

Package ‘MethodEvaluation’

April 15, 2016

Type Package

Title Package for evaluation of estimation methods

Version 0.0.2

Date 2016-04-12

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Description This package contains resources for the evaluation of the performance of methods that aim to estimate the magnitude (relative risk) of the effect of a drug on an outcome. These resources include reference sets for evaluating methods on real data, as well as functions for inserting simulated effects in real data based on negative control drug-outcome pairs. Further included are functions for the computation of the minimum detectable relative risks and functions for computing performance statistics such as predictive accuracy, error and bias.

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Depends R (>= 3.2.0),
DatabaseConnector (>= 1.3.0),
FeatureExtraction,
Cyclops

Imports ff,
ffbase (>= 0.12.1),
RJDBC,
SqlRender (>= 1.1.2),
pROC,
ggplot2

Suggests testthat

RoxygenNote 5.0.1

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computeAuc	<i>Compute the area under the ROC curve</i>
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Description

Compute the area under the ROC curve

Usage

```
computeAuc(methodResults, referenceSet, confidenceIntervals = TRUE)
```

computeAucsInjectedSignals	<i>Compute the AUCs for various injected signal sizes</i>
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Description

Compute the AUCs for various injected signal sizes

Usage

```
computeAucsInjectedSignals(logRr, trueLogRr)
```

Arguments

logRr	A vector containing the log of the relative risk as estimated by a method.
trueLogRr	A vector containing the injected log(relative risk) for each estimate.

Value

A data frame with per injected signal size the AUC and the 95 percent confidence interval of the AUC.

computeBias	<i>Compute the bias (mean error)</i>
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Description

Compute the bias (mean error)

Usage

```
computeBias()
```

computeMdr	<i>Compute minimal detectable relative risk (MDRR)</i>
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Description

computeMdr computes the minimal detectable relative risk (MDRR) for drug-outcome pairs.

Usage

```
computeMdr(connectionDetails, cdmDatabaseSchema,
  oracleTempSchema = cdmDatabaseSchema, exposureOutcomePairs,
  exposureDatabaseSchema = cdmDatabaseSchema, exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema, outcomeTable = "condition_era",
  outcomeConditionTypeConceptIds = c(), cdmVersion = "4")
```

Arguments

connectionDetails

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

cdmDatabaseSchema

Name of database schema that contains OMOP CDM and vocabulary.

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.

exposureOutcomePairs

A data frame with at least two columns:

- "exposureConceptId" containing the drug_concept_ID or cohort_definition_id of the exposure variable
- "outcomeConceptId" containing the condition_concept_ID or cohort_definition_id of the outcome variable

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.
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".

Details

Computes the MDRR using simple power-calculations using person-level statistics stratified by age and gender.

Value

A data frame containing the MDRRs for the given exposure-outcome pairs.

Examples

```
## Not run:
connectionDetails <- createConnectionDetails(dbms = "sql server",
                                             server = "RNDUSRDHIT07.jnj.com")
exposureOutcomePairs <- data.frame(exposureConceptId = c(767410, 1314924, 907879),
                                   outcomeConceptId = c(444382, 79106, 138825))
mdrrs <- computeMdrd(connectionDetails,
                     "cdm_truven_mdc",
                     exposureOutcomePairs,
                     outcomeTable = "condition_era")

## End(Not run)
```

computeMse

Compute the mean squared error

Description

Compute the mean squared error

Usage

```
computeMse()
```

createOutcomeCohorts *Create outcomes of interest*

Description

Create outcomes of interest

Usage

```
createOutcomeCohorts(connectionDetails, cdmDatabaseSchema,
  createNewCohortTable = FALSE, cohortDatabaseSchema = cdmDatabaseSchema,
  cohortTable = "cohort", referenceSet = "omopReferenceSet",
  cdmVersion = "4")
```

Arguments

connectionDetails	An R object of type ConnectionDetails created using the function createConnectionDetails in the DatabaseConnector package.
cdmDatabaseSchema	A database schema containing health care data in the OMOP Common Data Model. Note that for SQL Server, both the database and schema should be specified, e.g. 'cdm_schema.dbo'
createNewCohortTable	Should a new cohort table be created, or should the outcomes be inserted in a existing table?
cohortDatabaseSchema	The database schema where the target table is located. Note that for SQL Server, both the database and schema should be specified, e.g. 'cdm_schema.dbo'
cohortTable	The name of the table where the outcomes will be stored.
cdmVersion	Define the OMOP CDM version used: currently support "4" and "5".

