# Package 'CohortMethod'

July 28, 2015

```
Type Package

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

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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,
      ffbase (>= 0.12.1),
      plyr,
      Rcpp (>= 0.11.2),
      RJDBC,
      SqlRender (\geq 1.1.1),
      survival,
      rjson,
      OhdsiRTools
Suggests testthat,
```

pROC,

2 R topics documented:

gnm, knitr, rmarkdown LinkingTo Rcpp NeedsCompilation yes

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 ${\tt CohortMethod}$ 

CohortMethod

# Description

CohortMethod

 ${\it cohort} {\it MethodDataSimulationProfile} \\ {\it A simulation profile}$ 

# Description

A simulation profile

# Usage

data(cohortMethodDataSimulationProfile)

compute Covariate Balance

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.

# Usage

computeCovariateBalance(restrictedCohorts, cohortMethodData, outcomeId = NULL)

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### **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 date frame describing the covariate balance before and after matching/trimming.

computePsAuc Compute the area under the ROC curve

# **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 a 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

constructEras 5

#### 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)</pre>
```

constructEras

Build eras

### **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'.

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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 verion 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)
```

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createCmAnalysis Create a CohortMethod analysis specification	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**

matchOnPsAndCovariates

٤	guments		
	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	Type 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.	
	getDbCohortMeth	odDataArgs	
		An object representing the arguments to be used when calling the $\ensuremath{getDbCohortMethodData}$ function.	
	createPs	Should the createPs function be used in this analysis?	
	createPsArgs	An object representing the arguments to be used when calling the <pre>createPs</pre> function.	
	trimByPs	Should the trimByPs function be used in this analysis?	
	trimByPsArgs	An object representing the arguments to be used when calling the trimByPs function.	
	trimByPsToEquipoise		
		Should the trimByPsToEquipoise function be used in this analysis?	
	trimByPsToEquipoiseArgs		
		An object representing the arguments to be used when calling the trimByPsToEquipoise function.	
	matchOnPs	Should the matchOnPs function be used in this analysis?	
	matchOnPsArgs	An object representing the arguments to be used when calling the matchOnPs function.	

Should the matchOnPsAndCovariates function be used in this analysis?

matchOnPsAndCovariatesArgs

An object representing the arguments to be used when calling the matchOnPsAndCovariates
function.

stratifyByPs Should the stratifyByPs function be used in this analysis? stratifyByPsArgs

An object representing the arguments to be used when calling the stratifyByPs function.

stratifyByPsAndCovariates

Should the stratifyByPsAndCovariates function be used in this analysis?

stratify By Ps And Covariates Args

An object representing the arguments to be used when calling the stratifyByPsAndCovariates
function.

compute Covariate Balance

Should the computeCovariateBalance function be used in this analysis?

fitOutcomeModel

Should the fitOutcomeModel function be used in this analysis?

fitOutcomeModelArgs

An object representing the arguments to be used when calling the fitOutcomeModel function.

#### **Details**

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

 $create {\tt CohortMethodDataSimulationProfile}$ 

Create simulation profile

# **Description**

createCohortMethodDataSimulationProfile creates a profile based on the provided cohort-MethodData 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

 $An\ object\ of\ type\ cohort {\tt DataSimulationProfile}.$ 

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createCreatePsArgs

Create a parameter object for the function createPs

### **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 createPriorfor details.

control The control object used to control the cross-validation used todetermine the hy-

perparameters of the prior (if applicable). SeecreateControl for details.

# **Details**

Create an object defining the parameter values.

```
createDrugComparatorOutcomes
```

Create drug-comparator-outcomes combinations.

# 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 createCmAnalysis 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 createCmAnalysis 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 createCmAnalysis 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 expo-

sure 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 hy-

perparameters of the prior (if applicable). SeecreateControl for details.

#### **Details**

Create an object defining the parameter values.

 ${\tt createGetDbCohortMethodDataArgs}$ 

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 per-

son 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 anydescendant conditions/drugs/procedures occurring at least onceanytime 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 descendantconcepts) be excluded from the covariates? Note that this willwork if the drugs are actualy drug concept IDs (and not cohortIDs).

covariateSettings

An object of type covariateSettings as created using thecreateCovariateSettings 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  $\boldsymbol{0}$ 

means no caliper is used.

caliperScale The scale on which the caliper is defined. Two scales are supported: caliperScale

= 'propensity score' or caliper Scale = 'standardized'. On the standardized scale, the caliper is interpreted in standard deviations of the propensity scoredistribu-

tion.

maxRatio The maximum number of persons int the comparator arm to be matched to each-

person in the treatment arm. A maxRatio of 0 means no maximum: allcompara-

tors will be assigned to a treated person.

covariateIds One or more covariate IDs in the cohortMethodData object on whichsubjects

should be also matched.

