

Using FeatureExtraction

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1 Introduction

The **FeatureExtraction** package can be used to create features for a cohort, using the information stored in the Common Data Model. A cohort is defined a set of persons who satisfy one or more inclusion criteria for a duration of time. Features can for example be diagnoses observed prior to entering the cohort. Some people might also refer to such features as ‘baseline characteristics’, or to features in general as ‘covariates’, and we will use those terms interchangeably throughout this vignette.

This vignette describes how features can be constructed using the default covariate definitions embeded in the package. Although these definitions allow quite some customization through predefined parameters, it is possible that someone needs more customization. In this case, the reader is referred to the other vignettes included in this package that deal with constructing completely custom covariates.

This vignette will first describe how to specify which features to construct. In many situations, for example when using **FeatureExtraction** as part of another package such as **CohortMethod** or **PatientLevelPrediction**, that is all one needs to know about the **FeatureExtraction** package, as the actual calling of the package is done by the other package. However, it is also possible to use this package on its own, for example to create a descriptive characterization of a cohort to include in a paper.

2 Covariate settings

Users can specify which covariates to construct in three ways:

1. Choose the default set of covariates.
2. Choose from a set of prespecified analyses.
3. Create a set of custom analyses.

An **analysis** is a process that creates one or more similar covariates. For example, one analysis might create a binary covariate for each condition observed in the `condition_occurrence` table in the year prior to cohort start, and another analysis might create a single covariate representing the Charlson comorbidity index.

Note that it is always possible to specify a set of concept IDs that can or can't be used to construct features. When choosing the default set (option 1) or the prespecified analysis (option 2) this can only be done across all analysis. When creating custom analyses (option 3) this can be specified per analysis.

For **advanced users**: It is also possible to specify a set of covariate IDs that need to be constructed. A covariate ID identifies a specific covariate, for example the Charlson comorbidity index, or the occurrence of a specific condition concept in a specific time window. A covariate ID is therefore not to be confused with a concept ID. The typical scenario where one might want to specify covariate IDs to construct is when someone already constructed covariates in one population, found a subset of covariates to be of interest, and would like to have only those covariates constructed in another population.

2.1 Using the default set of covariates

Using the default set of covariates is straightforward:

```
settings <- createDefaultCovariateSettings()
```

This will create a wide array of features, ranging from demographics, through conditions and drugs, to several risk scores.

Note that we could specify a set of concepts that should not be used to create covariates, for example:

```
settings <- createDefaultCovariateSettings(excludedCovariateConceptIds = 1124300,  
                                          addDescendantsToExclude = TRUE)
```

This will create the default set of covariates, except those derived from concept 1124300 (the ingredient diclofenac) and any of its descendants (ie. all drugs containing the ingredient diclofenac).

2.2 Using prespecified analyses

The function `createCovariateSettings` allow the user to choose from a large set of predefined covariates. Type `?createCovariateSettings` to get an overview of the available options. For example:

```
settings <- createCovariateSettings(useDemographicsGender = TRUE,  
                                   useDemographicsAgeGroup = TRUE,  
                                   useConditionOccurrenceAnyTimePrior = TRUE)
```

This will create binary covariates for gender, age (in 5 year age groups), and each concept observed in the `condition_occurrence` table any time prior to (and including) the cohort start date.

Many of the prespecified analyses refer to a short, medium, or long term time window. By default, these windows are defined as:

- Long term: 365 days prior up to and including the cohort start date.
- Short term: 180 days prior up to and including the cohort start date.
- Medium term: 30 days prior up to and including the cohort start date.

However, the user can change these values. For example:

```
settings <- createCovariateSettings(useConditionEraLongTerm = TRUE,  
                                   useConditionEraShortTerm = TRUE,  
                                   useDrugEraLongTerm = TRUE,  
                                   useDrugEraShortTerm = TRUE,  
                                   longTermStartDays = -180,
```

```
shortTermStartDays = -14,
endDays = -1)
```

This redefines the long term window as 180 days prior up to (but not including) the cohort start date, and redefines the short term window as 14 days prior up to (but not including) the cohort start date.