Details

This function will create the outcomes of interest referenced in the various reference sets. The outcomes of interest are derived using information like diagnoses, procedures, and drug prescriptions. The outcomes are stored in a table on the database server.

euadrReferenceSet *The EU-ADR reference set*

Description

A reference set of 43 drug-outcome pairs where we believe the drug causes the outcome (positive controls) and 50 drug-outcome pairs where we believe the drug does not cause the outcome (negative controls). The controls involve 10 health outcomes of interest. Note that originally, there was an additional positive control (Nimesulide and acute liver injury), but Nimesulide is not in RxNorm, and is not available in many countries.

Usage

```
data(euadrReferenceSet)
```

Format

A data frame with 399 rows and 10 variables:

exposureConceptId Concept ID identifying the exposure

exposureConceptName Name of the exposure

outcomeConceptId Concept ID identifying the outcome

outcomeConceptName Name of the outcome

groundTruth 0 = negative control, 1 = positive control

indicationConceptId Concept Id identifying the (primary) indication of the drug. To be used when one wants to nest the analysis within the indication

indicationConceptName Name of the indication

comparatorDrugConceptId Concept ID identifying a comparator drug that can be used as a counterfactual

comparatorDrugConceptName Name of the comparator drug

comparatorType How the comparator was selected

References

Coloma PM, Avillach P, Salvo F, Schuemie MJ, Ferrajolo C, Pariente A, Fourrier-Reglat A, Molokhia M, Patadia V, van der Lei J, Sturkenboom M, TrifirC2 G. A reference standard for evaluation of methods for drug safety signal detection using electronic healthcare record databases. *Drug Safety* 36(1):13-23, 2013

filterOnMdr

Filter data based on MDRR

Description

Filters a dataset to those exposure-outcome pairs with sufficient power.

Usage

```
filterOnMdr(data, mdr, threshold = 1.25)
```

Arguments

data	A data frame with at least two columns: <ul style="list-style-type: none"> "exposureConceptId" containing the drug_concept_ID or cohort_definition_id of the exposure variable "outcomeConceptId" containing the condition_concept_ID or cohort_definition_id of the outcome variable
mdr	A data frame as generated by the computeMdr function.
threshold	The required minimum detectable relative risk.

Value

A subset of the data object.

injectSignals	<i>Inject signals in database</i>
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Description

Inject signals in database

Usage

```
injectSignals(connectionDetails, cdmDatabaseSchema,
  oracleTempSchema = cdmDatabaseSchema,
  exposureDatabaseSchema = cdmDatabaseSchema, exposureTable = "drug_era",
  outcomeDatabaseSchema = cdmDatabaseSchema, outcomeTable = "cohort",
  outputDatabaseSchema = outcomeDatabaseSchema, outputTable = outcomeTable,
  createOutputTable = FALSE, exposureOutcomePairs, modelType = "poisson",
  buildOutcomeModel = TRUE, minOutcomeCount = 100,
  covariateSettings = FeatureExtraction::createCovariateSettings(useCovariateDemographics
    = TRUE, useCovariateConditionOccurrence = TRUE,
  useCovariateConditionOccurrence365d = TRUE, useCovariateConditionOccurrence30d
    = TRUE, useCovariateConditionOccurrenceInpt180d = TRUE,
  useCovariateConditionEra = TRUE, useCovariateConditionEraEver = TRUE,
  useCovariateConditionEraOverlap = TRUE, useCovariateConditionGroup = TRUE,
  useCovariateDrugExposure = TRUE, useCovariateDrugExposure365d = TRUE,
  useCovariateDrugExposure30d = TRUE, useCovariateDrugEra = TRUE,
  useCovariateDrugEra365d = TRUE, useCovariateDrugEra30d = TRUE,
  useCovariateDrugEraEver = TRUE, useCovariateDrugEraOverlap = TRUE,
  useCovariateDrugGroup = TRUE, useCovariateProcedureOccurrence = TRUE,
  useCovariateProcedureOccurrence365d = TRUE, useCovariateProcedureOccurrence30d
    = TRUE, useCovariateProcedureGroup = TRUE, useCovariateObservation = TRUE,
  useCovariateObservation365d = TRUE, useCovariateObservation30d = TRUE,
  useCovariateObservationCount365d = TRUE, useCovariateMeasurement365d = TRUE,
  useCovariateMeasurement30d = TRUE, useCovariateMeasurementCount365d = TRUE,
  useCovariateMeasurementBelow = TRUE, useCovariateMeasurementAbove = TRUE,
  useCovariateConceptCounts = TRUE, useCovariateRiskScores = TRUE,
  useCovariateRiskScoresCharlson = TRUE, useCovariateRiskScoresDCSI = TRUE,
  useCovariateRiskScoresCHADS2 = TRUE, useCovariateRiskScoresCHADS2VAsc = TRUE,
  useCovariateInteractionYear = FALSE, useCovariateInteractionMonth = FALSE,
  excludedCovariateConceptIds = c(), deleteCovariatesSmallCount = 100),
  prior = createPrior("laplace", exclude = 0, useCrossValidation = TRUE),
  control = createControl(cvType = "auto", startingVariance = 0.1, noiseLevel
    = "quiet", threads = 10), firstExposureOnly = FALSE, washoutPeriod = 183,
  riskWindowStart = 0, riskWindowEnd = 0, addExposureDaysToEnd = TRUE,
  firstOutcomeOnly = FALSE, effectSizes = c(1, 1.25, 1.5, 2, 4),
  precision = 0.01, outputIdOffset = 1000,
  workFolder = "./SignalInjectionTemp", cdmVersion = "4")
```

