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Estimating disease prevalence from drug utilization data using the Random Forest algorithm

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Background: Aggregated claims data on medication are often used as a proxy for the prevalence of diseases, especially chronic diseases. However, linkage between medication and diagnosis tend to be theory based and not very precise. Modelling disease probability at an individual level using individual level data may yield more accurate results. Methods: Individual probabilities of having a certain chronic disease were estimated using the Random Forest (RF) algorithm. A training set was created from a general practitioners database of 276,723 cases that included diagnosis and claims data on medication. Model performance for 29 chronic diseases was evaluated using Receiver Operating Curves, by measuring the Area Under the Curve (AUC). Results: The diseases for which model performance was best were Parkinson’s disease (AUC = .89, 95% CI = .77–1.00), diabetes (AUC = .87, 95% CI = .85–.90), osteoporosis (AUC = .87, 95% CI = .81–.92) and heart failure (AUC = .81, 95% CI = .74–.88). Five other diseases had an AUC >.75: asthma, chronic enteritis, COPD, epilepsy and HIV/AIDS. For 16 of 17 diseases tested, the medication categories used in theory-based algorithms were also identified by our method, however the RF models included a broader range of medications as important predictors. Conclusion: Data on medication use can be a useful predictor when estimating the prevalence of several chronic diseases. To improve the estimates, for a broader range of chronic diseases, research should use better training data, include more details concerning dosages and duration of prescriptions, and add related predictors like hospitalizations.

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Methods

Random Forest

Estimating the probability that an individual has a certain disease could be considered a mathematical classification problem. RF is a non-parametric method to address classification problems. For implementation the R-package ‘Random Forest’ was used.

Data

Drug use data of the entire Dutch population is available from the National Health Care Institute (ZiN). The ZiN claims database covers all outpatient prescriptions reimbursed under the Dutch mandatory Health Insurance scheme. Drugs were classified in 204 pharmaceutical groups according to the four position ATC-code. To these groups, age and gender were added as predictors. The dataset contained 47 million individual prescription records in 2010, covering a population of 16.7 million, of which 70% had at least one prescription.

A training set with disease information was obtained from the primary care database of the Netherlands Institute for Health Services Research (NIVEL). As every citizen is required to have a GP—with the exception of those living institutionalized—this means the dataset is likely to cover the whole Dutch population, with the exception of the 80+ population of which in 2010 a significant part lived institutionalized. All patient contacts were labelled with a diagnostic code, ICPC. A person was defined to have a disease when he/she had at least one contact with a GP for this disease over a period of 3 years. All GP-patients with full data available over 2008–2010 were selected. This resulted in a training set of 276,723 individuals. The selection of 29 diseases was based on a list provided by O’Halloran et al. See Supplementary file S1 for details. We combined the available data (drug utilization, age and gender, and ICPC codes) at Statistics Netherlands within the System of Social Statistical Datasets (SSD). The SSD allows data from different administrative registers to be combined using an anonymous patient identifier for research purposes.

Implementation of RF

Usually, all observations in a training set and all predictors are combined in one RF-analysis. However, within the SSD system, computing power is limited, and analysis with our dataset (276,723 records with 206 variables) proved to be difficult. We therefore used a two-step approach. First, for each chronic disease, persons with the disease were randomly selected, up to a maximum of 5000 patients. To this set, an equal number of persons without the disease was randomly selected and added. For each of these smaller sets, the RF algorithm was applied. The variable importance measure, defined as the average decrease in accuracy when a predictor is left out of the analysis, was evaluated. For each disease, the 10 drug groups with the highest variable importance were selected. By selecting 10 drug groups, the most important predictors were included for all diseases, while limiting the computing times. Second, a new dataset was created for each disease based on the full training set, but only age, gender and the drug groups selected in the first step were added as predictors. We applied RF a second time. For each disease this second RF-model was then applied to obtain the probability of having this disease for each individual in the prescriptions database, hence for the 11.6 million Dutch inhabitants that were reimbursed a prescription drug in 2010. The model was also applied to the remaining 5 million Dutch individuals without any prescription. They received for each of the 29 diseases a probability equivalent to the age and gender specific probability in the training set for those diagnosed with the disease, but not receiving any prescription.
Percentage study population with multiple diseases: 11.5%
Percentage with at least one chronic disease: 33.2%
Persons with 5 or more chronic diseases: 1436
Persons with 4 chronic diseases: 2697
Persons with 3 chronic diseases: 7609
Persons with 2 chronic diseases: 20090
Persons with 1 chronic disease: 60065
Persons without disease: 184826

To measure the performance of the final RF-models, the area under the Receiver-Operator Curve (AUC) was measured for the training set for each disease separately. An AUC-value above .7 is generally considered useful.38 To prevent overfitting, 10-fold cross validation was applied.