#### **Details**

Create an object defining the parameter values.

createMatchOnPsArgs 13

	C	1-:4 C 41-	f 4: 4 - 1. O - D -
createMatchOnPsArgs	Create a paramete	er obiect for the	function matchOnPs

### **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 isacceptable 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 isinterpreted in standard deviations of the propensity scoredistribu-

tion.

maxRatio The maximum number of persons int the comparator arm to be matched toeach

person in the treatment arm. A maxRatio of 0 means no maximum:all compara-

tors will be assigned to a treated person.

stratificationColumns

Names or numbers of one or more columns in the data data.frameon which subjects should be stratified prior to matching. No personswill be matched with

persons outside of the strata identified by thevalues in these columns.

# **Details**

Create an object defining the parameter values.

createPs	Create propensity scores	

# **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 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**

createPs creates propensity scores using a regularized logistic regression.

# **Examples**

```
data(cohortDataSimulationProfile)
cohortMethodData <- simulateCohortMethodData(cohortDataSimulationProfile, n = 1000)
ps <- createPs(cohortMethodData)</pre>
```

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? Theboundaries of

the strata are automatically defined to contain equalnumbers of treated persons.

covariateIds One or more covariate IDs in the cohortMethodData object on whichsubjects

should also be stratified.

### **Details**

Create an object defining the parameter values.

createStratifyByPsArgs

 ${\tt createStratifyByPsArgs}$ 

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 whichsubjects should also be stratified in addition to stratification onpropensity 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**

 ${\it trim} {\it Fraction}$ 

This fraction will be removed from each treatment group. In the treatmentgroup, 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.

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```
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.

# **Description**

drawAttritionDiagram draws the attition 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 he createOutcomeMode

function.

treatmentLabel A label to us for the treated cohort.

comparatorLabel

A label to us for the comparator cohort.

fileName Name of the file where the plot should be saved, for example 'plot.png'. See the

function ggsave in the ggplot2 package for supported file formats.

### Value

A ggplot object. Use the ggsave function to save to file in a different format.

fitOutcomeModel 17

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 expo-

sure if the addExposureDaysToEnd parameter is specified).

 $add {\tt Exposure Days To End}$ 

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 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**

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

 ${\tt connectionDetails\ created\ using\ the\ function\ createConnectionDetails\ in\ the\ DatabaseConnector\ package.}$ 

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 exposure Table = DRUG ERA, comparatorId is a CONCEPT ID and all descendant concepts within that CON-CEPT\_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 mininum 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 outcome Table = 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 exposure Table = 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 exposure Table 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 exposure Table = 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 outcome Table <> CONDI-TION OCCURRENCE, then expectation is outcome Table has format of CO-HORT 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 actualy 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 ffdf object listing the outcomes per person, including the time to event, and the outcome conncept ID. Outcomes are not yet filtered based on risk window, since this is done at a later stage.

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

covariateRef An ffdf 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.

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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.

 $\verb"outcomeConditionTypeConceptIds"$ 

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.

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#### **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

Get the outcome model

# **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 he createOutcomeMode 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

Get the propensity model

# Description

getPsModel shows the propensity score model

# Usage

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

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### **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

Extract covariate names

### **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

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 $load {\tt CmAnalysisList}$ 

Load a list of cmAnalysis from file

# **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 Load the cohort data from a folder

# 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

 ${\tt loadDrugComparatorOutcomesList}$ 

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:

 ${\tt caliperScale = 'propensity \ score' \ or \ caliperScale = 'standardized'.}$  On the standardized scale, the caliper is interpreted in standard deviations of the

propensity score distribution.

maxRatio The maximum number of persons int the comparator arm to be matched to each

person in the treatment arm. A maxRatio of 0 means no maximum: all com-

parators 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 date 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**