Again, we can also specify which concept IDs should or should not be used to construct covariates:

```
settings <- createCovariateSettings(useConditionEraLongTerm = TRUE,
                                   useConditionEraShortTerm = TRUE,
                                   useDrugEraLongTerm = TRUE,
                                   useDrugEraShortTerm = TRUE,
                                   longTermStartDays = -180,
                                   shortTermStartDays = -14,
                                   endDays = -1,
                                   excludedCovariateConceptIds = 1124300,
                                   addDescendantsToExclude = TRUE)
```

2.3 Creating a set of custom covariates

This option should only be used by **advanced users**. It requires one to understand that at the implementation level, an analysis is a combination of a piece of highly parameterized SQL together with a specification of the parameter values. The best way to understand the available options is to take a prespecified analysis as starting point, and convert it to a detailed setting object:

```
settings <- createCovariateSettings(useConditionEraLongTerm = TRUE)
settings2 <- convertPrespecSettingsToDetailedSettings(settings)
settings2$analyses[[1]]
```

```
## $analysisId
## [1] 202
##
## $sqlFileName
## [1] "DomainConcept.sql"
##
## $parameters
## $parameters$analysisId
## [1] 202
##
## $parameters$analysisName
## [1] "ConditionEraLongTerm"
##
## $parameters$startDay
## [1] -365
##
## $parameters$endDay
## [1] 0
##
## $parameters$subType
## [1] "all"
##
## $parameters$domainId
## [1] "Condition"
##
```

```
## $parameters$domainTable
## [1] "condition_era"
##
## $parameters$domainConceptId
## [1] "condition_concept_id"
##
## $parameters$domainStartDate
## [1] "condition_era_start_date"
##
## $parameters$domainEndDate
## [1] "condition_era_end_date"
##
## $parameters$description
## [1] "One covariate per condition in the condition_era table overlapping with any part of the long te
##
##
## $includedCovariateConceptIds
## list()
##
## $includedCovariateIds
## list()
##
## $addDescendantsToInclude
## [1] FALSE
##
## $excludedCovariateConceptIds
## list()
##
## $addDescendantsToExclude
## [1] FALSE
```

One can create a detailed analysis settings object from scratch, and use it to create a detailed settings object:

```
analysisDetails <- createAnalysisDetails(analysisId = 1,
                                       sqlFileName = "DemographicsGender.sql",
                                       parameters = list(analysisId = 1,
                                                         analysisName = "Gender",
                                                         domainId = "Demographics"),
                                       includedCovariateConceptIds = c(),
                                       addDescendantsToInclude = FALSE,
                                       excludedCovariateConceptIds = c(),
                                       addDescendantsToExclude = FALSE,
                                       includedCovariateIds = c())

settings <- createDetailedCovariateSettings(list(analysisDetails))
```

2.4 Temporal covariates

Ordinarily, covariates are created for just a few time windows of interest, for example the short, medium, and long term windows described earlier. However, sometimes a more fine-grained temporal resolution is required, for example creating covariates for each day separately, in the 365 days prior to cohort start. We will refer to this type of covariates as *temporal covariates*. Temporal covariates share the same covariate ID across the time windows, and use a separate time ID to distinguish between time windows. There currently aren't many applications that are able to handle temporal covariates. For example, the `CohortMethod` package will

break when provided with temporal covariates. However, there are some machine learning algorithms in the `PatientLevelPrediction` package that require temporal covariates.

Again, we can just choose to use the default settings:

```
settings <- createDefaultTemporalCovariateSettings()
```

Or, we can choose from a set of prespecified temporal covariates:

```
settings <- createTemporalCovariateSettings(useConditionOccurrence = TRUE,  
                                           useMeasurementValue = TRUE)
```

In this case we've chosen to create binary covariates for each concept in the `condition_occurrence` table, and continuous covariates for each measurement - unit combination in the `measurement` table in the CDM. By default, temporal covariates are created for each day separately in the 365 days before (but not including) the cohort start date. Different time windows can also be specified, for example creating 7 day intervals instead:

```
settings <- createTemporalCovariateSettings(useConditionOccurrence = TRUE,  
                                           useMeasurementValue = TRUE,  
                                           temporalStartDays = seq(-364, -7, by = 7),  
                                           temporalEndDays = seq(-358, -1, by = 7))
```

Each time window includes the specified start and end day.

Similar to ordinary covariates, **advanced users** can also define custom analyses:

```
analysisDetails <- createAnalysisDetails(analysisId = 1,  
                                       sqlFileName = "MeasurementValue.sql",  
                                       parameters = list(analysisId = 1,  
                                                         analysisName = "MeasurementValue",  
                                                         domainId = "Measurement"),  
                                       includedCovariateConceptIds = c(),  
                                       addDescendantsToInclude = FALSE,  
                                       excludedCovariateConceptIds = c(),  
                                       addDescendantsToExclude = FALSE,  
                                       includedCovariateIds = c())  
  
settings <- createDetailedTemporalCovariateSettings(list(analysisDetails))
```

3 Constructing covariates for a cohort of interest

Here we will walk through an example, creating covariates for two cohorts of interest: new users of diclofenac and new users of celecoxib.