The AUC and a 95% confidence interval around the AUC-value were obtained using the R-package ‘cvAUC’39. If the lower boundary of this interval was above .5, we considered the model to perform better than a random prediction.

The predicted population prevalence by age and gender for the Netherlands was graphically compared with a prevalence estimate based on direct extrapolation of the training set prevalence. Correlations were computed as well for the six diseases with lower confidence bound (95%) of the AUC >.7. The age range considered was 30–80 years, since the prevalence below 30 is very low for most chronic diseases and the 80+ population was not well covered in our training set.

For a binary classification of each individual, a cut-off needs to be chosen. This was done by setting an age and disease-specific cut-off value. All persons with a probability higher than the cut-off were classified as ‘ill’. The cut-off was chosen to minimize the deviation between the observed and the predicted prevalence in the training set for each age, gender and disorder.

**Table 1** Pharmaceutical utilization in dataset

| Disease                                      | Persons with at least one recorded episode for each chronic disease in 2008–2010 | Total number of pharmaceutical groups utilized in 2010 | Average number of pharmaceutical groups utilized per person |
|----------------------------------------------|---------------------------------------------------------------------------------|------------------------------------------------------|----------------------------------------------------------|
| Persons without disease                      | 184826                                                                          | 328385                                               | 1.8                                                      |
| Persons with 1 chronic disease               | 60065                                                                           | 235032                                               | 3.9                                                      |
| Persons with 2 chronic diseases              | 20090                                                                           | 125335                                               | 6.2                                                      |
| Persons with 3 chronic diseases              | 7609                                                                            | 63064                                                | 8.2                                                      |
| Persons with 4 chronic diseases              | 2697                                                                            | 27190                                                | 9.9                                                      |
| Persons with 5 or more chronic diseases      | 1436                                                                            | 17219                                                | 11.6                                                     |
| Total                                        | 276723                                                                          | 796225                                               | 2.9                                                      |

Legend: Training set population has been divided into six strata, based on the number of chronic diseases present. First column presents stratum. Second column gives population size. Third column gives total number of pharmaceutical groups utilized. Pharmaceutical groups have been defined in terms of an ATC 4 position code: A01A, A02A, etc. Last column gives average utilization in stratum.

**Table 2** Model outcome AUC with confidence interval, ordered by mean AUC

| Disease                                      | AUC (95% conf. interval) | Prevalence in training set per 10 000 persons |
|----------------------------------------------|--------------------------|----------------------------------------------|
| Parkinson’s disease                          | .89 (.77–1.00)           | 15                                           |
| Diabetes mellitus                            | .87 (.85–.90)            | 421                                          |
| Osteoporosis                                 | .87 (.81–.92)            | 103                                          |
| Heart failure                                | .81 (.74–.88)            | 82                                           |
| Chronic obstructive pulmonary disease        | .79 (.75–.83)            | 209                                          |
| Chronic enteritis/colitis ulcerosa           | .79 (.68–.90)            | 31                                           |
| HIV/AIDS                                     | .78 (.39–1.00)           | 4                                            |
| Asthma                                       | .77 (.74–.80)            | 424                                          |
| Epilepsy                                     | .77 (.66–.87)            | 41                                           |
| Coronary heart disease                       | .70 (.66–.74)            | 255                                          |
| Visual disorder                              | .69 (.64–.73)            | 191                                          |
| Schizophrenia                                | .69 (.48–.89)            | 10                                           |
| Rheumatoid arthritis                         | .68 (.60–.76)            | 66                                           |
| Dementia                                     | .67 (.54–.80)            | 28                                           |
| Congenital neurological anomaly              | .67 (.01–1.00)           | 3                                            |
| Multiple sclerosis                           | .66 (.42–.90)            | 9                                            |
| Cancer                                       | .60 (.56–.64)            | 264                                          |
| Chronic alcohol abuse                        | .59 (.49–.69)            | 45                                           |
| Depressive disorder                          | .58 (.54–.63)            | 253                                          |
| Stroke (including TIA)                       | .57 (.52–.63)            | 137                                          |
| Congenital cardiovascular anomaly            | .57 (.37–.77)            | 7                                            |
| Chronic back or neck disorder                | .56 (.53–.60)            | 432                                          |
| Osteoarthritis                               | .56 (.52–.60)            | 282                                          |
| Anxiety disorder, neurosis, PTSS             | .56 (.50–.61)            | 154                                          |
| Mental retardiation                          | .55 (.36–.74)            | 13                                           |
| Hearing disorder                             | .52 (.44–.61)            | 62                                           |
| Anorexia                                     | .52 (.33–.71)            | 8                                            |
| Gastric or duodenal ulcer                    | .50 (.39–.62)            | 25                                           |
| Tuberculosis                                 | .50 (.06–.94)            | 2                                            |

