

400 375 201710 201711 201712 201801 201802 201803 201804 201805 201806 201807 201808 201809

201610 201611 201612 201701 201702 201703 201704 201705 201706 201707 201708 201709 Source: Healthware analysis based on NHIFA data

Changes to subsidized medicinal product categories, September 2018



in

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Applications for

reimbursement



Number of reimbursed products



Source: Healthware analysis based on NHIFA data



Market data

Average number of medical sales reps

Source: NHIFA data, Healthware analysis

TOP 10 active substances by number of patients (thousand patients)

234 228 220 219

194

184 183

330

294

perindopril and diuretics

o antoprazole

nebivolol

opurinol

250



TOP 10 brands by all reimbursement paid

			•			
CLEXANE						665 M Ft
JAKAVI				465 M Ft		
XARELTO			434 N	1 Ft		
TECFIDERA		364 M Ft				
XEPLION		346 M Ft				(0
GILENYA	305 M Ft					share
SUTENT	296 M Ft					of T
ELIQUIS	296 M Ft					OP 10
IBRANCE	291 M Ft					0 bra
XULTOPHY	288 M Ft				13%	nds
				Source: Pharmacy turno	ver data, Healthware ana	lysis



Classifying patient paths - Case study

In this case-study we are introducing a solution for the frequently occurring problem, how a patient path trajectory system (eg. laboratory values measured in fixed time frames for every patient) - which might seem to be chaotic for the first sight – can be classified into homogenous groups, clusters effectively.

The method is presented on simulated data. Each group is expected to contain patient paths having similar properties. In other words, they are expected to be close to each other (according to a specific definition of a distance between patient paths). The characteristics of the obtained groups, especially their temporal trends reveal the essential differences between the groups. As an advantage of clustering, it can be expected that after the execution of the method it is easier to compare the groups, and the obtained types of patient paths can also be used in later models.

The main properties of the clusters:

patient paths belonging to the same group are similar
patient paths of different groups differ markedly from each other

We applied the kmi package of K for longitudinal clustering". The algorithm implemented in the package is also capable to determine the optimal number of groups. Selecting too many groups would produce a complicated model. On the other hand, if too few groups were creat ed, then significantly differing patient paths would be classified into the same cluster.

For illustration, we demonstrate the method using simulated time series. We simulated two groups of patient paths including 200 patients in each group. The trend of the first group is increasing over time according to the square root function. The trend of the second group is decreasing, it is inversely proportional to the time elapsed. By adding individual, patient-specific noise processes - which are correlated in time - to the trends, each patient path became different from the common temporal trend. (The noise processes were chosen to be first-order autoregressive processes.)



Recoling Times In summary, we concluded that the algorithm reliably classified the patient paths into coherent clusters even in the presence of heavily correlated and highly scattered poice processes

This clustering method was successfully applied to a real problem, too. This year on the ISPOR conference we published it in our poster². We classified the trajectories of the disease activity index (DAS28) of patients diagnosed with rheumatoid arthritis. The obtained groups differed significantly from each other in the duration of their first biological therapy, the proportion of patients who reached remission, the proportion of knee and hip operations, as well as in the total healthcare costs.

Genolini Christophe; kml: K-Means for Longitudinal Data. R package version 2.4.1; <u>link</u> Tamás Balázs, Péter Juhász, Andrea Radnai, Károly Dede, Miklós Bacskai, Gyula Po Jisease Activity Score in Hungary; <u>link</u>