

# Package ‘CohortMethod’

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**Type** Package

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

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**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 (>= 3.6.0),  
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 ( $\geq 3.0.1$ ),  
 bit64,  
 checkmate,  
 EmpiricalCalibration

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

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

**LinkingTo** Rcpp

**NeedsCompilation** yes

**RoxygenNote** 7.2.3

**Roxygen** list(markdown = TRUE)

**Encoding** UTF-8

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adjustedKm

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

Compute a weight-adjusted Kaplan-Meier curve

**Usage**

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

**Arguments**

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

---

<code>checkCmInstallation</code>	<i>Check is CohortMethod and its dependencies are correctly installed</i>
----------------------------------	---

---

**Description**

Check is CohortMethod and its dependencies are correctly installed

**Usage**

```
checkCmInstallation(connectionDetails)
```

**Arguments**

<code>connectionDetails</code>	An R object of type connectionDetails created using the function createConnectionDetails in the DatabaseConnector package.
--------------------------------	--

**Details**

This function checks whether CohortMethod and its dependencies are correctly installed. This will check the database connectivity, large scale regression engine (Cyclops), and large data object handling (ff).

---

CohortMethodData-class	<i>Cohort Method Data</i>
------------------------	---------------------------

---

**Description**

CohortMethodData is an S4 class that inherits from [CovariateData](#), which in turn inherits from [Andromeda](#). It contains information on the cohorts, their outcomes, and baseline covariates. Information about multiple outcomes can be captured at once for efficiency reasons.

A CohortMethodData is typically created using [getDbCohortMethodData\(\)](#), can only be saved using [saveCohortMethodData\(\)](#), and loaded using [loadCohortMethodData\(\)](#).

**Usage**

```
## S4 method for signature 'CohortMethodData'
show(object)

## S4 method for signature 'CohortMethodData'
summary(object)
```

**Arguments**

object                    An object of type CohortMethodData.

---

```
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 to the method described in Austin et al (2008).

**Usage**

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

## Arguments

- population** A data frame containing the people that are remaining after matching and/or trimming.
- cohortMethodData** An object of type `CohortMethodData` as generated using `getDbCohortMethodData()`.
- 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.
- covariateFilter** Determines the covariates for which to compute covariate balance. Either a vector of covariate IDs, or a table 1 specifications object as generated for example using `FeatureExtraction::getDefaultTable1Specifications()`. If `covariateFilter = NULL`, balance will be computed for all variables found in the data.

## 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.

---

computeEquipoise	<i>Compute fraction in equipoise</i>
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---

## Description

Compute fraction in equipoise

## Usage

```
computeEquipoise(data, equipoiseBounds = c(0.3, 0.7))
```

**Arguments**

data	A data frame with at least the two columns described below.
equipoiseBounds	The bounds on the preference score to determine whether a subject is in equipoise.

**Details**

Computes the fraction of the population (the union of the target and comparator cohorts) who are in clinical equipoise (i.e. who had a reasonable chance of receiving either target or comparator, based on the baseline characteristics).

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 numeric value (fraction in equipoise) between 0 and 1.

**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

---

computeMdr	<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**

population	A data frame describing the study population as created using the <a href="#">createStudyPopulation</a> function. This should at least have these columns: personSeqId, treatment, outcomeCount, timeAtRisk.
alpha	Type I error.
power	1 - beta, where beta is the type II error.

twoSided	Consider a two-sided test?
modelType	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, maxRows = 1e+05)
```

### 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)
maxRows	Maximum number of rows to use. If the number of rows is larger, a random sample will be taken. This can increase speed, with minor cost to precision. Set to 0 to use all data.

### 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 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
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---

