

# Package ‘CohortMethod’

July 29, 2015

**Type** Package

**Title** New-user cohort method with large scale propensity and outcome models

**Version** 1.1.0

**Date** 2015-07-01

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**Description** CohortMethod is an R package for performing new-user cohort studies in an observational database in the OMOP Common Data Model. It extracts the necessary data from a database in OMOP Common Data Model format, and uses 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 and matching on 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.

**License** Apache License 2.0

**VignetteBuilder** knitr

**Depends** R (>= 3.1.0),  
DatabaseConnector (>= 1.3.0),  
Cyclops (>= 1.1.0),  
PatientLevelPrediction (>= 0.0.2)

**Imports** bit,  
ggplot2,  
ff,  
ffbase (>= 0.12.1),  
plyr,  
Rcpp (>= 0.11.2),  
RJDBC,  
SqlRender (>= 1.1.1),  
survival,  
rjson,  
OhdsiRTools

**Suggests** testthat,  
pROC,

gnm,  
knitr,  
rmarkdown,  
EmpiricalCalibration

**LinkingTo** Rcpp

**NeedsCompilation** yes

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CohortMethod	<i>CohortMethod</i>
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Description

CohortMethod

cohortMethodDataSimulationProfile	<i>A simulation profile</i>
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Description

A simulation profile

Usage

data(cohortMethodDataSimulationProfile)

computeCovariateBalance	<i>Compute covariate balance before and after matching and trimming</i>
-------------------------	---

Description

For every covariate, prevalence in treatment and comparator groups before and after matching/trimming are computed.

Usage

computeCovariateBalance(restrictedCohorts, cohortMethodData, outcomeId = NULL)

**Arguments**

restrictedCohorts	A data frame containing the people that are remaining after matching and/or trimming.
cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.
outcomeId	The concept ID of the outcome. Persons marked for removal for the outcome will be removed when computing the balance before matching/trimming.

**Details**

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

rowId	(integer)	A unique identifier for each row (e.g. the person ID)
treatment	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group

**Value**

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

---

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

---

**Description**

computePsAuc computes the area under the ROC curve of the propensity score

**Usage**

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

**Arguments**

data	A data frame with at least the two columns described below
confidenceIntervals	Compute 95 percent confidence intervals (computationally expensive for large data sets)

**Details**

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

treatment	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
propensityScore	(real)	Propensity score

**Value**

A data frame 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)
```

---

constructEras	<i>Build eras</i>
---------------	-------------------

---

**Description**

Constructs eras (continuous periods of exposure or disease).

**Usage**

```
constructEras(connectionDetails, sourceDatabaseSchema,
  sourceTable = "drug_exposure",
  targetDatabaseSchema = sourceDatabaseSchema, targetTable = "drug_era",
  createTargetTable = FALSE, cdmDatabaseSchema = sourceDatabaseSchema,
  gracePeriod = 30, rollUp = TRUE, rollUpConceptClassId = "Ingredient",
  rollUpVocabularyId = "RxNorm", cdmVersion = "5")
```

**Arguments**

connectionDetails

An R object of type connectionDetails created using the function createConnectionDetails in the DatabaseConnector package.

sourceDatabaseSchema

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

sourceTable      The name of the source table.

targetDatabaseSchema

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

targetTable      The name of the target table.

createTargetTable

Should the target table be created? If not, the data is inserted in an existing table.

cdmDatabaseSchema

Only needed when rolling up concepts to ancestors: The name of the database schema that contains the vocabulary files. Requires read permissions to this database. On SQL Server, this should specify both the database and the schema, so for example 'cdm\_instance.dbo'.

gracePeriod	The number of days allowed between periods for them to still be considered part of the same era.
rollUp	Should concepts be rolled up to their ancestors?
rollUpConceptClassId	The identifier of the concept class to which concepts should be rolled up.
rollUpVocabularyId	The identifier of the vocabulary to which concepts should be rolled up.
cdmVersion	The version of the CDM that is being used.

## Details

This function creates eras from source data. For example, one could use this function to create drug eras based on drug exposures. The function allows drugs to be rolled up to ingredients, and prescriptions to the same ingredient that overlap in time are merged into a single ingredient. Note that stockpiling is not assumed to take place (ie. overlap is discarded), but a grace period can be specified allowing for a small gap between prescriptions when merging. The user can specify the source and target table. These tables are assumed to have the same structure as the cohort table in the Common Data Model (CDM), except when the table names are 'drug\_exposure' or 'condition\_occurrence' for the source table, or 'drug\_era' or 'condition\_era' for the target table, in which case the tables are assumed to have the structure defined for those tables in the CDM. If both the source and target table specify a field for type\_concept\_id, the era construction will partition by the type\_concept\_id, in other words periods with different type\_concept\_ids will be treated independently.

## Examples

```
## Not run:
# Constructing drug eras in CDM v4:
constructEras(connectionDetails,
               sourceDatabaseSchema = cdmDatabaseSchema,
               sourceTable = "drug_exposure",
               targetTable = "drug_era",
               createTargetTable = FALSE,
               gracePeriod = 30,
               rollUpVocabularyId = 8,
               rollUpConceptClassId = "Ingredient",
               cdmVersion = "4")

