary report of the analyses

### Usage

```
summarizeAnalyses(referenceTable, outputFolder)
```

### Arguments

referenceTable A [dplyr::tibble](#) as created by the [runCmAnalyses](#) function.  
 outputFolder Name of the folder where all the outputs have been written to.

### Value

A tibble containing summary statistics for each target-comparator-outcome-analysis combination.

---

trimByIptw	<i>Remove subjects with a high IPTW</i>
------------	---

---

### Description

Compute the inverse probability of treatment weights (IPTW) using the propensity scores, and remove subjects having a weight higher than the user-specified threshold.

### Usage

```
trimByIptw(population, maxWeight = 10, estimator = "ate")
```

### Arguments

population A data frame with at least the three columns described below.  
 maxWeight The maximum allowed IPTW.  
 estimator The type of estimator. Options are estimator = "ate" for the average treatment effect, and estimator = "att" for the average treatment effect in the treated.

### Details

The data frame should have the following three columns:

- rowId (numeric): A unique identifier for each row (e.g. the person ID).
- treatment (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- propensityScore (numeric): Propensity score.

**Value**

Returns a tibble with the same columns as the input, as well as a weights column containing the IPTW.

**Examples**

```
rowId <- 1:2000
treatment <- rep(0:1, each = 1000)
propensityScore <- c(runif(1000, min = 0, max = 1), runif(1000, min = 0, max = 1))
data <- data.frame(rowId = rowId, treatment = treatment, propensityScore = propensityScore)
result <- trimByIptw(data)
```

---

trimByPs

*Trim persons by propensity score*


---

**Description**

Use the provided propensity scores to trim subjects with extreme scores.

**Usage**

```
trimByPs(population, trimFraction = 0.05)
```

**Arguments**

population	A data frame with the three columns described below
trimFraction	This fraction will be removed from each treatment group. In the target group, persons with the highest propensity scores will be removed, in the comparator group person with the lowest scores will be removed.

**Details**

The data frame should have the following three columns:

- rowId (numeric): A unique identifier for each row (e.g. the person ID).
- treatment (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- propensityScore (numeric): Propensity score.

**Value**

Returns a tibble with the same three columns as the input.

**Examples**

```
rowId <- 1:2000
treatment <- rep(0:1, each = 1000)
propensityScore <- c(runif(1000, min = 0, max = 1), runif(1000, min = 0, max = 1))
data <- data.frame(rowId = rowId, treatment = treatment, propensityScore = propensityScore)
result <- trimByPs(data, 0.05)
```

---

trimByPsToEquipoise	<i>Keep only persons in clinical equipoise</i>
---------------------	--

---

### Description

Use the preference score to trim subjects that are not in clinical equipoise

### Usage

```
trimByPsToEquipoise(population, bounds = c(0.3, 0.7))
```

### Arguments

population	A data frame with at least the three columns described below.
bounds	The upper and lower bound on the preference score for keeping persons.

### Details

The data frame should have the following three columns:

- `rowId` (numeric): A unique identifier for each row (e.g. the person ID).
- `treatment` (integer): Column indicating whether the person is in the target (1) or comparator (0) group.
- `propensityScore` (numeric): Propensity score.

### Value

Returns a tibble with the same three columns as the input.

### References

Walker AM, Patrick AR, Lauer MS, Hornbrook MC, Marin MG, Platt R, Roger VL, Stang P, and Schneeweiss S. (2013) A tool for assessing the feasibility of comparative effectiveness research, *Comparative Effective Research*, 3, 11-20

### Examples

```
rowId <- 1:2000
treatment <- rep(0:1, each = 1000)
propensityScore <- c(runif(1000, min = 0, max = 1), runif(1000, min = 0, max = 1))
data <- data.frame(rowId = rowId, treatment = treatment, propensityScore = propensityScore)
result <- trimByPsToEquipoise(data)
```