## Description

Create a CohortMethod analysis specification

## Usage

```
createCmAnalysis(
  analysisId = 1,
  description = "",
  getDbCohortMethodDataArgs,
  createStudyPopArgs,
  createPsArgs = NULL,
  trimByPsArgs = NULL,
  trimByPsToEquipoiseArgs = NULL,
  trimByIptwArgs = NULL,
  truncateIptwArgs = NULL,
  matchOnPsArgs = NULL,
  matchOnPsAndCovariatesArgs = NULL,
  stratifyByPsArgs = NULL,
  stratifyByPsAndCovariatesArgs = NULL,
  computeSharedCovariateBalanceArgs = NULL,
  computeCovariateBalanceArgs = NULL,
  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.
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.
createPsArgs	An object representing the arguments to be used when calling the <a href="#">createPs()</a> function.
trimByPsArgs	An object representing the arguments to be used when calling the <a href="#">trimByPs()</a> function.

trimByPsToEquipoiseArgs	An object representing the arguments to be used when calling the <code>trimByPsToEquipoise()</code> function.
trimByIptwArgs	An object representing the arguments to be used when calling the <code>trimByIptw()</code> function.
truncateIptwArgs	An object representing the arguments to be used when calling the <code>truncateIptw()</code> function.
matchOnPsArgs	An object representing the arguments to be used when calling the <code>matchOnPs()</code> function.
matchOnPsAndCovariatesArgs	An object representing the arguments to be used when calling the <code>matchOnPsAndCovariates()</code> function.
stratifyByPsArgs	An object representing the arguments to be used when calling the <code>stratifyByPs()</code> function.
stratifyByPsAndCovariatesArgs	An object representing the arguments to be used when calling the <code>stratifyByPsAndCovariates()</code> function.
computeSharedCovariateBalanceArgs	An object representing the arguments to be used when calling the <code>computeCovariateBalance()</code> function per target-comparator-analysis.
computeCovariateBalanceArgs	An object representing the arguments to be used when calling the <code>computeCovariateBalance()</code> function per target-comparator-outcome-analysis.
fitOutcomeModelArgs	An object representing the arguments to be used when calling the <code>fitOutcomeModel()</code> function.

## Details

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

Providing a NULL value for any of the argument applies the corresponding step will not be executed. For example, if `createPsArgs = NULL`, no propensity scores will be computed.

---

createCmTable1	<i>Create a table 1</i>
----------------	-------------------------

---

## Description

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

## Usage

```
createCmTable1(
  balance,
  specifications = getDefaultCmTable1Specifications(),
  beforeTargetPopSize = NULL,
  beforeComparatorPopSize = NULL,
  afterTargetPopSize = NULL,
```

```

    afterComparatorPopSize = NULL,
    beforeLabel = "Before matching",
    afterLabel = "After matching",
    targetLabel = "Target",
    comparatorLabel = "Comparator",
    percentDigits = 1,
    stdDiffDigits = 2
  )

```

### Arguments

<code>balance</code>	A data frame created by the <code>computeCovariateBalance</code> function.
<code>specifications</code>	Specifications of which covariates to display, and how.
<code>beforeTargetPopSize</code>	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.
<code>beforeComparatorPopSize</code>	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.
<code>afterTargetPopSize</code>	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.
<code>afterComparatorPopSize</code>	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.
<code>beforeLabel</code>	Label for identifying columns before matching / stratification / trimming.
<code>afterLabel</code>	Label for identifying columns after matching / stratification / trimming.
<code>targetLabel</code>	Label for identifying columns of the target cohort.
<code>comparatorLabel</code>	Label for identifying columns of the comparator cohort.
<code>percentDigits</code>	Number of digits to be used for percentages.
<code>stdDiffDigits</code>	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.

---

```
createComputeCovariateBalanceArgs
```

*Create a parameter object for the function computeCovariateBalance*

---

**Description**

Create a parameter object for the function computeCovariateBalance

**Usage**

```
createComputeCovariateBalanceArgs(
  subgroupCovariateId = NULL,
  maxCohortSize = 250000,
  covariateFilter = NULL
)
```

**Arguments**

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 downsampled before computing covariate balance to save time. Setting this number to 0 means no downsampling will be applied.

covariateFilter

Determines the covariates for which to compute covariate balance. Either a vector of covariate IDs, or a table 1 specifications object as generated for example using `FeatureExtraction::getDefaultTable1Specifications()`. If covariateFilter = NULL, balance will be computed for all variables found in the data.

**Details**

Create an object defining the parameter values.

---

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,
    resetCoefficients = TRUE, tolerance = 2e-07, cvRepetitions = 10, startingVariance =
    0.01),
  estimator = "att"
)
```

## Arguments

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 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.
estimator	The type of estimator for the IPTW. Options are estimator = "ate" for the average treatment effect, estimator = "att" for the average treatment effect in the treated, and estimator = "ato" for the average treatment effect in the overlap population.