3.1 Configuring the connection to the server

We need to tell R how to connect to the server where the data are. `CohortMethod` uses the `DatabaseConnector` package, which provides the `createConnectionDetails` function. Type `?createConnectionDetails` for the specific settings required for the various database management systems (DBMS). For example, one might connect to a PostgreSQL database using this code:

```
connectionDetails <- createConnectionDetails(dbms = "postgresql",  
                                             server = "localhost/ohdsi",  
                                             user = "joe",
```

```

password = "supersecret")

cdmDatabaseSchema <- "my_cdm_data"
resultsDatabaseSchema <- "my_results"

```

The last two lines define the `cdmDatabaseSchema` and `resultSchema` variables. We'll use these later to tell R where the data in CDM format live, and where we want to write intermediate and result tables. Note that for Microsoft SQL Server, databaseschemas need to specify both the database and the schema, so for example `cdmDatabaseSchema <- "my_cdm_data.dbo"`.

3.2 Creating a cohort of interest

FeatureExtraction requires the cohorts to be instantiated in the `cohort` table in the Common Data Model, or in a table that has the same structure as the `cohort` table. We could create cohorts using a cohort definition tool, but here we'll just use some simple SQL to find the first drug era per person. Note that because we will be creating covariates based on data before cohort start, we are requiring 365 days of observation before the first exposure. FeatureExtraction will not check if a subject is observed during the specified time windows.

```

/*****
File cohortsOfInterest.sql
*****/

IF OBJECT_ID('@resultsDatabaseSchema.cohorts_of_interest', 'U') IS NOT NULL
  DROP TABLE @resultsDatabaseSchema.cohorts_of_interest;

SELECT first_use.*
INTO @resultsDatabaseSchema.cohorts_of_interest
FROM (
  SELECT drug_concept_id AS cohort_definition_id,
         MIN(drug_era_start_date) AS cohort_start_date,
         MIN(drug_era_end_date) AS cohort_end_date,
         person_id
  FROM @cdmDatabaseSchema.drug_era
  WHERE drug_concept_id = 1118084 -- celecoxib
         OR drug_concept_id = 1124300 -- diclofenac
  GROUP BY drug_concept_id,
         person_id
) first_use
INNER JOIN @cdmDatabaseSchema.observation_period
  ON first_use.person_id = observation_period.person_id
  AND cohort_start_date >= observation_period_start_date
  AND cohort_end_date <= observation_period_end_date
WHERE DATEDIFF(DAY, observation_period_start_date, cohort_start_date) >= 365;

```

This is parameterized SQL which can be used by the `SqlRender` package. We use parameterized SQL so we do not have to pre-specify the names of the CDM and result schemas. That way, if we want to run the SQL on a different schema, we only need to change the parameter values; we do not have to change the SQL code. By also making use of translation functionality in `SqlRender`, we can make sure the SQL code can be run in many different environments.

```

library(SqlRender)
sql <- readSql("cohortsOfInterest.sql")
sql <- renderSql(sql,
                 cdmDatabaseSchema = cdmDatabaseSchema,

```

```

        resultsDatabaseSchema = resultsDatabaseSchema)$sql
sql <- translateSql(sql, targetDialect = connectionDetails$dbms)$sql

connection <- connect(connectionDetails)
executeSql(connection, sql)

```

In this code, we first read the SQL from the file into memory. In the next line, we replace the two parameter names with the actual values. We then translate the SQL into the dialect appropriate for the DBMS we already specified in the `connectionDetails`. Next, we connect to the server, and submit the rendered and translated SQL.

If all went well, we now have a table with the cohorts of interest. We can see how many events per type:

```

sql <- paste("SELECT cohort_definition_id, COUNT(*) AS count",
            "FROM @resultsDatabaseSchema.cohorts_of_interest",
            "GROUP BY cohort_definition_id")
sql <- renderSql(sql, resultsDatabaseSchema = resultsDatabaseSchema)$sql
sql <- translateSql(sql, targetDialect = connectionDetails$dbms)$sql

querySql(connection, sql)

```

```

## cohort_concept_id count
## 1          1124300 240761
## 2          1118084  47293

```

3.3 Creating per-person covariates for a cohort of interest

We can create per-person covariates for one of the cohorts of interest, for example using the default settings:

```

covariateSettings <- createDefaultCovariateSettings()

covariateData <- getDbCovariateData(connectionDetails = connectionDetails,
                                   cdmDatabaseSchema = cdmDatabaseSchema,
                                   cohortDatabaseSchema = resultsDatabaseSchema,
                                   cohortTable = "cohorts_of_interest",
                                   cohortId = 1118084,
                                   rowIdField = "subject_id",
                                   covariateSettings = covariateSettings)

summary(covariateData)