Legend: First column gives name of chronic disease. Second column gives model outcome as AUC with 95% confidence interval, in order of decreasing AUC. Third column states prevalence of chronic disease or condition in the training set. (n = 276723).

**Outcome measures**

For each disease, the most important drugs according to the variable importance were compared with theoretical drug classifications included in relevant guidelines. For 13 of the 29 chronic diseases pharmaceutical groups used in the Dutch insurance system were available.32 In addition, for four other diseases the drugs found were compared with Dutch treatment guidelines: tuberculosis,33 MS,34 chronic back or neck disorder35,36 and gastric or duodenal ulcer.37

To measure the performance of the final RF-models, the area under the Receiver-Operator Curve (AUC) was measured for the training set for each disease separately. An AUC-value above .7 is generally considered useful.38 To prevent overfitting, 10-fold cross validation was applied.

The AUC and a 95% confidence interval around the AUC-value were obtained using the R-package ‘cvAUC’.39 If the lower boundary of this interval was above .5, we considered the model to perform better than a random prediction.

The predicted population prevalence by age and gender for the Netherlands was graphically compared with a prevalence estimate based on direct extrapolation of the training set prevalence. Correlations were computed as well for the six diseases with lower confidence bound (95%) of the AUC >.7. The age range considered was 30–80 years, since the prevalence below 30 is very low for most chronic diseases and the 80+ population was not well covered in our training set.

For a binary classification of each individual, a cut-off needs to be chosen. This was done by setting an age and disease-specific cut-off value. All persons with a probability higher than the cut-off were classified as ‘ill’. The cut-off was chosen to minimize the deviation between the observed and the predicted prevalence in the training set for each age, gender and disorder.

**Results**

Table 1 gives descriptives for the training set. The average annual number of different pharmaceutical drugs taken by patients in the training set was 2.9, which is very comparable with the utilization in the total Dutch population in the same year (2.8). Table 1 also shows that the number of ATC groups utilized by an individual patient rises proportionally with the number of chronic diseases present.

Table 2 lists the AUC values produced by our analysis, sorted by average AUC. For 17 diseases the lower boundary of the 95% AUC confidence interval was >.5. For 10 diseases the average AUC was .7 or higher, but for only six the lower boundary of the AUC 95% confidence interval was >=.7: Parkinson’s disease, diabetes mellitus, osteoporosis, heart failure, asthma and chronic obstructive pulmonary disease (COPD).