## 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 = "keep all",
  removeSubjectsWithPriorOutcome = TRUE,
  priorOutcomeLookback = 99999,
  minDaysAtRisk = 1,
  maxDaysAtRisk = 99999,
  riskWindowStart = 0,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  endAnchor = "cohort end",
  censorAtNewRiskWindow = FALSE
)
```

## Arguments

<code>firstExposureOnly</code>	Should only the first exposure per subject be included?
<code>restrictToCommonPeriod</code>	Restrict the analysis to the period when both exposures are observed?
<code>washoutPeriod</code>	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>removeSubjectsWithPriorOutcome</code>	Remove subjects that have the outcome prior to the risk window start?
<code>priorOutcomeLookback</code>	How many days should we look back when identifying prior outcomes?
<code>minDaysAtRisk</code>	The minimum required number of days at risk. Risk windows with fewer days than this number are removed from the analysis.
<code>maxDaysAtRisk</code>	The maximum allowed number of days at risk. Risk windows that are longer will be truncated to this number of days.
<code>riskWindowStart</code>	The start of the risk window (in days) relative to the startAnchor.
<code>startAnchor</code>	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.
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.

---

createDefaultMultiThreadingSettings
<i>Create default CohortMethod multi-threading settings</i>

---

## Description

Create CohortMethod multi-threading settings based on the maximum number of cores to be used.

## Usage

```
createDefaultMultiThreadingSettings(maxCores)
```

## Arguments

maxCores	Maximum number of CPU cores to use.
----------	-------------------------------------

## Value

An object of type CmMultiThreadingSettings.

## See Also

[createMultiThreadingSettings\(\)](#)

## Examples

```
settings <- createDefaultMultiThreadingSettings(10)
```

---

```
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,
  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, resetCoefficients = TRUE,
    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)
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 = "",
  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.

**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      *Create a parameter object for the function matchOnPs*

---

## 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.

---

```
createMultiThreadingSettings
```

*Create CohortMethod multi-threading settings*

---

## Description

Create CohortMethod multi-threading settings

## Usage

```
createMultiThreadingSettings(
  getDbCohortMethodDataThreads = 1,
  createPsThreads = 1,
  psCvThreads = 1,
  createStudyPopThreads = 1,
  trimMatchStratifyThreads = 1,
  computeSharedBalanceThreads = 1,
  computeBalanceThreads = 1,
  prefilterCovariatesThreads = 1,
  fitOutcomeModelThreads = 1,
  outcomeCvThreads = 1,
  calibrationThreads = 1
)
```

## Arguments

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.

computeSharedBalanceThreads

The number of parallel threads to use for computing shared covariate balance.

computeBalanceThreads

The number of parallel threads to use for computing covariate balance.

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.

calibrationThreads

The number of parallel threads to use for empirical calibration.

### Value

An object of type CmMultiThreadingSettings.

### See Also

[createDefaultMultiThreadingSettings\(\)](#)

---

createOutcome	<i>Create outcome definition</i>
---------------	----------------------------------

---

### Description

Create outcome definition

### Usage

```
createOutcome(
  outcomeId,
  outcomeOfInterest = TRUE,
  trueEffectSize = NA,
  priorOutcomeLookback = NULL,
  riskWindowStart = NULL,
  startAnchor = NULL,
  riskWindowEnd = NULL,
  endAnchor = NULL
)
```

### Arguments

outcomeId	An integer used to identify the outcome in the outcome cohort table.
outcomeOfInterest	Is this an outcome of interest? If not, creation of non-essential files will be skipped, including outcome=specific covariate balance files. This could be helpful to speed up analyses with many controls, for which we're only interested in the effect size estimate.
trueEffectSize	For negative and positive controls: the known true effect size. To be used for empirical calibration. Negative controls have trueEffectSize = 1. If the true effect size is unknown, use trueEffectSize = NA
priorOutcomeLookback	How many days should we look back when identifying prior. outcomes?
riskWindowStart	The start of the risk window (in days) relative to the startAnchor.
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.
endAnchor	The anchor point for the end of the risk window. Can be "cohort start" or "cohort end".

## Details

Any settings here that are not NULL will override any values set in `createCreateStudyPopulationArgs()`.

## Value

An object of type `outcome`, to be used in `createTargetComparatorOutcomes()`.

---

createPs	<i>Create propensity scores</i>
----------	---------------------------------

---

## Description

Creates propensity scores and inverse probability of treatment weights (IPTW) 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,
    resetCoefficients = TRUE, tolerance = 2e-07, cvRepetitions = 10, startingVariance =
    0.01),
  estimator = "att"
)
```

## Arguments

- `cohortMethodData`  
An object of type `CohortMethodData` as generated using `getDbCohortMethodData()`.
- `population`  
A data frame describing the population. This should at least have a `rowId` column corresponding to the `rowId` column in the `CohortMethodData` covariates object and a treatment column. If population is not specified, the full population in the `CohortMethodData` 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.
estimator	The type of estimator for the IPTW. Options are <code>estimator = "ate"</code> for the average treatment effect, <code>estimator = "att"</code> for the average treatment effect in the treated, and <code>estimator = "ato"</code> for the average treatment effect in the overlap population.

### 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).

### 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

### 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 = NULL,
  firstExposureOnly = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  removeDuplicateSubjects = "keep all",
  removeSubjectsWithPriorOutcome = TRUE,
  priorOutcomeLookback = 99999,
  minDaysAtRisk = 1,
  maxDaysAtRisk = 99999,
  riskWindowStart = 0,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  endAnchor = "cohort end",
  censorAtNewRiskWindow = FALSE
)
```