```

```

## CovariateData object summary
##
## Number of covariates: 41330
## Number of non-zero covariate values: 25892630

```

3.3.1 Per-person covariate output format

The main component of the `covariateData` object is `covariates`:

```

covariateData$covariates

## ffd f (all open) dim=c(25892630,3), dimorder=c(1,2) row.names=NULL
## ffd virtual mapping
## PhysicalName VirtualVmode PhysicalVmode AsIs VirtualIsMatrix PhysicalIsMatrix PhysicalIsMatrix

```

## rowId	rowId	integer	integer FALSE	FALSE	FALSE
## covariateId	list...1	double	double FALSE	FALSE	FALSE
## covariateValue	list...2	double	double FALSE	FALSE	FALSE

ffd data

##	rowId	covariateId	covariateValue
## 1	1	4150129212	1
## 2	2	4211231212	1
## 3	3	4122924212	1
## 4	4	436785212	1
## 5	5	4231363212	1
## 6	6	4031662212	1
## 7	7	37016775212	1
## 8	8	4180170212	1
## :	:	:	:
## 25892623	25892623	1903	1
## 25892624	25892624	1903	3
## 25892625	25892625	1903	1
## 25892626	25892626	1903	1
## 25892627	25892627	1903	2
## 25892628	25892628	1903	2
## 25892629	25892629	1903	2
## 25892630	25892630	1903	2

The columns are defines as follows:

- **rowId** uniquely identifies a cohort entry. When calling `getDbCovariateData` we defined `rowIdField = "subject_id"`, so in this case the `rowId` is the same as the `subject_id` in the cohort table. In cases where a single subject can appear in the cohort more than once it is up to the user to create a field in the cohort table that uniquely identifies each cohort entry, and use that as `rowIdField`.
- **covariateId** identifies the covariate, and definitions of covariates can be found in the `cohortData$covariateRef` object.
- **covariateValue** field provides the value.

3.3.2 Saving the data to file

Creating covariates can take considerable computing time, and it is probably a good idea to save it for future sessions. Because `covariateData` objects use `ff`, we cannot use R's regular save function. Instead, we'll have to use the `saveCovariateData()` function:

```
saveCovariateData(covariateData, "covariates")
```

We can use the `loadCovariateData()` function to load the data in a future session.

3.3.3 Normalizing and removing redundancy

One reason for generating per-person covariates may be to use them in some form of machine learning. In that case it may be necessary to tidy the data before proceeding. The `tidyCovariateData` function can perform two tasks:

1. **Normalization:** scale all covariate values to a value between 0 and 1 (by dividing by the max value for each covariate).
2. **Removal of redundancy:** If every person in the cohort has the same value for a covariate (e.g. a cohort that is restricted to women will have the same gender covariate value for all) that value is redundant. Redundant values may wreak havoc with some machine learning algorithms, for example causing a simple regression to no longer have a single solution. Similarly, groups of covariates may be

redundant (e.g. every person will belong to at least one age group, making one group redundant as it can be defined as the absence of the other groups).

```
tidyCovariates <- tidyCovariateData(covariateData,
                                   normalize = TRUE,
                                   removeRedundancy = TRUE)
```

If we want to know which redundant covariates were removed we can query the `metaData` object:

```
deletedCovariateIds <- tidyCovariates$metaData$deletedCovariateIds
deletedCovariateIds
```

```
## [1] 1118084410 1118084412 1118084413 21603931410 21603931412 21603931413 21603932410 21603932412
## [16] 900000010802 8532001 10003 8527004 2015006 1007
```

If we want to know what these numbers mean, we can use the `covariateRef` object that is part of any `covariateData` object. Because `covariateRef`, like other parts of `covariateData`, uses the `ff` package, it is advised to load the `ffbase` package which allows `ff` objects to be used as data frames:

```
library(ffbase)
covariateData$covariateRef[covariateData$covariateRef$covariateId %in% deletedCovariateIds, ]
```

```
##      covariateId      covariateName analysisId conceptId
## 1      8527004      race = White           4      8527
## 2      8532001      gender = FEMALE        1      8532
## 3      2015006      index year: 2015        6         0
## 4       1007      index month: 1            7         0
## 5       10003      age group: 50-51          3         0
## 6 900000010802 ...tance Abuse Coverage Indicator 802 900000010
## 7 21603933412 ...EUMATIC PRODUCTS, NON-STEROIDS 412 21603933
## 8 21603932412 ...ORY AND ANTIRHEUMATIC PRODUCTS 412 21603932
## 9 21603932410 ...ORY AND ANTIRHEUMATIC PRODUCTS 410 21603932
## 10 1118084412 ...s relative to index: celecoxib 412 1118084
## 11 1118084410 ...s relative to index: celecoxib 410 1118084
## 12 21603931412 ...index: MUSCULO-SKELETAL SYSTEM 412 21603931
## 13 21603991410 ...days relative to index: Coxibs 410 21603991
## 14 21603991412 ...days relative to index: Coxibs 412 21603991
## 15 21603931410 ...index: MUSCULO-SKELETAL SYSTEM 410 21603931
## 16 21603933410 ...EUMATIC PRODUCTS, NON-STEROIDS 410 21603933
## 17 21603933413 ...EUMATIC PRODUCTS, NON-STEROIDS 413 21603933
## 18 21603932413 ...ORY AND ANTIRHEUMATIC PRODUCTS 413 21603932
## 19 1118084413 ...s relative to index: celecoxib 413 1118084
## 20 21603991413 ...days relative to index: Coxibs 413 21603991
## 21 21603931413 ...index: MUSCULO-SKELETAL SYSTEM 413 21603931
```