There is some association between the frequency of the disease and the prediction of the AUC. For almost all 12 diseases with a prevalence in the training set higher than 100 per 10 000 persons, the prediction is better than a random assignment. The only exception is anxiety disorder (154 cases per 10 000 persons), with a very poor performance and AUC of .56 (95% cf. = .50–.61). For 11 out of 17 diseases with a frequency below 100 per 10 000 persons, performance is poor, i.e. the lower boundary of the AUC 95% confidence interval was below .5. A notable exception was Parkinson’s disease which...
Table 3 Predictors of chronic diseases in Random Forest analysis

| Disease                                      | ATC4 groups with strongest relation with disease in RF-analysis |
|----------------------------------------------|---------------------------------------------------------------|
|                                              | [1]= strongest relation, [10]= weakest relation               |
|                                              | [1]  | [2]  | [3]  | [4]  | [5]  | [6]  | [7]  | [8]  | [9]  | [10] |
| Parkinson's disease*                         | N04B | N04A | Birthyear | L04A | A07E | C10A | C09B | N05A | C03C | N06D |
| Diabetes mellitus*                           | A10A | A10B | C10A | B01A | H04A | C09A | C09B | C08C | C03C | C07A |
| Osteoporosis                                 | M05B | A12A | A11C | L04A | C10A | B01A | C09D | D01A | C09C | D06A |
| Heart failure                                | C03C | C03D | C032 | A12B | C08D | C08C | C01D | C03A | C10A | C07B |
| Chronic obstructive pulmonary disease*       | R03A | R03B | R06A | A01A | R01A | N06B | J01F | R05D | J06A | J01C |
| Chronic enteritis/colitis ulcerosa           | A07E | L04A | L01B | B03B | A06A | J01M | A11C | A07D | N02A | M01A |
| HIV/AIDS*                                    | J05A | J01E | J04A | J01F | N01B | J07B | D06B | A02B | J01C | J01A |
| Asthma*                                      | R03A | R03B | R03C | Birthyear | H02A | A07A | S01G | R01A | R06A | R05D |
| Epilepsy*                                    | N03A | N05B | Birthyear | A03F | N05A | B01A | N06A | D04A | D11A | N05C |
| Coronary heart disease*                      | C01D | C08D | C03C | C01B | C03A | C09A | D06A | C01E | C09C | B03A |
| Visual disorder                              | S01E | S01B | Birthyear | S01C | A10B | S01F | D02A | S01A | A10A | S01X |
| Schizophrenia*                               | N05A | N05B | N05C | N04A | N06A | N06B | Birthyear | N03A | N07A | N07C |
| Rheumatoid arthritis*                        | L04A | P01B | A07E | B03B | N02A | L01B | H02A | D02B | M05B | D06A |
| Dementia*                                    | N06D | Birthyear | N05A | A12A | C03C | N03A | M05B | Y | D03D | C09D |
| Congenital neurological anomaly              | M03B | G04B | N03A | J01X | D07X | N05A | J01E | A12A | D01A | N05B |
| Multiple sclerosis*                          | L03A | M03B | G04B | N03A | N06A | N04B | B03B | S01A | J01X | C03C |
| Cancer*                                      | Birthyear | L02B | H03A | Y | D06A | A04A | Gender | L02A | G03C | A12A |
| Chronic alcohol abuse*                       | N07B | N05A | Birthyear | N05B | G04C | A02B | N06A | M04A | A10B | N05C |
| Depressive disorder*                         | N06A | N05A | N05B | N06B | N05C | N07B | N03A | A03F | A11C | G04B |
| Stroke (including TIA)                       | B01A | V03A | C01D | C01B | C07A | Birthyear | C08C | C01A | S01C | S01E |
| Congenital cardiovascular anomaly            | B01A | C07A | J01C | N03A | C09A | D06A | R03B | Y | D02A | S02C |
| Chronic back or neck disorder b              | N02A | M01A | A02B | A06A | N02B | C05A | S02C | N03A | H02A | R05D |
| Osteoarthritis                               | Birthyear | B01A | Gender | M04A | C10A | N02B | C10B | C03A | S01E | N02A |
| Anxiety disorder, neurosis, PTSS             | N06A | N05B | N05A | C07A | N01B | N05C | A03F | A06A | N03A | D05A |
| Mental retardaliation                         | N05A | N03A | D02A | D06A | A06A | N05B | Y | D10A | S01F | N01B |
| Hearing disorder                             | Birthyear | L02A | B02A | S01X | D05A | G04C | H02A | A10B | C08B | C07B |
| Anorexia                                     | Gender | G03A | A06A | A01A | Y | J01X | G03H | R05D | G01A | A12B |
| Gastric or duodenal ulcer b                  | A02B | D05A | G04B | A07A | A03A | A03F | M01A | A11C | D06B | A06A |
| Tuberculosis b                               | J04A | D02A | D07X | C01B | J01A | C08C | C03C | S01C | S02C | C03E |