Arguments

connectionDetails

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

<code>cdmDatabaseSchema</code>	Name of database schema that contains OMOP CDM and vocabulary.
<code>oracleTempSchema</code>	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.
<code>exposureDatabaseSchema</code>	The name of the database schema that is the location where the exposure data used to define the exposure cohorts is available. If <code>exposureTable = DRUG_ERA</code> , <code>exposureDatabaseSchema</code> is not used by assumed to be <code>cdmSchema</code> . Requires read permissions to this database.
<code>exposureTable</code>	The table name that contains the exposure cohorts. If <code>exposureTable <> DRUG_ERA</code> , then expectation is <code>exposureTable</code> has format of COHORT table: <code>cohort_concept_id</code> , <code>SUBJECT_ID</code> , <code>COHORT_START_DATE</code> , <code>COHORT_END_DATE</code> .
<code>outcomeDatabaseSchema</code>	The name of the database schema that is the location where the data used to define the outcome cohorts is available. If <code>exposureTable = CONDITION_ERA</code> , <code>exposureDatabaseSchema</code> is not used by assumed to be <code>cdmSchema</code> . Requires read permissions to this database.
<code>outcomeTable</code>	The table name that contains the outcome cohorts. When the table name is not <code>CONDITION_ERA</code> This table is expected to have the same format as the COHORT table: <code>SUBJECT_ID</code> , <code>COHORT_START_DATE</code> , <code>COHORT_END_DATE</code> , <code>COHORT_CONCEPT_ID</code> (CDM v4) or <code>COHORT_DEFINITION_ID</code> (CDM v5 and higher).
<code>outputDatabaseSchema</code>	The name of the database schema that is the location of the tables containing the new outcomes Requires write permissions to this database.
<code>outputTable</code>	The name of the table names that will contain the generated outcome cohorts.
<code>createOutputTable</code>	Should the output table be created prior to inserting the outcomes? If TRUE and the tables already exists, it will first be deleted. If FALSE, the table is assumed to exist and the outcomes will be inserted. Any existing outcomes with the same IDs will first be deleted.
<code>exposureOutcomePairs</code>	A data frame with at least two columns: <ul style="list-style-type: none"> • "exposureId" containing the <code>drug_concept_ID</code> or <code>cohort_concept_id</code> of the exposure variable • "outcomeId" containing the <code>condition_concept_ID</code> or <code>cohort_concept_id</code> of the outcome variable
<code>modelType</code>	Can be either "poisson" or "survival"
<code>buildOutcomeModel</code>	Should an outcome model be created to predict outcomes. New outcomes will be inserted based on the predicted probabilities according to this model, and this will help preserve the observed confounding when injecting signals.
<code>minOutcomeCount</code>	Minimum number of outcome events required to build a model and inject a signal.
<code>covariateSettings</code>	An object of type <code>covariateSettings</code> as created using the <code>createCovariateSettings</code> function in the <code>FeatureExtraction</code> package.