3.4 Creating aggregated covariates for a cohort of interest

Often we do not need to have per-person covariates, but instead we are interested in aggregated statistics instead. For example, we may not need to know which persons are male, but would like to know what proportion of the cohort is male. We can aggregate per-person covariates:

```
covariateData2 <- aggregateCovariates(covariateData)
```

Of course, if all we wanted was aggregated statistics it would have been more efficient to aggregate them during creation:

```

covariateSettings <- createDefaultCovariateSettings()

covariateData2 <- getDbCovariateData(connectionDetails = connectionDetails,
                                     cdmDatabaseSchema = cdmDatabaseSchema,
                                     cohortDatabaseSchema = resultsDatabaseSchema,
                                     cohortTable = "cohorts_of_interest",
                                     cohortId = 1118084,
                                     covariateSettings = covariateSettings,
                                     aggregated = TRUE)

summary(covariateData2)

```

```

## CovariateData object summary
##
## Number of covariates: 41330
## Number of non-zero covariate values: 41330

```

Note that we specified `aggregated = TRUE`. Also, we are no longer required to define a `rowIdField` because we will no longer receive per-person data.

3.4.1 Aggregated covariate output format

The two main components of the aggregated `covariateData` object are `covariates` and `covariatesContinuous`, for binary and continous covariates respectively:

```
covariateData2$covariates
```

```

## ffd (all open) dim=c(41326,3), dimorder=c(1,2) row.names=NULL
## ffd virtual mapping
##
##      PhysicalName VirtualVmode PhysicalVmode AsIs VirtualIsMatrix PhysicalIsMatrix Physical
## covariateId      list..      double      double FALSE              FALSE              FALSE
## sumValue         list...1      double      double FALSE              FALSE              FALSE
## averageValue     list...2      double      double FALSE              FALSE              FALSE
## ffd data
##      covariateId      sumValue averageValue
## 1      4.078214e+09 2.000000e+00 4.228956e-05
## 2      4.368402e+08 1.000000e+00 2.114478e-05
## 3      1.388962e+08 1.400000e+01 2.960269e-04
## 4      3.734882e+08 1.000000e+00 2.114478e-05
## 5      4.328982e+08 3.900000e+01 8.246464e-04
## 6      4.340782e+08 1.000000e+00 2.114478e-05
## 7      4.331342e+08 1.000000e+01 2.114478e-04
## 8      4.394172e+08 1.000000e+00 2.114478e-05
## :      :      :      :
## 41319 2.314271e+09 1.970000e+02 4.165521e-03
## 41320 2.314285e+09 1.331000e+03 2.814370e-02
## 41321 4.075713e+10 5.000000e+00 1.057239e-04
## 41322 4.274253e+10 1.000000e+00 2.114478e-05
## 41323 4.481643e+10 2.000000e+00 4.228956e-05
## 41324 4.625748e+10 1.000000e+00 2.114478e-05
## 41325 4.625749e+10 7.600000e+01 1.607003e-03
## 41326 4.625753e+10 1.160000e+02 2.452794e-03