Legend: First column gives name of chronic disease. The next 10 columns list the predictors used in the final RF-model, in order of decreasing importance. To facilitate comparison, diseases are presented in the same order as in table 2.

a: For these diseases, comparison is possible with pharmaceutical groups listed in risk adjustment compulsory insurance. The shaded groups are also used for the detection of these diseases by Dutch insurers.32

b: These diseases have been compared with ATC-groups mentioned in the relevant Dutch treatment guidelines. The shaded groups are included in these guidelines.33–37

despite a low frequency (15 per 10 000 persons) seems to be very predictable from drugs utilization.

In table 3 predictors of all model output from the RF-analysis are ranked by importance. The shaded areas denote drugs that are also mentioned as indicator drugs for these diseases in the theoretical drug classifications we compared with. Only for cancer we found no similarities. For all other diseases, the ATC codes mentioned by insurers and guidelines are also strong predictors for the corresponding diseases in our RF-models. However, our models show a number of additional predictors for most disorders. Supplementary file S2 gives more information.

The actual prevalence in the training set and the calculated prevalence based on applying the final RF-models have been compared for the six diseases with a lower bound of the AUC 95% confidence interval >.7. Except Asthma, correlations are above .9. Asthma shows correlations of .43 for males and .66 for females, indicating poor performance. Looking at the graphs for osteoporosis, a large discrepancy exists between predicted and observed prevalence around the age of 70. Figure 1 gives an example (COPD, male). A full set of figures is found in Supplementary file S3.

Discussion

For a broad range of 29 diseases, RF was used to predict disease prevalence based on medication use. Predictive performance was acceptable for 6 out of 29 diseases and would result in reliable estimates of population prevalence. Furthermore, we find that theory-based indicator drugs were included in the range of diseases identified by the RF model. This seems to be independent from the performance of the models, which indicates that the RF algorithm can also be used to identify suitable predictors, even in those cases where the predictive performance is low. Especially for diabetes, heart failure and COPD we observe a high correlation between estimated and observed population.

Our outcomes can be compared with a few other studies. Chaudhry20 predicted the presence of diabetes with an AUC of .95 and dementia with an AUC of .875, higher than the .87 and .67 we found. However, for dementia he used dementia-coded doctor visits as predictors, while we use this as our definition of disease. Khalilia et al.21 used data on hospital stays as predictors, on a very large set (8 million records). A training set was generated by bootstrapping. The average AUC he reports (.88) is much higher than those we found. For the two diseases which could be directly compared (diabetes and osteoporosis) he finds almost the same AUC (.879 and .870 respectively) as we found, .87 for both. Compared with these two previous studies, we included a relatively broad range of diseases and added the comparison with theory-based models.

While the method seems useful for some diseases, the predictive performance is still low for most diseases. This could have multiple causes. First, for some diseases, there is no standard pattern of drugs included in all treatment options. In addition, drugs might be
prescribed for multiple diseases. For instance, the two strongest predictors for asthma and COPD are the same (R03A and R03B, table 3). As a result, misclassification of asthma and COPD patients is likely to occur, which has not been further investigated in this study. Furthermore, patients and GPs may deal with diseases in different ways. Based on patient characteristics a GP will sometimes advise lifestyle changes instead of drugs, but will treat similar cases in other instances immediately with drugs. In addition, the patient may have treatment preferences. The relationship between diagnosis and drugs can also change over time. Innovation or policy changes can strongly influence prescription behaviour, making regular calibration of the algorithms necessary.

Second, the predictive power is likely limited due to weaknesses of the current data. In the current training set, only 3 years of diagnoses are used. While many patients with a chronic disease are visiting a GP more than once every 3 years, some patients who visit less frequently will not occur as diseased in the training set. Furthermore, some diseases might not be treated primarily by a GP, but directly in the hospital, also resulting in missing diagnoses in the training set. As the training set serves as a ‘golden standard’, any diagnosis errors in the training set will translate into the final predictions. Investing in a smaller set of persons for which disease diagnosis is even more reliable, e.g. through the use of cohort studies may provide a training set with better performance. The disadvantage of such a cohort, and the advantage of our current approach is that for rare diseases, relationships between disease and drugs would have to be derived from only a very limited number of disease cases.