# Constructing drug eras in CDM v5:
constructEras(connectionDetails,
               sourceDatabaseSchema = cdmDatabaseSchema,
               sourceTable = "drug_exposure",
               targetTable = "drug_era",
               createTargetTable = FALSE,
               gracePeriod = 30,
               rollUpVocabularyId = "RxNorm",
               rollUpConceptClassId = "Ingredient",
               cdmVersion = "5")

## End(Not run)
```

---

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

---

## Description

Create a CohortMethod analysis specification

## Usage

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

## Arguments

analysisId	An integer that will be used later to refer to this specific set of analysis choices.
description	A short description of the analysis.
comparatorType	If more than one comparator is provided for each drugComparatorOutcome, this field should be used to select the specific comparator to use in this analysis.
indicationType	If more than one indication is provided for each drugComparatorOutcome, this field should be used to select the specific indication to use in this analysis.
getDbCohortMethodDataArgs	An object representing the arguments to be used when calling the <a href="#">getDbCohortMethodData</a> function.
createPs	Should the <a href="#">createPs</a> function be used in this analysis?
createPsArgs	An object representing the arguments to be used when calling the <a href="#">createPs</a> function.
trimByPs	Should the <a href="#">trimByPs</a> function be used in this analysis?
trimByPsArgs	An object representing the arguments to be used when calling the <a href="#">trimByPs</a> function.
trimByPsToEquipoise	Should the <a href="#">trimByPsToEquipoise</a> function be used in this analysis?
trimByPsToEquipoiseArgs	An object representing the arguments to be used when calling the <a href="#">trimByPsToEquipoise</a> function.
matchOnPs	Should the <a href="#">matchOnPs</a> function be used in this analysis?
matchOnPsArgs	An object representing the arguments to be used when calling the <a href="#">matchOnPs</a> function.
matchOnPsAndCovariates	Should the <a href="#">matchOnPsAndCovariates</a> function be used in this analysis?

matchOnPsAndCovariatesArgs	An object representing the arguments to be used when calling the <a href="#">matchOnPsAndCovariates</a> function.
stratifyByPs	Should the <a href="#">stratifyByPs</a> function be used in this analysis?
stratifyByPsArgs	An object representing the arguments to be used when calling the <a href="#">stratifyByPs</a> function.
stratifyByPsAndCovariates	Should the <a href="#">stratifyByPsAndCovariates</a> function be used in this analysis?
stratifyByPsAndCovariatesArgs	An object representing the arguments to be used when calling the <a href="#">stratifyByPsAndCovariates</a> function.
computeCovariateBalance	Should the <a href="#">computeCovariateBalance</a> function be used in this analysis?
fitOutcomeModel	Should the <a href="#">fitOutcomeModel</a> function be used in this analysis?
fitOutcomeModelArgs	An object representing the arguments to be used when calling the <a href="#">fitOutcomeModel</a> function.

## Details

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

---

createCohortMethodDataSimulationProfile  
*Create simulation profile*

---

## Description

createCohortMethodDataSimulationProfile creates a profile based on the provided cohortMethodData object, which can be used to generate simulated data that has similar characteristics.

## Usage

```
createCohortMethodDataSimulationProfile(cohortMethodData)
```

## Arguments

cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.
------------------	--

## Details

The output of this function is an object that can be used by the simulateCohortMethodData function to generate a cohortMethodData object.

## Value

An object of type cohortDataSimulationProfile.



---

createCreatePsArgs	<i>Create a parameter object for the function createPs</i>
--------------------	--

---

### Description

Create a parameter object for the function createPs

### Usage

```
createCreatePsArgs(excludeCovariateIds = NULL,
  prior = createPrior("laplace", exclude = c(0), useCrossValidation = TRUE),
  control = createControl(noiseLevel = "silent", cvType = "auto",
    startingVariance = 0.1))
```

### Arguments

excludeCovariateIds	Exclude these covariates from the propensity model.
prior	The prior used to fit the model. See createPrior for details.
control	The control object used to control the cross-validation used to determine the hyperparameters of the prior (if applicable). See createControl for details.

### Details

Create an object defining the parameter values.

---

createDrugComparatorOutcomes	<i>Create drug-comparator-outcomes combinations.</i>
------------------------------	--

---

### Description

Create drug-comparator-outcomes combinations.

### Usage

