Building patient-level predictive models

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Warning: package 'rJava' was built under R version 3.2.1

1 Introduction

This vignette describes how you can use the PatientLevelPrediction package to build patient-level prediction models. We will walk through all the steps needed to build an exemplar model, and we have selected the well-studied topic of predicting re-hospitalization. To reduce the scope a bit, we have limited ourselves to a diabetes type 2 population.

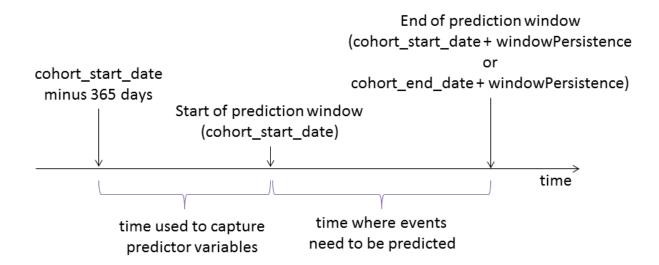
1.1 Specifying the cohort of interest and outcomes

The PatientLevelPrediction package requires longitudinal observational healthcare data in the OMOP Common Data Model format. The user will need to specify two things:

- 1. Time periods for which we wish to predict the occurrence of an outcome. We will call this the **cohort of interest** or cohort for short. One person can have multiple time periods, but time periods should not overlap.
- 2. Outcomes for which we wish to build a predictive model.

The cohort and outcomes should be provided as data in a table on the server, where the table should have the same structure as the cohort table in the OMOP CDM, meaning it should have the following columns:

- cohort_concept_id (CDM v4) or cohort_concept_id (CDM v5+), a unique identifier for distinguishing between different types of cohorts, e.g. cohorts of interest and outcome cohorts.
- subject_id, a unique identifier corresponding to the person_id in the CDM.
- cohort_start_date, the start of the time period where we wish to predict the occurrence of the outcome.
- cohort_end_date, which can be used to determine the end of the prediction window. Can be NULL for outcomes.



The prediction window always starts on the cohort_end_date. When the useCohortEndDate argument is set to TRUE, the prediction window will end on the cohort_end_date plus the number of days specified using the windowPersistence argument. If the useCohortEndDate argument is set to FALSE, the prediction window will end on the cohort_start_date plus the number of days specified using the windowPersistence argument.

The package will use data from the time period preceding (and including) the cohort_start_date to build a large set of features that can be used to predict outcomes during the prediction window (including the first and last day of the prediction window). These features can include binary indicators for the occurrence of any individual drug, condition, procedures, as well as demographics and comorbidity indices.

Important: Currently, PatientLevelPrediction does not check or limit the input data in any way. It is up to the user to make sure there are at least 365 days of observation time preceding the cohort_start_date for the cohort of interest. Furthermore, since the cohort_start_date is included in both the time used to construct predictors, and the time when outcomes can occurr, it is up to the user to either remove outcomes occurring on the cohort_start_date, or to remove predictors that are in fact indicators of the occurrence of an outcome on the cohort_start_date.

2 Installation instructions

Before installing the PatientLevelPrediction package make sure you have Java available. Java can be downloaded from www.java.com. For Windows users, RTools is also necessary. RTools can be downloaded from CRAN.

The PatientLevelPrediction package is currently maintained in a Github repository, and has dependencies on other packages in Github. All of these packages can be downloaded and installed from within R using the devtools package:

```
install.packages("devtools")
library(devtools)
install_github("ohdsi/SqlRender")
install_github("ohdsi/DatabaseConnector")
install_github("ohdsi/Cyclops")
install_github("ohdsi/PatientLevelPrediction")
```

Once installed, you can type library (PatientLevelPrediction) to load the package.

3 Data extraction

The first step in running the PatientLevelPrediction is extracting all necessary data from the database server holding the data in the Common Data Model (CDM) format.

3.1 Configuring the connection to the server

We need to tell R how to connect to the server where the data are. PatientLevelPrediction 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:

The last three lines define the cdmDatabaseSchema and resultSchema variables, as well as the CDM version. We'll use these later to tell R where the data in CDM format live, where we want to write intermediate and result tables, and what version CDM is used. 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 Preparing the cohort and outcome of interest

Before we can start using the PatientLevelPrediction package itself, we need to construct a cohort of interest for which we want to perform the prediction, and the outcome, the event that we would like to predict. We do this by writing SQL statements against the CDM that populate a table containing the persons and events of interest. The resulting table should have the same structure as the cohort table in the CDM. For CDM v4, this means it should have the fields cohort_concept_id, cohort_start_date, cohort_end_date, and subject_id. For CDM v4, the cohort_concept_id field must be called cohort_definition_id.