 ${\tt matchOnPsAndCovariates}$ 

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)
```

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### **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 int the comparator arm to be matched to each

person in the treatment arm. A maxRatio of 0 means no maximum: all com-

parators will be assigned to a treated person.

cohortMethodData

An object of type cohortMethodData as generated using getDbCohortMethodData.

covariateIds One or more covariate IDs in the cohortMethodData object on which subjects

should be also matched.

#### 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 date 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**

balance A data frame created by the computeCovariateBalance funcion.

n Count of variates to plot.

maxNameWidth Covariate names longer than this number of characters are truncated to create a

nicer 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.

 $\verb|plotCovariateBalanceScatterPlot|$ 

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**

balance A data frame created by the compute Covariate Balance function.

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.

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plotKaplanMeier

Plot the Kaplan-Meier curve

### **Description**

plotKaplanMeier creates the Kaplain-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 he fitOutcomeModel func-

tion.

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.

 ${\tt treatmentLabel} \ \ A \ label \ to \ us \ for \ the \ treated \ cohort.$ 

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.

# **Examples**

# todo

plotPs

Plot the propensity score distribution

### **Description**

plotPs shows the propensity (or preference) score distribution

# Usage

```
plotPs(data, unfilteredData = NULL, scale = "preference",
  type = "density", binWidth = 0.05, 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. Two possible values: type = 'density' or type = 'histogram'

binWidth For histograms, the width of the bins

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.

### **Details**

The data frame should have a 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 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)</pre>
```

recomputePsForFullData

Recompute the PS for the full data set

# **Description**

Recompute the PS for the full data set

# Usage

 $recompute Ps For Full Data (ps Sample, \ cohort Method Data Sample, \ cohort Method Data)$ 

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### **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

Run a list of analyses

# **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.

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#### 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 createCmAnalysis function.

drugComparatorOutcomesList

A list of objects of type drugComparatorOutcomes as created using the createdrugComparatorOutc

#### 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.

### under Sample Comparator To Treated Ratio

function.

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.

### ${\tt getDbCohortMethodDataThreads}$

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'.

### trimMatchStratifyThreads

 $\label{threads} The number of parallel threads to use for trimming, matching and stratifying. \\ {\tt computeCovarBalThreads}$ 

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'.

sampleCohorts 33

#### **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.

sampleCohorts

Sample the target or comparator group down

### **Description**

Sample the target or comparator group down

# Usage

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

### **Arguments**

cohortMethodData

The original cohortMethodData.

treatedSampleSize

The sampe 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.

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sampleComparator

Sample the comparator group down

# 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 to comparator group is sampled down to a size relative to the treated group.

### Value

An object of type cohortMethodData with the sampled populations.

save CmAnalysis List

Save a list of cmAnalysis to file

# **Description**

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

# Usage

```
{\tt 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 35

saveCohortMethodData Save the cohort data to folder

# Description

 ${\tt saveCohortMethodData}\ saves\ an\ object\ of\ type\ cohortMethodData\ to\ folder.$ 

# 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

 ${\tt saveDrugComparatorOutcomesList}$ 

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

save Drug Comparator Outcomes List (drug Comparator Outcomes List, file)

# **Arguments**

 ${\tt drugComparatorOutcomesList}$ 

The drugComparatorOutcomes list to be written to file

file The name of the file where the results will be written

36 stratifyByPs

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

Stratify persons by propensity score

### **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 date 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)</pre>
```

stratifyByPsAndCovariates

Stratify persons by propensity score and other covariates

# Description

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

### Usage

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

### **Arguments**

data A data frame with the three columns described below

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.

 ${\tt cohortMethodData}$ 

An object of type cohort Method Data as generated using getDbCohort Method Data.

covariateIds One or more covariate IDs in the cohortMethodData object on which subjects

should also be stratified.

38 trimByPs

### **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 date frame with the same columns as the input data plus one extra column: stratumId.

### **Examples**

# todo

summarizeAnalyses

Create a summary report of the analyses

# **Description**

Create a summary report of the analyses

# Usage

summarizeAnalyses(outcomeReference)

# **Arguments**

outcomeReference

A data.frame as created by the runAnalyses function.

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.

trimByPsToEquipoise 39

### **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 date 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)</pre>
```

trimByPsToEquipoise

Keep only persons in clinical equipoise

### **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 date 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)</pre>
```

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)
```

 $\verb|vignetteOutcomeModel2|| Outcome \ data \ for \ the \ vignette$ 

# Description

Outcome data for the vignette

# Usage

```
data(vignetteOutcomeModel2)
```

 ${\tt vignetteOutcomeModel3} \ \ \textit{Outcome data for the vignette}$ 

# Description

Outcome data for the vignette

# Usage

data(vignetteOutcomeModel3)

vignettePs

Propensity scores for the vignette

# Description

Propensity scores for the vignette

# Usage

data(vignettePs)

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