```
createDrugComparatorOutcomes(targetId, comparatorId, outcomeIds,
  indicationConceptIds = c(), exclusionConceptIds = c(),
  excludedCovariateConceptIds = c(), includedCovariateConceptIds = c())
```

### Arguments

targetId	A concept ID indentifying the target drug in the exposure table. If multiple strategies for picking the target will be tested in the analysis, a named list of numbers can be provided instead. In the analysis, the name of the number to be used can be specified using the #' targetType parameter in the <a href="#">createCmAnalysis</a> function.
----------	---

comparatorId	A concept ID indentifying the comparator drug in the exposure table. If multiple strategies for picking the comparator will be tested in the analysis, a named list of numbers can be provided instead. In the analysis, the name of the number to be used can be specified using the #' comparatorType parameter in the <a href="#">createCmAnalysis</a> function.
outcomeIds	A vector of concept IDs indentifying the outcome(s) in the outcome table.
indicationConceptIds	A vector of concept IDs identifying conditions that are required to appear prior to or on the index date. If multiple strategies for picking the indication will be tested in the analysis, a named list of vectors can be provided instead. In the analysis, the name of the vector to be used can be specified using the indicationType parameter in the <a href="#">createCmAnalysis</a> function.
exclusionConceptIds	A list of concept IDs that cannot appear on or before the index date. This argument is to be used only for exclusion criteria that are specific to the drug-comparator combination.
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 drug-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 drug-comparator combination.

## Details

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

---

createFitOutcomeModelArgs

*Create a parameter object for the function fitOutcomeModel*

---

## Description

Create a parameter object for the function fitOutcomeModel

## Usage

```
createFitOutcomeModelArgs(stratifiedCox = TRUE, riskWindowStart = 0,
  riskWindowEnd = 9999, addExposureDaysToEnd = FALSE,
  useCovariates = TRUE, fitModel = TRUE, modelType = "cox",
  prior = createPrior("laplace", useCrossValidation = TRUE),
  control = createControl(cvType = "auto", startingVariance = 0.1,
  selectorType = "byPid", noiseLevel = "quiet"))
```

**Arguments**

stratifiedCox	Specifically for Cox regressions: specify whether to use the stratadefined in subPopulation in the analysis. For Poissonregression and logistic regression, this is implied in 'clr' and 'cpr'.
riskWindowStart	The start of the risk window (in days) relative to the index data.
riskWindowEnd	The end of the risk window (in days) relative to the index data (+days of exposure if the addExposureDaysToEnd parameter isspecified).
addExposureDaysToEnd	Add the length of exposure the risk window?
useCovariates	Whether to use the covariate matrix in the cohortMethodData in theoutcome model.
fitModel	If false, the model will not be fit, and only summary statistics areavailable.
modelType	The type of model to be fitted. See details for options.
prior	The prior used to fit the model. SeecreatePrior for details.
control	The control object used to control the cross-validation used todetermine the hyperparameters of the prior (if applicable). SeecreateControl 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(washoutWindow = 183,
  indicationLookbackWindow = 183, studyStartDate = "", studyEndDate = "",
  exclusionConceptIds = c(), outcomeConditionTypeConceptIds = c(),
  cdmVersion = "4", excludeDrugsFromCovariates = TRUE, covariateSettings)
```

**Arguments**

washoutWindow	The minimum required continuous observation time prior to indexdate for a person to be included in the cohort.
indicationLookbackWindow	NA
studyStartDate	A calendar date specifying the minimum date that a cohort indexdate can appear. Date format is 'yyyymmdd'.
studyEndDate	A calendar date specifying the maximum date that a cohort indexdate can appear. Date format is 'yyyymmdd'.

exclusionConceptIds	A list of CONCEPT_IDs used to restrict the cohorts, based on any descendant conditions/drugs/procedures occurring at least once anytime prior to the cohort index date.
outcomeConditionTypeConceptIds	A list of TYPE_CONCEPT_ID values that will restrict condition occurrences. Only applicable if outcomeTable = CONDITION_OCCURRENCE.
cdmVersion	Define the OMOP CDM version used: currently support "4" and "5".
excludeDrugsFromCovariates	Should the target and comparator drugs (and their descendant concepts) be excluded from the covariates? Note that this will work if the drugs are actually drug concept IDs (and not cohort IDs).
covariateSettings	An object of type covariateSettings as created using the createCovariateSettings function in the PatientLevelPrediction package..

## 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 = 0.25,
  caliperScale = "standardized", maxRatio = 1, 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. Two scales are supported: caliperScale = 'propensity score' or caliperScale = 'standardized'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution.
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 treated person.
covariateIds	One or more covariate IDs in the cohortMethodData object on which subjects should be also matched.

## Details

Create an object defining the parameter values.

---

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

---

## Description

Create a parameter object for the function matchOnPs

## Usage

```
createMatchOnPsArgs(caliper = 0.25, caliperScale = "standardized",
  maxRatio = 1, 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. Two scales are supported: caliperScale = 'propensity score' or caliperScale = 'standardized'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution.
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 treated person.
stratificationColumns	Names or numbers of one or more columns in the data data.frame on which subjects should be stratified prior to matching. No persons will be matched with persons outside of the strata identified by the values in these columns.

## Details

Create an object defining the parameter values.

---

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

---

## Description

createPs creates propensity scores using a regularized logistic regression.

## Usage

```
createPs(cohortMethodData, outcomeId = NULL, excludeCovariateIds = NULL,
  prior = createPrior("laplace", exclude = c(0), useCrossValidation = TRUE),
  control = createControl(noiseLevel = "silent", cvType = "auto",
    startingVariance = 0.1))
```

**Arguments**

cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.
outcomeId	The concept ID of the outcome. Persons marked for removal for the outcome will be removed prior to creating the propensity score model.
excludeCovariateIds	Exclude these covariates from the propensity model.
prior	The prior used to fit the model. See <a href="#">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="#">createControl</a> for details.

**Details**

createPs creates propensity scores using a regularized logistic regression.

**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, 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 treated persons.
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())
```

**Arguments**

`numberOfStrata` How many strata? The boundaries of the strata are automatically defined to contain equal numbers of treated 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.