For our example study, we need to create the cohort of diabetics that have been hospitalized and have a minimum amount of observation time available before and after the hospitalization. We also need to defined re-hospitalizations, which we define as any hospitalizations occurring after the original hospitalization.

For this purpose we have created a file called HospitalizationCohorts.sql with the following contents:

```
/**********
File HospitalizationCohorts.sql
************
IF OBJECT_ID('@resultsDatabaseSchema.rehospitalization', 'U') IS NOT NULL
DROP TABLE @resultsDatabaseSchema.rehospitalization;
SELECT visit_occurrence.person_id AS subject_id,
MIN(visit start date) AS cohort start date,
DATEADD(DAY, @post_time, MIN(visit_start_date)) AS cohort_end_date,
1 AS cohort_concept_id
INTO @resultsDatabaseSchema.rehospitalization
FROM @cdmDatabaseSchema.visit_occurrence
INNER JOIN @cdmDatabaseSchema.observation_period
ON visit_occurrence.person_id = observation_period.person_id
INNER JOIN @cdmDatabaseSchema.condition_occurrence
ON condition_occurrence.person_id = visit_occurrence.person_id
WHERE place_of_service_concept_id IN (9201, 9203)
AND DATEDIFF(DAY, observation period start date, visit start date) > @pre time
AND visit_start_date > observation_period_start_date
AND DATEDIFF(DAY, visit_start_date, observation_period_end_date) > @post_time
AND visit_start_date < observation_period_end_date</pre>
AND DATEDIFF(DAY, condition_start_date, visit_start_date) > @pre_time
AND condition_start_date <= visit_start_date
AND condition concept id IN (
SELECT descendant concept id
FROM @cdmDatabaseSchema.concept_ancestor
WHERE ancestor_concept_id = 201826) /* Type 2 DM */
GROUP BY visit_occurrence.person_id;
INSERT INTO @resultsDatabaseSchema.rehospitalization
SELECT visit_occurrence.person_id AS subject_id,
visit_start_date AS cohort_start_date,
visit_end_date AS cohort_end_date,
2 AS cohort_concept_id
FROM @resultsDatabaseSchema.rehospitalization
INNER JOIN @cdmDatabaseSchema.visit occurrence
ON visit_occurrence.person_id = rehospitalization.subject_id
WHERE place_of_service_concept_id IN (9201, 9203)
AND visit_start_date > cohort_start_date
AND visit_start_date <= cohort_end_date</pre>
AND cohort_concept_id = 1;
```

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.

In this code, we first read the SQL from the file into memory. In the next line, we replace four 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 events of interest. We can see how many events per type:

3.3 Extracting the data from the server

Now we can tell PatientLevelPrediction to extract all necessary data for our analysis:

```
cohortData <- getDbCohortData(connectionDetails$dbms,</pre>
                               cdmDatabaseSchema = cdmDatabaseSchema,
                               cohortDatabaseSchema = resultsDatabaseSchema,
                               cohortTable = "rehospitalization",
                               cohortIds = 1,
                               useCohortEndDate = TRUE,
                               windowPersistence = 0,
                               cdmVersion = cdmVersion)
covariateSettings <- createCovariateSettings(useCovariateDemographics = TRUE,</pre>
                                              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,
                                              useCovariateDrugEraOverlap = TRUE,
                                              useCovariateDrugEraEver = TRUE,
                                              useCovariateDrugGroup = TRUE,
                                              useCovariateProcedureOccurrence = TRUE,
                                              useCovariateProcedureOccurrence365d = TRUE,
                                              useCovariateProcedureOccurrence30d = TRUE,
                                              useCovariateProcedureGroup = TRUE,
                                              useCovariateObservation = TRUE,
                                              useCovariateObservation365d = TRUE,
                                              useCovariateObservation30d = TRUE,
                                              useCovariateObservationCount365d = TRUE,
                                              useCovariateMeasurement = 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 = 1)
covariateData <- getDbCovariateData(connectionDetails$dbms,</pre>
                                     cdmDatabaseSchema = cdmDatabaseSchema,
                                     cohortDatabaseSchema = resultsDatabaseSchema,
                                     cohortTable = "rehospitalization",
                                     cohortIds = 1,
                                     covariateSettings = covariateSettings,
                                     cdmVersion = cdmVersion)
outcomeData <- getDbOutcomeData(connectionDetails$dbms,</pre>
                                 cdmDatabaseSchema = cdmDatabaseSchema,
                                 cohortDatabaseSchema = resultsDatabaseSchema,
                                 cohortTable = "rehospitalization",
                                 cohortIds = 1,
                                useCohortEndDate = TRUE,
                                windowPersistence = 0,
                                outcomeDatabaseSchema = resultsDatabaseSchema,
                                outcomeTable = "rehospitalization",
                                 outcomeConceptIds = 2,
                                 cdmVersion = cdmVersion)
```