prior	The prior used to fit the outcome 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.
firstExposureOnly	Should signals be injected only for the first exposure? (ie. assuming an acute effect)
washoutPeriod	Number of days at the start of observation for which no signals will be injected, but will be used to determine whether exposure or outcome is the first one, and for extracting covariates to build the outcome model.
riskWindowStart	The start of the risk window relative to the start of the exposure (in days). When 0, risk is assumed to start on the first day of exposure.
riskWindowEnd	The end of the risk window relative to the start of the exposure. Note that typically the length of exposure is added to this number (when the addExposureDaysToEnd parameter is set to TRUE).
addExposureDaysToEnd	Should length of exposure be added to the risk window?
firstOutcomeOnly	Should only the first outcome per person be considered when modeling the outcome?
effectSizes	A numeric vector of effect sizes that should be inserted.
precision	The allowed ratio between target and injected signal size.
outputIdOffset	What should be the first new outcome ID that is to be created?
workFolder	Path to a folder where intermediate data will be stored.
cdmVersion	Define the OMOP CDM version used: currently support "4" and "5".

Details

This function will insert additional outcomes for a given set of drug-outcome pairs. It is assumed that these drug-outcome pairs represent negative controls, so the true relative risk before inserting any outcomes should be 1. There are two models for inserting the outcomes during the specified risk window of the drug: a Poisson model assuming multiple outcomes could occur during a single exposure, and a survival model considering only one outcome per exposure. For each

Value

A data.frame listing all the drug-pairs in combination with requested effect sizes and the real inserted effect size (might be different from the requested effect size because of sampling error).

MethodEvaluation

MethodEvaluation

Description

MethodEvaluation

omopReferenceSet	<i>The OMOP reference set A reference set of 165 drug-outcome pairs where we believe the drug causes the outcome (positive controls) and 234 drug-outcome pairs where we believe the drug does not cause the outcome (negative controls). The controls involve 4 health outcomes of interest: acute liver injury, acute kidney injury, acute myocardial infarction, and GI bleeding.</i>
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Description

The OMOP reference set A reference set of 165 drug-outcome pairs where we believe the drug causes the outcome (positive controls) and 234 drug-outcome pairs where we believe the drug does not cause the outcome (negative controls). The controls involve 4 health outcomes of interest: acute liver injury, acute kidney injury, acute myocardial infarction, and GI bleeding.

Usage

```
data(omopReferenceSet)
```

Format

A data frame with 399 rows and 10 variables:

exposureConceptId Concept ID identifying the exposure

exposureConceptName Name of the exposure

outcomeConceptId Concept ID identifying the outcome

outcomeConceptName Name of the outcome

groundTruth 0 = negative control, 1 = positive control

indicationConceptId Concept Id identifying the (primary) indication of the drug. To be used when one wants to nest the analysis within the indication

indicationConceptName Name of the indication

comparatorDrugConceptId Concept ID identifying a comparator drug that can be used as a counterfactual

comparatorDrugConceptName Name of the comparator drug

comparatorType How the comparator was selected

References

Ryan PB, Schuemie MJ, Welebob E, Duke J, Valentine S, Hartzema AG. Defining a reference set to support methodological research in drug safety. Drug Safety 36 Suppl 1:S33-47, 2013

plotBias	<i>Plot the error distribution</i>
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Description

Plot the error distribution

Usage

```
plotBias()
```

plotRocsInjectedSignals	<i>Plot the ROC curves for various injected signal sizes</i>
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Description

Plot the ROC curves for various injected signal sizes

Usage

```
plotRocsInjectedSignals(logRr, trueLogRr, showAucs, fileName = NULL)
```

Arguments

logRr	A vector containing the log of the relative risk as estimated by a method.
trueLogRr	A vector containing the injected log(relative risk) for each estimate.
showAucs	Should the AUCs be shown in the plot?
fileName	Name of the file where the plot should be saved, for example 'plot.png'. See the function ggsave in the ggplot2 package for supported file formats.

Value

A Ggplot object. Use the ggsave function to save to file.

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