**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 treatment 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(0.25, 0.75))
```

### Arguments

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

### Details

Create an object defining the parameter values.

---

```
drawAttritionDiagram    Draw the attrition diagram
```

---

### Description

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

### Usage

```
drawAttritionDiagram(outcomeModel, treatmentLabel = "Treated",
  comparatorLabel = "Comparator", fileName = NULL)
```

### Arguments

outcomeModel    An object of type outcomeModel as generated using the createOutcomeMode function.

treatmentLabel   A label to use for the treated cohort.

comparatorLabel   A label to use for the comparator cohort.

fileName          Name of the file where the plot should be saved, for example 'plot.png'. See the function ggsave in the ggplot2 package for supported file formats.

### Value

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



---

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

---

## Description

fitOutcomeModel creates an outcome model, and computes the relative risk

## Usage

```
fitOutcomeModel(outcomeId, cohortMethodData, subPopulation = NULL,
  stratifiedCox = TRUE, riskWindowStart = 0, riskWindowEnd = 9999,
  addExposureDaysToEnd = FALSE, useCovariates = TRUE, fitModel = TRUE,
  modelType = "cox", prior = createPrior("laplace", useCrossValidation =
    TRUE), control = createControl(cvType = "auto", startingVariance = 0.1,
  selectorType = "byPid", noiseLevel = "quiet"))
```

## Arguments

outcomeId	The concept ID of the outcome. Persons marked for removal for the outcome will be removed prior to creating the outcome model.
cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.
subPopulation	A data frame specifying the (matched and/or trimmed) subpopulation to be used in the study, as well as their strata (for conditional models). This data frame should have at least a RowId, and a StratumId when including stratification.
stratifiedCox	Specifically for Cox regressions: specify whether to use the strata defined in subPopulation in the analysis. For Poisson regression and logistic regression, this is implied in 'clr' and 'cpr'.
riskWindowStart	The start of the risk window (in days) relative to the index data.
riskWindowEnd	The end of the risk window (in days) relative to the index data (+ days of exposure if the addExposureDaysToEnd parameter is specified).
addExposureDaysToEnd	Add the length of exposure the risk window?
useCovariates	Whether to use the covariate matrix in the cohortMethodData in the outcome model.
fitModel	If false, the model will not be fit, and only summary statistics are available.
modelType	The type of model to be fitted. See details for options.
prior	The prior used to fit the model. See <a href="#">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="#">createControl</a> for details.

## Details

The model type can be one of these:

lr	Logistic regression
clr	Conditional logistic regression

cox Cox regression (stratified or not, depending on whether stata is specified)  
 pr Poisson regression  
 cpr Conditional Poisson regression

### Value

An object of class `outcomeModel`. Generic function `summary`, `coef`, and `confint` are available.

### Examples

```
# todo
```

---

```
getDbCohortMethodData Get the cohort data from the server
```

---

### Description

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

### Usage