```

```
covariateData2$covariatesContinuous
```

```
## ffd (all open) dim=c(4,11), dimorder=c(1,2) row.names=NULL
```

```
## ffdv virtual mapping
##           PhysicalName VirtualVmode PhysicalVmode  AsIs VirtualIsMatrix PhysicalIsMatrix Phy
## covariateId      list..      double      double FALSE              FALSE              FALSE
## countValue       list...1     double      double FALSE              FALSE              FALSE
## minValue         list...2     double      double FALSE              FALSE              FALSE
## maxValue         list...3     double      double FALSE              FALSE              FALSE
## averageValue     list...4     double      double FALSE              FALSE              FALSE
## standardDeviation list...5     double      double FALSE              FALSE              FALSE
## medianValue      list...6     double      double FALSE              FALSE              FALSE
## p10Value         list...7     double      double FALSE              FALSE              FALSE
## p25Value         list...8     double      double FALSE              FALSE              FALSE
## p75Value         list...9     double      double FALSE              FALSE              FALSE
## p90Value         list...10    double      double FALSE              FALSE              FALSE
## ffdv data
##   covariateId  countValue  minValue  maxValue  averageValue  standardDeviation  medianValue
## 1  1901.000000 33013.000000  0.000000  21.000000   2.321168         2.657008     1.000000
## 2  1904.000000 31301.000000  0.000000   6.000000   1.307128         1.144543     1.000000
## 3  1902.000000 27951.000000  0.000000  13.000000   2.646671         2.723938     1.000000
## 4  1903.000000 30898.000000  0.000000   5.000000   1.234157         1.055174     1.000000
```

The columns are defines as follows:

- `covariateId` identifies the covariate, and definitions of covariates can be found in the `cohortData$covariateRef` object.
- `sumValue` is the sum of the covariate values. Because these are binary features, this is equivalent to the number of people that have the covariate with a value of 1.
- `averageValue` is the average covariate value. Because these are binary features, this is equivalent to the proportion of people that have the covariate with a value of 1.
- `countValue` is the number of people that have a value (for continous variables).
- `minValue`, `maxValue`, `averageValue`, `standardDeviation`, `medianValue`, `p10Value`, `p25Value`, `p75Value`, and `p90Value` all inform on the distribution of covariate values. Note that for some covariates (such as the Charlson comorbidity index) a value of 0 is interpreted as the value 0, while for other covariates (Such as blood pressure) 0 is interpreted as missing, and the distribution statistics are only computed over non-missing values. To learn which continuous covariates fall into which category one can consult the `missingMeansZero` field in the `covariateData$analysisRef` object.

3.5 Creating a table 1

One task supported by the `FeatureExtraction` package is creating a table of overall study population characteristics that can be include in a paper. Since this is typically the first table in a paper we refer to such a table as ‘table 1’. A default table 1 is available in the `FeatureExtraction` package:

```
result <- createTable1(covariateData2)
print(result, row.names = FALSE, right = FALSE)
```

Characteristic	% (n = 47,293)	Characteristic	% (n = 47,293)
Age group		Heart failure	7
0-1	0	Ischemic heart disease	7
5-6	0	Peripheral vascular disease	13
10-11	1	Pulmonary embolism	1
15-16	2	Venous thrombosis	3
20-21	2	Medical history: Neoplasms	
25-26	4	Hematologic neoplasm	1
30-31	6	Malignant lymphoma	0
35-36	7	Malignant neoplasm of anorectum	0
40-41	9	Malignant neoplastic disease	6
45-46	11	Malignant tumor of breast	2
50-51	14	Malignant tumor of colon	0
55-56	13	Malignant tumor of lung	0
60-61	11	Malignant tumor of urinary bladder	0
65-66	6	Primary malignant neoplasm of prostate	0
70-71	4	Medication use	
75-76	3	Agents acting on the renin-angiotensin system	47
80-81	3	Antibacterials for systemic use	83
85-86	2	Antidepressants	64

90-91	0	Antiepileptics	77
Gender: female	72	Antiinflammatory and antirheumatic products	100
Race		Antineoplastic agents	27
race = Black or African American	19	Antipsoriatics	9
race = Other Race	19	Antithrombotic agents	22
race = White	62	Beta blocking agents	54
Ethnicity		Calcium channel blockers	48
ethnicity = Hispanic or Latino	2	Diuretics	63
ethnicity = Not Hispanic or Latino	17	Drugs for acid related disorders	83
Medical history: General		Drugs for obstructive airway diseases	60
Acute respiratory disease	37	Drugs used in diabetes	31
Attention deficit hyperactivity disorder	2	Immunosuppressants	4
Chronic liver disease	5	Lipid modifying agents	55
Chronic obstructive lung disease	19	Opioids	80
Crohn's disease	1	Psycholeptics	86
Dementia	2	Psychostimulants, agents used for adhd and nootropics	78
Depressive disorder	34		
Diabetes mellitus	25	Characteristic	Value
Gastroesophageal reflux disease	25	Charlson comorbidity index	
Gastrointestinal hemorrhage	4	Mean	2
Human immunodeficiency virus infection	1	Minimum	0
Hyperlipidemia	36	25th percentile	0
Hypertensive disorder	50	Median	1
Lesion of liver	1	75th percentile	3
Obesity	17	Maximum	21
Osteoarthritis	45	CHADS2Vasc	
Pneumonia	6	Mean	1
Psoriasis	1	Minimum	0
Renal impairment	5	25th percentile	0
Rheumatoid arthritis	5	Median	1
Schizophrenia	4	75th percentile	2
Ulcerative colitis	0	Maximum	6
Urinary tract infectious disease	15	DCSI	
Viral hepatitis C	4	Mean	3
Visual system disorder	36	Minimum	0
Medical history: Cardiovascular disease		25th percentile	0
Atrial fibrillation	3	Median	1
Cerebrovascular disease	6	75th percentile	5
Coronary arteriosclerosis	10	Maximum	13
Heart disease	26		