Next to errors in disease diagnosis, drugs use measures could also be improved. Drug use often varies between years. Grouping multiple years of drug use could improve results. Also, more complete drug utilization data could be obtained by including inpatient drugs. For some diseases, utilizing more detailed pharmaceutical predictors, such as ATC4 or ATC5 groups, would improve results.

Even though improvements might be needed to obtain reliable prevalence estimates for most diseases, for 16 out of 17 diseases for which theory-based predictors were found within existing guidelines, important similarities were found. This means that even though the predictive power of the algorithm on the current data is insufficient, it is still possible to identify relevant drug groups. Compared with purely theory-based models, the RF algorithms have the important advantage of coming with confidence intervals and information about model performance. From this similarity we also infer that Dutch general practitioners broadly follow existing pharmaceutical guidelines. Cancer was the only disease for which the drugs found using the RF algorithm differ from theory. This could be the result of grouping all cancers together, and many drugs used in cancer treatment were not covered by the dataset as they were prescribed in a hospital setting.

We do not want to suggest that prediction models can entirely replace current GP registers or population surveys. On the contrary, since without these registries the models cannot be built or validated. However, even in countries like the Netherlands which are covered by both population surveys and GP networks the method is of practical value, as it allows for analysis on subgroups, such as regions or stratifications by socio-economic status. The primary care database used as training set has been enlarged in recent years, but still covers at this moment only 10% of the population and the Dutch GPs. Using drug use will allow for better prevalence estimates for the 90% not covered.

Because the full population is covered in the prescription data we use, and the model provides estimates of the probability of having a disease at the individual level, other useful applications would be pre-selecting subjects for medical trials, or making case-mix corrections, e.g. for comparing hospital performance.

To conclude, combining diagnosis data and drug use by the RF algorithm provides can be a useful tool to predict population prevalence. Applications include situations where the diagnosis data is not necessarily representative for the population of interest.
but the relation found between diagnosis and drug use is representative. Furthermore, it can be used to select relevant drug use groups in almost all cases.

Supplementary data

Supplementary data are available at EURPUB online.

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Introduction

Cannabis is illegal in France but, as in many countries, legalization is under debate. Survey data could be used to evaluate the prevalence of cannabis use in the general population whereas emergency department (ED) visits for cannabis exposure (CE) are symptomatic of pathologic situations associated with cannabis. Zhu used ED visits in the United States to report an increase in CE visits in adults.1 In the USA, various studies2–6 using Poison Control Center (PCC) data2–6 or hospital data2–5 have reported that legalization of cannabis was associated with an increase in un-intentional cannabis ingestions by young children. In France, without any change in cannabis law, an increase in phone calls to PCC was first reported in 2009.9 In 2017, Claudet also reported an increase in hospital admissions for CE in a retrospective observational study (2004–14) using hospital data and retrospective reading of patients’ medical file.10 However, this study could have over-estimated the increase, since during the first years of the study, only inpatients’ medical file was computerized. Children who had not been admitted to hospital following the ED visit were thus not caught in the study in many EDs. Moreover, the methodology used did not allow for an easy analysis repetition to survey the trend. In France, each ED admission has to be reported daily through Electronic Emergency Department Abstracts (EEDA). These EEDA are transmitted to OSCOUR network included in the French Syndromic Surveillance System (SSS) System SurSaUD3 coordinated by Public Health France.11 EEDA have previously been used for various epidemiological studies in EDs.12–15 Our study aimed to measure, in southern France, the trend of CE visits in EDs, in children and young adults, using the daily available data included in the French SSS.

Methods

Data source

Our cross-sectional study analysed EEDA related to patients younger than 27 years old. EEDA have been included in the French SSS since 2004. They are directly collected from patients’ computerized medical files filled in during medical consultations. Details of this network have been published elsewhere.16,17 The Provence-Alpes-Côte d’Azur...