```
getDbCohortMethodData(connectionDetails, cdmDatabaseSchema,
  oracleTempSchema = cdmDatabaseSchema, targetId, comparatorId,
  indicationConceptIds = c(), washoutWindow = 183,
  indicationLookbackWindow = 183, studyStartDate = "", studyEndDate = "",
  exclusionConceptIds = c(), outcomeIds,
  outcomeConditionTypeConceptIds = c(),
  exposureDatabaseSchema = cdmDatabaseSchema, exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence", cdmVersion = "4",
  excludeDrugsFromCovariates = TRUE, covariateSettings)
```

### Arguments

`connectionDetails`

An R object of type `connectionDetails` created using the function `createConnectionDetails` in the `DatabaseConnector` package.

`cdmDatabaseSchema`

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

`oracleTempSchema`

For Oracle only: the name of the database schema where you want all temporary tables to be managed. Requires create/insert permissions to this database.

`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_concept_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_concept_id in the cohort-like table.
indicationConceptIds	A list of CONCEPT_IDs used to restrict the target and comparator cohorts, based on any descendant condition of this list occurring at least once within the indicationLookbackWindow prior to the cohort index date. If no concept IDs are specified, the cohorts are not restricted to any indication.
washoutWindow	The minimum required continuous observation time prior to index date for a person to be included in the cohort.
indicationLookbackWindow	The window to look back prior to cohort index date to identify records of a indication condition. Only applicable if indicationConceptIds != "".
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'.
exclusionConceptIds	A list of CONCEPT_IDs used to restrict the cohorts, based on any descendant conditions/drugs/procedures occurring at least once anytime prior to the cohort index date.
outcomeConditionTypeConceptIds	A list of TYPE_CONCEPT_ID values that will restrict condition occurrences. Only applicable if outcomeTable = CONDITION_OCCURRENCE.
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_concept_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".
excludeDrugsFromCovariates	Should the target and comparator drugs (and their descendant concepts) be excluded from the covariates? Note that this will work if the drugs are actually drug concept IDs (and not cohort IDs).
covariateSettings	An object of type covariateSettings as created using the createCovariateSettings function in the PatientLevelPrediction package..

**outcomeConceptIds**

A list of CONCEPT\_IDs used to define outcomes. If outcomeTable = CONDITION\_OCCURRENCE, the list is a set of ancestor CONCEPT\_IDs, and all occurrences of all descendant concepts will be selected. If outcomeTable <> CONDITION\_OCCURRENCE, the list contains records found in COHORT\_DEFINITION\_ID field.

**Details**

Based on the parameters, the treatment and comparator cohorts are constructed. Baseline covariates at or before the index date are extracted, as well as outcomes occurring on or after the index date. 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 instance or in a separate schema. Similarly, outcomes are identified using the condition\_occurrence or condition\_era table, or through user-defined cohorts in a cohort table either inside the CDM instance or in a separate schema. Covariates are automatically extracted from the appropriate tables within the CDM. This function calls the getDbCovariates and getDbOutcomes functions. Important: The target and comparator drug must not be included in the covariates, including any descendant concepts. If the targetId and comparatorId arguments represent real concept IDs, you can set the excludeDrugsFromCovariates argument to TRUE and automatically the drugs and their descendants will be excluded from the covariates. However, if the targetId and comparatorId arguments do not represent concept IDs, you will need to manually add the drugs and descendants to the excludedCovariateConceptIds of the covariateSettings argument.

**Value**

Returns an object of type cohortMethodData, containing information on the cohorts, their outcomes, and baseline covariates. Information about multiple outcomes can be captured at once for efficiency reasons. This object is a list with the following components:

**outcomes** An ffdof object listing the outcomes per person, including the time to event, and the outcome concept ID. Outcomes are not yet filtered based on risk window, since this is done at a later stage.

**cohorts** An ffdof object listing the persons in each cohort, listing their exposure status as well as the time to the end of the observation period and time to the end of the cohort (usually the end of the exposure era).

**covariates** An ffdof object listing the baseline covariates per person in the two cohorts. This is done using a sparse representation: covariates with a value of 0 are omitted to save space.

**exclude** An ffdof object listing for each outcome concept ID the persons that need to be excluded from the analysis because of prior outcomes.

**covariateRef** An ffdof object describing the covariates that have been extracted.

**metaData** A list of objects with information on how the cohortMethodData object was constructed.

The generic summary() function has been implemented for this object.

getDbOutcomes

*Get outcomes for persons in the cohorts***Description**

Gets the outcomes for the cohorts in the cohortMethodData object.

**Usage**

```
getDbOutcomes(connectionDetails = NULL, connection = NULL,
  cdmDatabaseSchema, oracleTempSchema = cdmDatabaseSchema, cohortMethodData,
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence", outcomeIds = "",
  outcomeConditionTypeConceptIds = "", cdmVersion = "4")
```

**Arguments**

connectionDetails

An R object of type ConnectionDetails created using the function createConnectionDetails in the DatabaseConnector package.

connection

A connection to the server containing the schema as created using the connect function in the DatabaseConnector package.

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'.

cohortMethodData

An object of type cohortMethodData as generated using getDbCohortMethodData.

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.

outcomeConditionTypeConceptIds

A list of TYPE\_CONCEPT\_ID values that will restrict condition occurrences. Only applicable if outcomeTable = CONDITION\_OCCURRENCE.

cdmVersion

Define the OMOP CDM version used: currently support "4" and "5".

oracleTempSchemaA

schema where temp tables can be created in Oracle.

outcomeConceptIds

A list of CONCEPT\_IDs used to define outcomes. If outcomeTable = CONDITION\_OCCURRENCE, the list is a set of ancestor CONCEPT\_IDs, and all occurrences of all descendant concepts will be selected. If outcomeTable <> CONDITION\_OCCURRENCE, the list contains records found in COHORT\_DEFINITION\_ID field.

**Details**

If the connection parameter is specified, the cohorts are already assumed to be on the server in the appropriate temp table. Else, the temp table will be created by loading the cohorts from the cohortMethodData object to the server. This function can be used to add additional outcomes to an existing cohortMethodData object.

**Value**

The original cohortMethodData object with the new outcome data added.

---

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

---

**Description**

getOutcomeModel shows 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 createOutcomeModel function.
cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.

**Details**

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

**Examples**

```
# todo
```

---

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

---

**Description**

getPsModel shows the propensity score model

**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.

**Details**

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

**Examples**

```
# todo
```

---

grepCovariateNames	<i>Extract covariate names</i>
--------------------	--------------------------------

---

**Description**

Extracts covariate names using a regular-expression.

**Usage**

```
grepCovariateNames(pattern, object)
```

**Arguments**

- pattern** A regular expression with which to name covariate names
- object** An R object of type cohortMethodData or covariateData.

**Details**

This function extracts covariate names that match a regular-expression for a cohortMethodData or covariateData object.

**Value**

Returns a data.frame containing information about covariates that match a regular expression. This data.frame has the following columns:

- covariateId** Numerical identifier for use in model fitting using these covariates
- covariateName** Text identifier
- analysisId** Analysis identifier
- conceptId** OMOP common data model concept identifier, or 0

---

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 data from a folder</i>
----------------------	---

---

**Description**

loadCohortMethodData loads an object of type cohortMethodData from a folder in the file system.

**Usage**