There are many parameters, but they are all documented in the PatientLevelPrediction manual. In short, we are pointing the functions to the table created earlier and indicating which concept IDs in that table

identify the cohort and outcome of interest. We construct cohortData, an object representing the cohort of interest, covariateData, holding all covariate information that will be used as predictors, and outcomeData as the set of outcomes during cohort time. All three objects use the package ff to store information in a way that ensures R does not run out of memory, even when the data are large.

Important: Make sure the same prediction window definition (e.g. useCohortEndDate and windowPersistence) is used when calling both getDbCohortData and getDbOutcomeData.

We can get some overall statistics using the generic summary() method:

```
summary(cohortData)
## CohortData object summary
##
     Cohort count Person count
## 1
           446375
                         446375
summary(covariateData)
## CovariateData object summary
##
## Number of covariates: 70539
## Number of non-zero covariate values: 591071684
summary(outcomeData)
## OutcomeData object summary
##
##
     Cohort count Person count
## 2
           118294
                         118294
```

3.3.1 Saving the data to file

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

```
saveCohortData(cohortData, "rehosp_cohorts")
saveCovariateData(covariateData, "rehosp_covariates")
saveOutcomeData(outcomeData, "rehosp_outcomes")
```

We can use the loadCohortData(), loadCovariateData(), and loadOutcomeData() function to load the data in a future session.

4 Fitting the model

4.1 Train-test split

We typically not only want to build our model, we also want to know how good it is. Because evaluation using the same data on which the model was fitted can lead to overestimation, one uses a train-test split of the data or cross-validation. We can use the splitData() function to split the data in a 75%-25% split:

```
parts <- splitData(cohortData, covariateData, outcomeData, c(0.75, 0.25))</pre>
```

The parts variable is a list, where each item holds the data for a subset of the full cohort of interest. For example, parts [[1]] \$cohortData contains the cohort data for 75% of the people in the cohort of interest. We can now fit the model on the first part of the data (the training set):

4.2 Fitting the model on the training data

```
model <- fitPredictiveModel(parts[[1]]$cohortData, parts[[1]]$covariateData,
    parts[[1]]$outcomeData, modelType = "logistic")</pre>
```

The fitPredictiveModel() function uses the Cyclops package to fit a large-scale regularized regression. To fit the model, Cyclops needs to know the hyperparameter value which specifies the variance of the prior. By default Cyclops will use cross-validation to estimate the optimal hyperparameter. However, be aware that this can take a really long time. You can use the prior and control parameters of the fitPredictiveModel() to specify Cyclops behaviour, including using multiple CPUs to speed-up the cross-validation.

4.3 Model evaluation

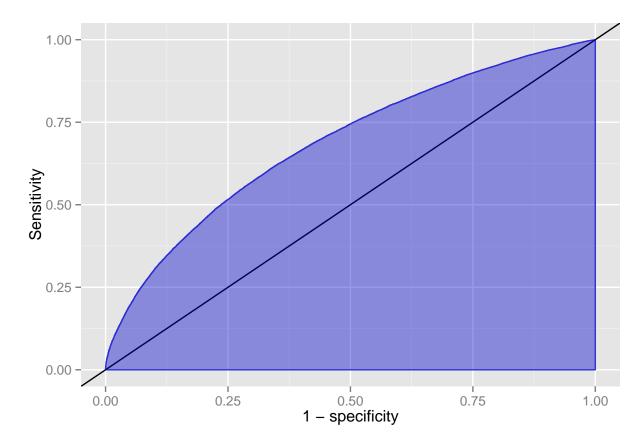
We can evaluate how well the model is able to predict the outcome, for example on the test data we can compute the area under the ROC curve:

prediction <- predictProbabilities(model, parts[[2]]\$cohortData, parts[[2]]\$covariateData)
computeAuc(prediction, parts[[2]]\$outcomeData)</pre>