Where applicable, these characteristics are drawn from analyses pertaining the ‘long-term’ windows, so concepts observed in the 365 days before up to and included the cohort start date.

The `createTable1` function requires a simple specification of what variables to include in the table. The default specifications included in the package can be reviewed by calling the `getDefaultTable1Specifications` function. The specification reference analysis IDs and covariate IDs, and in the default specification these IDs refer to those in the default covariate settings. It is possible to create custom table 1 specifications and use those instead.

Here we based table 1 on a `covariateData` object containing all default covariates, even though only a small fraction of covariates are used in the table. If we only want to extract those covariates needed for the table, we can use the `createTable1CovariateSettings` function:

```

covariateSettings <- createTable1CovariateSettings()

covariateData2b <- getDbCovariateData(connectionDetails = connectionDetails,
                                       cdmDatabaseSchema = cdmDatabaseSchema,
                                       cohortDatabaseSchema = resultsDatabaseSchema,
                                       cohortTable = "cohorts_of_interest",
                                       cohortId = 1118084,
                                       covariateSettings = covariateSettings,
                                       aggregated = TRUE)

summary(covariateData2b)

```

```

## CovariateData object summary
##
## Number of covariates: 90
## Number of non-zero covariate values: 90

```

3.6 Comparing two cohorts

Another task supported by the `FeatureExtraction` package is comparing two cohorts of interest. Suppose we want to compare two cohorts only on the variables included in the default table 1:

```

settings <- createTable1CovariateSettings(excludedCovariateConceptIds = c(1118084, 1124300),
                                         addDescendantsToExclude = TRUE)

covCelecoxib <- getDbCovariateData(connectionDetails = connectionDetails,
                                   cdmDatabaseSchema = cdmDatabaseSchema,
                                   cohortDatabaseSchema = resultsDatabaseSchema,
                                   cohortTable = "cohorts_of_interest",
                                   cohortId = 1118084,
                                   covariateSettings = settings,
                                   aggregated = TRUE)

covDiclofenac <- getDbCovariateData(connectionDetails = connectionDetails,
                                    cdmDatabaseSchema = cdmDatabaseSchema,
                                    cohortDatabaseSchema = resultsDatabaseSchema,
                                    cohortTable = "cohorts_of_interest",
                                    cohortId = 1124300,
                                    covariateSettings = settings,
                                    aggregated = TRUE)

std <- computeStandardizedDifference(covCelecoxib, covDiclofenac)

```

In this example we have chosen to exclude any covariates derived from the two concepts that were used to define the two cohorts: celecoxib (1118084), and diclofenac (1124300). We compute the standardized difference between the remaining covariates.

```
head(std)
```

```

##      covariateId      mean1      sd1      mean2      sd2      sd      stdDiff      covariateName
## 90          1904  1.30712790  1.1445434  0.9100477  1.0777444  1.5721045 -0.2525787      CHADS2Vasc
## 4           3003  0.02404161  0.1550552  0.1183165  0.3439724  0.3773051  0.2498638      age group: 15-16
## 24      80180210  0.44879792  0.6699309  0.2619693  0.5118305  0.8430764 -0.2216034 ...ative to index: Osteoarthritis
## 89          1902  2.64667075  2.7239376  1.8278085  2.6827256  3.8231993 -0.2141825 ...orbidity Severity Index (DCSI)
## 88          1901  2.32116804  2.6570083  1.6216871  2.5198808  3.6618974 -0.1910160 ...lson index - Romano adaptation
## 74  21601853410  0.48421542  0.6958632  0.3182368  0.5641259  0.8958034 -0.1852847 ... index: LIPID MODIFYING AGENTS

```

The `stdDiff` column contains the standardized difference. By default the data is ranked in descending order of the absolute value of the standardized difference, showing the covariate with the largest difference first.