```
loadCohortMethodData(file, readOnly = FALSE)
```

**Arguments**

file	The name of the folder containing the data.
readOnly	If true, the data is opened read only.

**Details**

The data will be written to a set of files in the folder specified by the user.

**Value**

An object of class cohortMethodData.

**Examples**

```
# todo
```



---

loadDrugComparatorOutcomesList

*Load a list of drugComparatorOutcomes from file*


---

**Description**

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

**Usage**

```
loadDrugComparatorOutcomesList(file)
```

**Arguments**

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

**Value**

A list of objects of type drugComparatorOutcome.

---

matchOnPs

*Match persons by propensity score*


---

**Description**

matchOnPs uses the provided propensity scores to match treated to comparator persons.

**Usage**

```
matchOnPs(data, caliper = 0.25, caliperScale = "standardized",
  maxRatio = 1, stratificationColumns = c())
```

**Arguments**

data	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. Two scales are supported: caliperScale = 'propensity score' or caliperScale = 'standardized'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution.
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 treated person.
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 at least the following three columns:

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

This function implements the greedy variable-ratio matching algorithm described in Rassen et al (2012).

## 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.

## 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

matchOnPsAndCovariates uses the provided propensity scores and a set of covariates to match treated to comparator persons.

## Usage

```
matchOnPsAndCovariates(data, caliper = 0.25, caliperScale = "standardized",
                        maxRatio = 1, cohortMethodData, covariateIds)
```

**Arguments**

<code>data</code>	A data frame with the three columns described below.
<code>caliper</code>	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.
<code>caliperScale</code>	The scale on which the caliper is defined. Two scales are supported: <code>caliperScale = 'propensity score'</code> or <code>caliperScale = 'standardized'</code> . On the standardized scale, the caliper is interpreted in standard deviations of the propensity score distribution.
<code>maxRatio</code>	The maximum number of persons in the comparator arm to be matched to each person in the treatment arm. A <code>maxRatio</code> of 0 means no maximum: all comparators will be assigned to a treated person.
<code>cohortMethodData</code>	An object of type <code>cohortMethodData</code> as generated using <code>getDbCohortMethodData</code> .
<code>covariateIds</code>	One or more covariate IDs in the <code>cohortMethodData</code> object on which subjects should be also matched.

**Details**

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

<code>rowId</code>	(integer)	A unique identifier for each row (e.g. the person ID)
<code>treatment</code>	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
<code>propensityScore</code>	(real)	Propensity score

This function implements the greedy variable-ratio matching algorithm described in Rassen et al (2012).

**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.

**Examples**

```
# todo
```

---

```
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,
  fileName = NULL)
```

### Arguments

<code>balance</code>	A data frame created by the <code>computeCovariateBalance</code> function.
<code>n</code>	Count of variates to plot.
<code>maxNameWidth</code>	Covariate names longer than this number of characters are truncated to create a nicer 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.

### Value

A ggplot object. Use the [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, fileName = NULL)
```

### Arguments

<code>balance</code>	A data frame created by the <code>computeCovariateBalance</code> function.
<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.

### 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 survival plot

**Usage**

```
plotKaplanMeier(outcomeModel, censorMarks = FALSE,
  confidenceIntervals = TRUE, includeZero = TRUE, dataCutoff = 0.99,
  treatmentLabel = "Treated", comparatorLabel = "Comparator",
  fileName = NULL)
```

**Arguments**

outcomeModel	An object of type outcomeModel as generated using the fitOutcomeModel function.
censorMarks	Whether or not to include censor marks in the plot.
confidenceIntervals	Plot 95 percent confidence intervals?
includeZero	Should the y axis include zero, or only go down to the lowest observed survival?
dataCutoff	Fraction of the data (number censored) after which the graph will not be shown.
treatmentLabel	A label to use for the treated cohort.
comparatorLabel	A label to use 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.

**Examples**

```
# todo
```

---

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

---

**Description**

plotPs shows the propensity (or preference) score distribution

**Usage**

```
plotPs(data, unfilteredData = NULL, scale = "preference",
  type = "density", binWidth = 0.05, fileName = NULL)
```

**Arguments**

<code>data</code>	A data frame with at least the two columns described below
<code>unfilteredData</code>	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 <code>data</code> .
<code>scale</code>	The scale of the graph. Two scales are supported: <code>scale = 'propensity'</code> or <code>scale = 'preference'</code> . The preference score scale is defined by Walker et al (2013).
<code>type</code>	Type of plot. Two possible values: <code>type = 'density'</code> or <code>type = 'histogram'</code>
<code>binWidth</code>	For histograms, the width of the bins
<code>fileName</code>	Name of the file where the plot should be saved, for example <code>'plot.png'</code> . See the function <code>ggsave</code> in the <code>ggplot2</code> package for supported file formats.