[1] 0.6858641

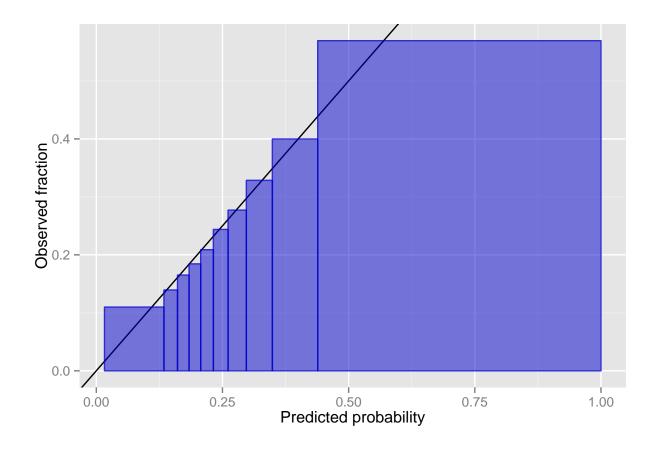
And we can plot the ROC curve itself:

plotRoc(prediction, parts[[2]]\$outcomeData)



We can also look at the calibration:

plotCalibration(prediction, parts[[2]]\$outcomeData, numberOfStrata = 10)



4.4 Inspecting the model

Now that we have some idea of the operating characteristics of our predictive model, we can investigate the model itself:

```
modelDetails <- getModelDetails(model, covariateData)
head(modelDetails)</pre>
```

```
##
             coefficient
                                  id
                                                              covariateName
## 0
              -1.0843708
                                                                  Intercept
## 4092289152 -0.5719409 4092289152 ...condition group:
                                                          4092289-Livebirth
             -0.4719253 4092289252 ...condition group:
## 4092289252
                                                          4092289-Livebirth
              -0.2873652
                          321042102 ...ohort index: 321042-Cardiac arrest
## 321042102
## 2313815702
              -0.2692642 2313815702 ..., without interpretation and report
               0.2530679 2110284702 ...dex: 2110284-Fetal non-stress test
## 2110284702
```

This shows the strongest predictors in the model with their corresponding betas.

5 Acknowledgments

Considerable work has been dedicated to provide the PatientLevelPrediction package.

citation("PatientLevelPrediction")

```
##
## To cite package 'PatientLevelPrediction' in publications use:
##
     Martijn J. Schuemie, Marc A. Suchard and Patrick B. Ryan (2015).
##
##
     PatientLevelPrediction: Package for patient level prediction
##
     using data in the OMOP Common Data Model. R package version
     0.0.3.
##
##
## A BibTeX entry for LaTeX users is
##
##
     @Manual{,
##
       title = {PatientLevelPrediction: Package for patient level prediction using data in the OMOP Com
       author = {Martijn J. Schuemie and Marc A. Suchard and Patrick B. Ryan},
##
##
       year = \{2015\},\
##
       note = {R package version 0.0.3},
##
     }
##
## ATTENTION: This citation information has been auto-generated from
## the package DESCRIPTION file and may need manual editing, see
## 'help("citation")'.
```

Further, PatientLevelPrediction makes extensive use of the Cyclops package.

citation("Cyclops")

```
##
## To cite Cyclops in publications use:
##
## Suchard MA, Simpson SE, Zorych I, Ryan P and Madigan D (2013).
## "Massive parallelization of serial inference algorithms for
## complex generalized linear models." _ACM Transactions on Modeling
## and Computer Simulation_, *23*, pp. 10. <URL:
## http://dl.acm.org/citation.cfm?id=2414791>.
## A BibTeX entry for LaTeX users is
##
##
     @Article{,
##
       author = {M. A. Suchard and S. E. Simpson and I. Zorych and P. Ryan and D. Madigan},
##
       title = {Massive parallelization of serial inference algorithms for complex generalized linear m
       journal = {ACM Transactions on Modeling and Computer Simulation},
##
##
       volume = \{23\},
##
       pages = \{10\},
##
       year = \{2013\},\
##
       url = {http://dl.acm.org/citation.cfm?id=2414791},
     }
##
```

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