## 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 NULL, 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.
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.
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,
  outcomes,
  excludedCovariateConceptIds = c(),
  includedCovariateConceptIds = c()
)
```

**Arguments**

targetId	A cohort ID identifying the target exposure in the exposure table.
comparatorId	A cohort ID identifying the comparator exposure in the exposure table.
outcomes	A list of object of type outcome as created by <a href="#">createOutcome()</a> .
excludedCovariateConceptIds	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 target-comparator combination.
includedCovariateConceptIds	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 target-comparator combination.

**Details**

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

**Value**

An object of type targetComparatorOutcomes.

---

createTrimByIptwArgs    *Create a parameter object for the function trimByIptw*

---

**Description**

Create a parameter object for the function trimByIptw

**Usage**

```
createTrimByIptwArgs(maxWeight = 10)
```

**Arguments**

maxWeight	The maximum allowed IPTW.
-----------	---------------------------

**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**

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**

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.

---

```
createTruncateIptwArgs
```

*Create a parameter object for the function truncateIptw*

---

### Description

Create a parameter object for the function truncateIptw

### Usage

```
createTruncateIptwArgs(maxWeight = 10)
```

### Arguments

maxWeight      The maximum allowed IPTW.

### 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.

---

exportToCsv	<i>Export cohort method results to CSV files</i>
-------------	--

---

### Description

Export cohort method results to CSV files

### Usage

```
exportToCsv(
  outputFolder,
  exportFolder = file.path(outputFolder, "export"),
  databaseId = 1,
  minCellCount = 5,
  maxCores = 1
)
```

### Arguments

outputFolder	The folder where runCmAnalyses() generated all results.
exportFolder	The folder where the CSV files will written.
databaseId	A unique ID for the database. This will be appended to most tables.
minCellCount	To preserve privacy: the minimum number of subjects contributing to a count before it can be included in the results. If the count is below this threshold, it will be set to -minCellCount.
maxCores	How many parallel cores should be used?

### Details

This requires that `runCmAnalyses()` has been executed first. It exports all the results in the outputFolder to CSV files for sharing with other sites.

### Value

Does not return anything. Is called for the side-effect of populating the exportFolder with CSV files.

---

fitOutcomeModel	<i>Create an outcome model, and compute the relative risk</i>
-----------------	---

---

### Description

Create an outcome model, and computes the relative risk

**Usage**

```

fitOutcomeModel(
  population,
  cohortMethodData = NULL,
  modelType = "logistic",
  stratified = FALSE,
  useCovariates = FALSE,
  inversePtWeighting = FALSE,
  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, resetCoefficients = TRUE,
    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)
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**

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.

---

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 [CohortMethodData](#), a population object generated by functions like [createStudyPopulation\(\)](#), 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	<i>Get the cohort data from the server</i>
-----------------------	--

---

**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,
  tempEmulationSchema = getOption("sqlRenderTempEmulationSchema"),
  targetId,
  comparatorId,
  outcomeIds,
  studyStartDate = "",
  studyEndDate = "",
```



```

    exposureDatabaseSchema = cdmDatabaseSchema,
    exposureTable = "drug_era",
    outcomeDatabaseSchema = cdmDatabaseSchema,
    outcomeTable = "condition_occurrence",
    cdmVersion = "5",
    firstExposureOnly = FALSE,
    removeDuplicateSubjects = FALSE,
    restrictToCommonPeriod = FALSE,
    washoutPeriod = 0,
    maxCohortSize = 0,
    covariateSettings
  )

```

## Arguments

connectionDetails

An R object of type connectionDetails created using the `DatabaseConnector::createConnectionDetails` 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'.

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".
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 <a href="#">createStudyPopulation</a> 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."

If the covariateSettings include cohort-based covariates, and the covariateCohortTable is NULL, the covariateCohortDatabaseSchema and covariateCohortTable will be set to the exposureDatabaseSchema and exposureTable, respectively .

## 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.

---

```
getFileReference
```

*Get file reference*

---

**Description**

Get file reference

**Usage**

```
getFileReference(outputFolder)
```

**Arguments**

`outputFolder`    Name of the folder where all the outputs have been written to.

**Value**

A tibble containing file names of artifacts generated for each target-comparator-outcome-analysis combination.

---

`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.

---

`getInteractionResultsSummary`*Get a summary report of the analyses results*

---

**Description**

Get a summary report of the analyses results

**Usage**