**Details**

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

<code>treatment</code>	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
<code>propensityScore</code>	(real)	Propensity score

**Value**

A ggplot object. Use the [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)
```

---

```
recomputePsForFullData
```

*Recompute the PS for the full data set*

---

**Description**

Recompute the PS for the full data set

**Usage**

```
recomputePsForFullData(psSample, cohortMethodDataSample, cohortMethodData)
```

**Arguments**

psSample            The propensity scores as created on the sample data.  
 cohortMethodDataSample  
                     The sample data on which the PS model was fitted.  
 cohortMethodData  
                     The full data.

**Details**

After using the [sampleCohorts](#) or [sampleComparator](#) to reduce the population size, this function can be used to apply a propensity model fitted on the sample to the full data.

**Value**

A new propensity score object.

---

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

---

**Description**

Run a list of analyses

**Usage**

```
runCmAnalyses(connectionDetails, cdmDatabaseSchema,
  oracleTempSchema = cdmDatabaseSchema,
  exposureDatabaseSchema = cdmDatabaseSchema, exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema,
  outcomeTable = "condition_occurrence",
  outputFolder = "./CohortMethodOutput", cmAnalysisList,
  drugComparatorOutcomesList, refitPsForEveryOutcome = FALSE,
  underSampleComparatorToTreatedRatio = 0, getDbCohortMethodDataThreads = 1,
  createPsThreads = 1, psCvThreads = 1, trimMatchStratifyThreads = 1,
  computeCovarBalThreads = 1, fitOutcomeModelThreads = 1,
  outcomeCvThreads = 1)
```

**Arguments**

connectionDetails  
                     An R object of type connectionDetails created using the function createConnectionDetails in the DatabaseConnector package.

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

oracleTempSchema  
                     For Oracle only: the name of the database schema where you want all temporary tables to be managed. Requires create/insert permissions to this database.

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.
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.
drugComparatorOutcomesList	A list of objects of type drugComparatorOutcomes as created using the <a href="#">createdrugComparatorOutcomes</a> function.
refitPsForEveryOutcome	Should the propensity model be fitted for every outcome (i.e. after people who already had the outcome are removed)? If false, a single propensity model will be fitted, and people who had the outcome previously will be removed afterwards.
underSampleComparatorToTreatedRatio	If the comparator group size exceeds the treated group size by this factor, the comparator group will be down-sampled before fitting the PS model. This can be useful when the comparator group is extremely large.
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'.
trimMatchStratifyThreads	The number of parallel threads to use for trimming, matching and stratifying.
computeCovarBalThreads	The number of parallel threads to use for computing the covariate balance.
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'.



## Details

Run a list of analyses for the drug-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(drugComparatorOutcomesList)` (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.

## Value

A data frame with the following columns:

<code>analysisId</code>	The unique identifier for a set of analysis choices.
<code>targetId</code>	The ID of the target drug.
<code>comparatorId</code>	The ID of the comparator group.
<code>indicationConceptIds</code>	The ID(s) of indications in which to nest to study.
<code>exclusionConceptIds</code>	The ID(s) of concepts used to exclude subjects.
<code>excludedCovariateConceptIds</code>	The ID(s) of concepts that cannot be used to construct covariates.
<code>includedCovariateConceptIds</code>	The ID(s) of concepts that should be used to construct covariates.
<code>cohortMethodDataFolder</code>	The ID of the outcome.
<code>sharedPsFile</code>	The name of the file containing the propensity scores of the shared propensity model. This model is used to create the outcome-specific propensity scores by removing people with prior outcomes.
<code>psFile</code>	The name of file containing the propensity scores for a specific outcomes (ie after people with prior outcomes have been removed).
<code>subPopFile</code>	The name of the file containing the identifiers of the population after any trimming, matching or stratifying, including their strata.
<code>covariateBalanceFile</code>	The name of the file containing the covariate balance (ie. the output of the <code>computeCovariateBalance</code> function).
<code>outcomeModelFile</code>	The name of the file containing the outcome model.

---

<code>sampleCohorts</code>	<i>Sample the target or comparator group down</i>
----------------------------	---

---

## Description

Sample the target or comparator group down

## Usage

```
sampleCohorts(cohortMethodData, treatedSampleSize = 10000,
               comparatorSampleSize = 20000)
```

## Arguments

`cohortMethodData`  
The original `cohortMethodData`.

`treatedSampleSize`  
The sample size for the treated cohort.

comparatorSampleSize

The sample size for the comparator cohort.

### Details

When the target and/or comparator cohorts are extremely large, it may be more efficient to only use a sample to fit the propensity model. This function creates a new cohortMethodData object with sampled populations.

### Value

An object of type cohortMethodData with the sampled populations.

---

sampleComparator	<i>Sample the comparator group down</i>
------------------	---

---

### Description

Sample the comparator group down

### Usage