# Index

- \* **datasets**
  - cohortMethodDataSimulationProfile,  
5
- adjustedKm, 3
- Andromeda, 4
- checkCmInstallation, 4
- CohortMethodData, 5, 11, 18, 20, 25, 26,  
29–32, 35, 43, 48, 49, 51
- CohortMethodData  
(CohortMethodData-class), 4
- CohortMethodData-class, 4
- cohortMethodDataSimulationProfile, 5
- computeCovariateBalance, 5
- computeMdr, 6
- computePsAuc, 7
- CovariateData, 4
- createCmAnalysis, 8, 46
- createCmAnalysis(), 22
- createCmTable1, 10
- createCohortMethodDataSimulationProfile,  
11
- createCohortMethodDataSimulationProfile(),  
49
- createCreatePsArgs, 11
- createCreateStudyPopulationArgs, 12
- createFitOutcomeModelArgs, 14
- createGetDbCohortMethodDataArgs, 15
- createMatchOnPsAndCovariatesArgs, 16
- createMatchOnPsArgs, 17
- createPs, 18
- createPs(), 9, 31
- createStratifyByPsAndCovariatesArgs,  
19
- createStratifyByPsArgs, 19
- createStudyPopulation, 6, 20, 30, 39
- createStudyPopulation(), 8, 25, 26, 28, 44
- createTable1, 29
- createTargetComparatorOutcomes, 22, 46
- createTrimByIptwArgs, 23
- createTrimByPsArgs, 23
- createTrimByPsToEquipoiseArgs, 24
- Cyclops::createControl(), 18, 26
- Cyclops::createPrior(), 18, 26
- DatabaseConnector::createConnectionDetails(),  
27, 45
- dplyr::tibble, 52
- drawAttritionDiagram, 24
- FeatureExtraction::createCovariateSettings(),  
29
- fitOutcomeModel, 25
- fitOutcomeModel(), 9, 30
- getAttritionTable, 26
- getDbCohortMethodData, 27
- getDbCohortMethodData(), 4, 5, 8, 11, 18,  
20, 25, 30, 31, 35, 43, 48, 51
- getDefaultCmTable1Specifications, 29
- getFollowUpDistribution, 30
- getOutcomeModel, 30
- getPsModel, 31
- ggplot2::ggsave, 36–38
- ggplot2::ggsave(), 42, 44
- ggsave, 24, 39, 41
- isCohortMethodData, 31
- loadCmAnalysisList, 32
- loadCohortMethodData, 32
- loadCohortMethodData(), 4
- loadTargetComparatorOutcomesList, 33
- matchOnPs, 33
- matchOnPs(), 9
- matchOnPsAndCovariates, 35
- matchOnPsAndCovariates(), 9
- plotCovariateBalanceOfTopVariables, 36
- plotCovariateBalanceScatterPlot, 37
- plotCovariatePrevalence, 38
- plotFollowUpDistribution, 39
- plotKaplanMeier, 40
- plotPs, 41
- plotTimeToEvent, 43
- runCmAnalyses, 45, 52

`runCmAnalyses()`, [9](#), [23](#)

`saveCmAnalysisList`, [48](#)

`saveCohortMethodData`, [48](#)

`saveCohortMethodData()`, [4](#)

`saveTargetComparatorOutcomesList`, [49](#)

`show`, CohortMethodData-method  
(CohortMethodData-class), [4](#)

`simulateCohortMethodData`, [49](#)

`simulateCohortMethodData()`, [11](#)

`stratifyByPs`, [50](#)

`stratifyByPs()`, [9](#)

`stratifyByPsAndCovariates`, [51](#)

`stratifyByPsAndCovariates()`, [9](#)

`summarizeAnalyses`, [52](#)

`summary`, CohortMethodData-method  
(CohortMethodData-class), [4](#)

`trimByIptw`, [52](#)

`trimByIptw()`, [9](#)

`trimByPs`, [53](#)

`trimByPs()`, [9](#)

`trimByPsToEquipoise`, [54](#)

`trimByPsToEquipoise()`, [9](#)