```
getInteractionResultsSummary(outputFolder)
```

**Arguments**

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.

---

getOutcomeModel	<i>Get the outcome model</i>
-----------------	------------------------------

---

### 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 [fitOutcomeModel\(\)](#) function.

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

### 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.

---

getResultsSummary	<i>Get a summary report of the analyses results</i>
-------------------	---

---

**Description**

Get a summary report of the analyses results

**Usage**

```
getResultsSummary(outputFolder)
```

**Arguments**

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.

---

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).



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

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 <a href="#">ggsave</a> in the <a href="#">ggplot2</a> 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	<i>Plot the Kaplan-Meier curve</i>
-----------------	------------------------------------

---

## 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 = NULL,
  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 <code>ggplot2::ggsave()</code> 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 = NULL,
  firstExposureOnly = FALSE,
  restrictToCommonPeriod = FALSE,
  washoutPeriod = 0,
  removeDuplicateSubjects = "keep all",
  minDaysAtRisk = 1,
  riskWindowStart = 0,
  startAnchor = "cohort start",
  riskWindowEnd = 0,
  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 NULL, 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.

removeDuplicateSubjects	Remove subjects that are in both the target and comparator cohort? See details for allowed values.
minDaysAtRisk	The minimum required number of days at risk.
riskWindowStart	The start of the risk window (in days) relative to the startAnchor.
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.
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?
periodLength	The length in days of each period shown in the plot.
numberOfPeriods	Number of periods to show in the plot. The periods are equally divided before and after the index date.
highlightExposedEvents	(logical) Highlight event counts during exposure in a different color?
includePostIndexTime	(logical) Show time after the index date?
showFittedLines	(logical) Fit lines to the proportions and show them in the plot?
targetLabel	A label to us for the target cohort.
comparatorLabel	A label to us for the comparator cohort.
title	Optional: the main title for the plot.
fileName	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

*Run a list of analyses***Description**

Run a list of analyses

**Usage**

```
runCmAnalyses(
  connectionDetails,
  cdmDatabaseSchema,
  tempEmulationSchema = getOption("sqlRenderTempEmulationSchema"),
  exposureDatabaseSchema = cdmDatabaseSchema,
  exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence",
  cdmVersion = "5",
  outputFolder = "./CohortMethodOutput",
  cmAnalysisList,
  targetComparatorOutcomesList,
  analysesToExclude = NULL,
  refitPsForEveryOutcome = FALSE,
  refitPsForEveryStudyPopulation = TRUE,
  multiThreadingSettings = createMultiThreadingSettings()
)
```

**Arguments**

connectionDetails

An R object of type connectionDetails created using the `DatabaseConnector::createConnectionDetails` 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'.

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.
analysesToExclude	Analyses to exclude. See the Analyses to Exclude section for details.
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.
multiThreadingSettings	An object of type CmMultiThreadingSettings as created using the <a href="#">createMultiThreadingSettings</a> or <a href="#">createDefaultMultiThreadingSettings()</a> functions.

## 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.

After completion, a tibble containing references to all generated files can be obtained using the [getFileReference\(\)](#) function. A summary of the analysis results can be obtained using the [getResultsSummary\(\)](#) function.

### 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**

cmAnalysisList    The cmAnalysis list to be written to file  
file                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**

cohortMethodData    An object of type CohortMethodData as generated using getDbCohortMethodData().  
file                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`.

---

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

---

### Description

Remove subjects having a weight higher than the user-specified threshold.

### Usage

```
trimByIptw(population, maxWeight = 10)
```

### Arguments

population	A data frame with at least the two columns described in the details
maxWeight	The maximum allowed IPTW.

### Details

The data frame should have the following two columns:

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

### Value

Returns a tibble with the same columns as the input.

### Examples

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

---

trimByPs	<i>Trim persons by propensity score</i>
----------	---

---

### 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)
```

---

<code>trimByPsToEquipoise</code>	<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)
```

---

truncateIptw	<i>Truncate IPTW values</i>
--------------	-----------------------------

---

**Description**

Set the inverse probability of treatment weights (IPTW) to the user-specified threshold if it exceeds said threshold.

**Usage**

```
truncateIptw(population, maxWeight = 10)
```

**Arguments**

population	A data frame with at least the two columns described in the details
maxWeight	The maximum allowed IPTW.

**Details**

The data frame should have the following two columns:

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

**Value**

Returns a tibble with the same columns as the input.

**Examples**

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

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