We can also show the comparison as a standard table 1:

```

result <- createTable1(covCelecoxib, covDiclofenac)
print(result, row.names = FALSE, right = FALSE)

```

Characteristic	% (n = 47,293)	% (n = 240,761)	Std.Diff	Characteristic	% (n = 47,293)	% (n = 240,761)	Std.Diff
Age group				Heart failure	7	4	-0.09
0-1	0	0	-0.01	Ischemic heart disease	7	4	-0.09
5-6	0	0	0.00	Peripheral vascular disease	13	8	-0.12
10-11	1	3	0.11	Pulmonary embolism	1	1	-0.05
15-16	2	12	0.25	Venous thromboemb	3	2	-0.06
20-21	2	6	0.14	Medical history: Neoplasms			
25-26	4	9	0.14	Hematologic neoplasm	1	1	-0.04
30-31	6	10	0.11	Malignant lymphoma	0	0	-0.02
35-36	7	10	0.07	Malignant neoplasm of anorectum	0	0	-0.02
40-41	9	9	0.00	Malignant neoplastic disease	6	4	-0.08
45-46	11	9	-0.05	Malignant tumor of breast	2	1	-0.04
50-51	14	10	-0.09	Malignant tumor of colon	0	0	-0.02
55-56	13	9	-0.09	Malignant tumor of lung	0	0	-0.03
60-61	11	6	-0.13	Malignant tumor of urinary bladder	0	0	-0.02
65-66	6	2	-0.12	Primary malignant neoplasm of prostate	0	0	-0.02
70-71	4	2	-0.11	Medication use			
75-76	3	1	-0.10	Agents acting on the renin-angiotensin system	47	33	-0.16
80-81	3	1	-0.10	Antibacterials for systemic use	81	80	0.00
85-86	2	1	-0.07	Antidepressants	64	53	-0.10
90-91	0	0	-0.01	Antiepileptics	77	74	-0.03
Gender: female	72	73	0.00	Antiinflammatory and antirheumatic products	73	72	-0.01
Race				Antineoplastic agents	17	13	-0.08
race = Black or African American	19	28	0.14	Antipsoriatics	9	7	-0.05
race = Other Race	19	15	-0.06	Antithrombotic agents	22	11	-0.18
race = White	62	57	-0.05	Beta blocking agents	54	40	-0.15
Ethnicity				Calcium channel blockers	48	32	-0.17
ethnicity = Hispanic or Latino	2	2	0.03	Diuretics	63	48	-0.14
ethnicity = Not Hispanic or Latino	17	13	-0.08	Drugs for acid related disorders	83	80	-0.03
Medical history: General				Drugs for obstructive airway diseases	60	57	-0.02
Acute respiratory disease	37	39	0.03	Drugs used in diabetes	31	21	-0.14
Attention deficit hyperactivity disorder	2	4	0.07	Immunosuppressants	4	2	-0.07
Chronic liver disease	5	4	-0.05	Lipid modifying agents	48	32	-0.19
Chronic obstructive lung disease	19	11	-0.14	Opioids	80	79	0.00
Crohn's disease	1	0	-0.02	Psycholeptics	86	81	-0.04
Dementia	2	1	-0.06	Psychostimulants, agents used for adhd and nootropics	78	78	0.00
Depressive disorder	34	29	-0.06				
Diabetes mellitus	25	17	-0.12	Characteristic	Value	Value	Std.Diff

Gastroesophageal reflux disease	25	19	-0.09	Charlson comorbidity index			
Gastrointestinal hemorrhage	4	3	-0.04	Mean	2	2	-0.19
Human immunodeficiency virus infection	1	1	0.01	Minimum	0	0	
Hyperlipidemia	36	25	-0.14	25th percentile	0	0	
Hypertensive disorder	50	37	-0.14	Median	1	1	
Lesion of liver	1	0	-0.02	75th percentile	3	2	
Obesity	17	17	-0.01	Maximum	21	27	
Osteoarthritis	45	26	-0.22	CHADS2Vasc			
Pneumonia	6	4	-0.07	Mean	1	1	-0.25
Psoriasis	1	1	-0.02	Minimum	0	0	
Renal impairment	5	3	-0.05	25th percentile	0	0	
Rheumatoid arthritis	5	2	-0.09	Median	1	0	
Schizophrenia	4	3	-0.04	75th percentile	2	1	
Ulcerative colitis	0	0	-0.02	Maximum	6	6	
Urinary tract infectious disease	15	16	0.01	DCSI			
Viral hepatitis C	4	3	-0.04	Mean	3	2	-0.21
Visual system disorder	36	31	-0.06	Minimum	0	0	
Medical history: Cardiovascular disease				25th percentile	0	0	
Atrial fibrillation	3	1	-0.06	Median	1	0	
Cerebrovascular disease	6	4	-0.06	75th percentile	5	3	
Coronary arteriosclerosis	10	5	-0.11	Maximum	13	13	
Heart disease	26	17	-0.15				