```
sampleComparator(cohortMethodData, comparatorToTreatedRatio = 2)
```

### Arguments

cohortMethodData

The original cohortMethodData.

comparatorToTreatedRatio

The ratio between comparator and treated group.

### Details

When the comparator group is extremely large, it may be more efficient to only use a sample to fit the propensity model. This function creates a new cohortMethodData object where the comparator group is sampled down to a size relative to the treated group.

### Value

An object of type cohortMethodData with the sampled populations.

---

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 data to folder</i>
----------------------	---------------------------------------

---

**Description**

saveCohortMethodData saves an object of type cohortMethodData to folder.

**Usage**

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

**Arguments**

cohortMethodData	An object of type cohortMethodData as generated using getDbCohortMethodData.
file	The name of the folder where the data will be written. The folder should not yet exist.

**Details**

The data will be written to a set of files in the folder specified by the user.

**Examples**

```
# todo
```

---

```
saveDrugComparatorOutcomesList
```

*Save a list of drugComparatorOutcome to file*

---

### Description

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

### Usage

```
saveDrugComparatorOutcomesList(drugComparatorOutcomesList, file)
```

### Arguments

drugComparatorOutcomesList	The drugComparatorOutcomes list to be written to file
file	The name of the file where the results will be written

---

```
simulateCohortMethodData
```

*Generate simulated data*

---

### Description

simulateCohortMethodData creates a cohortMethodData object with simulated data.

### Usage

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

### Arguments

cohortDataSimulationProfile	An object of type cohortDataSimulationProfile 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

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

### Usage

```
stratifyByPs(data, numberOfStrata = 5, stratificationColumns = c())
```

### Arguments

data	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 treated 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.

### Details

The data frame should have the following three columns:

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

### Value

Returns a data frame 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	<i>Stratify persons by propensity score and other covariates</i>
---------------------------	--

---

### Description

stratifyByPsAndCovariates uses the provided propensity scores and covariates to stratify persons.

**Usage**

```
stratifyByPsAndCovariates(data, numberOfStrata = 5, cohortMethodData,
  covariateIds)
```

**Arguments**

<code>data</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 treated persons.
<code>cohortMethodData</code>	An object of type <code>cohortMethodData</code> as generated using <code>getDbCohortMethodData</code> .
<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:

<code>rowId</code>	(integer)	A unique identifier for each row (e.g. the person ID)
<code>treatment</code>	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
<code>propensityScore</code>	(real)	Propensity score

**Value**

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

**Examples**

```
# todo
```

---

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

---

**Description**

Create a summary report of the analyses

**Usage**

```
summarizeAnalyses(outcomeReference)
```

**Arguments**

<code>outcomeReference</code>	A data.frame as created by the <code>runAnalyses</code> function.
-------------------------------	---

**Value**

A data frame with the following columns:

analysisId	The unique identifier for a set of analysis choices.
targetId	The ID of the target drug.
comparatorId	The ID of the comparator group.
indicationConceptIds	The ID(s) of indications in which to nest to study.
outcomeId	The ID of the outcome.
rr	The estimated effect size.
ci95lb	The lower bound of the 95 percent confidence interval.
ci95ub	The upper bound of the 95 percent confidence interval.
treated	The number of subjects in the treated group (after any trimming and matching).
comparator	The number of subjects in the comparator group (after any trimming and matching).
eventsTreated	The number of outcomes in the treated group (after any trimming and matching).
eventsComparator	The number of outcomes in the comparator group (after any trimming and matching).
logRr	The log of the estimated relative risk.
seLogRr	The standard error of the log of the estimated relative risk.

---

trimByPs

*Trim persons by propensity score*


---

## Description

trimByPs uses the provided propensity scores to trim subjects with extreme scores.

## Usage

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

## Arguments

data	A data frame with the three columns described below
trimFraction	This fraction will be removed from each treatment group. In the treatment 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	(integer)	A unique identifier for each row (e.g. the person ID)
treatment	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
propensityScore	(real)	Propensity score

## Value

Returns a data frame with the same three columns as the input.

**Examples**

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

---

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

---

**Description**

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

**Usage**

```
trimByPsToEquipoise(data, bounds = c(0.25, 0.75))
```

**Arguments**

data	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	(integer)	A unique identifier for each row (e.g. the person ID)
treatment	(integer)	Column indicating whether the person is in the treated (1) or comparator (0) group
propensityScore	(real)	Propensity score

**Value**

Returns a data frame 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)
```



---

`vignetteAnalysisSummary`*Analysis summary data for the vignette*

---

**Description**

Analysis summary data for the vignette

**Usage**

```
data(vignetteAnalysisSummary)
```

---

`vignetteBalance`*Balance data for the vignette*

---

**Description**

Balance data for the vignette

**Usage**

```
data(vignetteBalance)
```

---

`vignetteOutcomeModel1` *Outcome data for the vignette*

---

**Description**

Outcome data for the vignette

**Usage**

```
data(vignetteOutcomeModel1)
```

---

`vignetteOutcomeModel2` *Outcome data for the vignette*

---

**Description**

Outcome data for the vignette

**Usage**

```
data(vignetteOutcomeModel2)
```

---

vignetteOutcomeModel3	<i>Outcome data for the vignette</i>
-----------------------	--------------------------------------

---

**Description**

Outcome data for the vignette

**Usage**

```
data(vignetteOutcomeModel3)
```

---

vignettePs	<i>Propensity scores for the vignette</i>
------------	---

---

**Description**

Propensity scores for the vignette

**Usage**

```
data(vignettePs)
```

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