Newsletter

Actualities of Hungarian pharmaceutical financing market

HEALTHWARE

News, current issues

- Legislations come into force between 01/02/2017 and 01/03/2017: Act LXXXIII of 1997 (01.02.2017,03.02.2017)
- NEWS [HUN]: "Hungarian clinics in the European Reference Network" link
- NEWS [EN]: "World's most admired pharmaceutical companies 2017" link
- NEWS [EN]: "European and US regulators agree on mutual recognition of inspections of medicines manufacturers" link
- NEWS [HUN]: "New things to get familiar with in healthcare" link
- NEWS [HUN]: "How many billions do we leave in pharmacies?" link
- NEWS [HUN]: "How much would really cover our healthcare?" link
- STUDY [HUN]: "Healthcare expenditures in Hungary, 2010-2015" link

Macro approach to financing healthcare and medicinal products

Balance of the Health Insurance Fund

Billion HUF 2017 2017 Health Security Fund 2016. I-XII. appropriation % of % of I. months (1 Jan) appropriation last year 112,0% **Total of Budgetary Expenditures** 2 133,1 2 139,5 178,0 99,8% 1 089.9 1 121,4 101,0% 120.8% Curative preventive provisions 94,3 Contracted specialty care 683,3 801,3 58,9 88,2% 114,5% Medicine subsidies (pharmacy) 327.9 313.0 25.3 97.2% 96.4% **Total of Budgetary Revenues** 2 043.9 2 059.1 189.5 110.4% 105.5% **Social Security Contributions** 1 479,5 1 532,4 145,5 113,9% 108,5% Contribution of Pharmaceutical 71,6 66,0 4,6 83.7% 96,9% Manufacturers and Wholesalers Balance 11.5 55.2%

Tell us your opinion!

We are renewing our Newsletter.

We kindly ask you to share your opinion to help us improve the Newsletter.

You can fill the questionnaire and write feedback with following the link below.

Thankfully, The Healthware Team

Our questionnaire is here: link

Questionnaire

In expenditures and revenues of 2017 budget, there is 4.86% increase compared to appropriation of 2016 but only 0.3% increase compared to fulfilment, despite that the appropriation of expenditures were raised with 80 billion HUF. Revenues of Social security contributions are 52.9 billion HUF (3.6%) higher, while Contribution of manufacturers and wholesalers are 26 billion HUF (6.2%) lower in the appropriation of 1st of January, than in the last year's fulfilment. The pharmaceutical budget was planned to be 23.6 billion HUF (8.2%) higher than the last year appropriation (without the special budget drugs), and 9.2 billion HUF (2.9%) lower than the last year fulfilment.

In the first month of 2017, we can see 6.43% surplus in Health Security Fund, compared to the prorated appropriation of expenditures. Fulfilment of medicine subsidies is 2.8% lower than periodic appropriation. We can see only technical reasons, because on the contrary of the 25.3 billion HUF financial fulfilment, more than 28 billion HUF monthly reimbursement turnover was issued, based on the public real-world data of the December-January period.

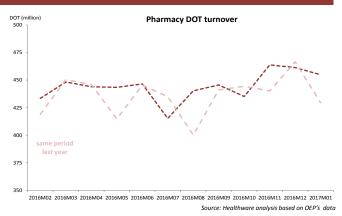
Changes to subsidised medicinal product categories

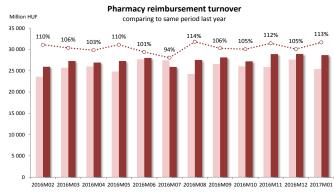
Changes in the public drug list	2016 Sep.	2016 Oct.	2016 Nov.	2016 Dec.	2017 Jan.	2017 Feb.	2017
Number of new products	31	32	12	25	13	12	50
Number of new Al	3	15	0	6	1	2	9
Number of delisted products	10	28	33	21	228	10	259
Prices							
Decrease	98	11	5	11	4	4	19
Increase	1	1	0	3	3	0	6

Changes in the public drug list	2016 Sep.	2016 Oct.	2016 Nov.	2016 Dec.	2017 Jan.	2017 Feb.	2017
Reimbursement							
Decrease	237	5	4	27	2	4	33
Increase	28	5	0	6	3	0	9
Co-payment							
Decrease	150	19	5	20	8	5	33
Increase	152	1	0	17	3	0	20

Source: Healthware analysis based on OEP-PUPHA data

Dynamics of the sales/circulation of prescription-only-medicine





Prescription drugs' DOT turnover in 2016 was 1.18% higher than in 2015, so the trend of drug consumption is still increasing, but in slower rate than in 2014 (2.74%) or 2013 (2.23%). Meanwhile, the reimbursement turnover was higher with 5.56%, because of the additional 14.2 billion HUF fulfillment of special permission appropriation, the 6% growth of reimbursement turnover of out of-fix group products, and stagnation of fixed market. The average reimbursement per DOT was higher with 4.33% than the 2015's average. New ATCs that got authorized in 2014-2016 generated 7.6% of annual reimbursement turnover, while only 1.1% of annual DOT turnover. Drug sales in the first month of 2017 was 5.94% higher than the same period last year, while the average reimbursement per DOT increased with 6.49%. The reimbursement turnover was higher with 12.82% for this period compared to last year.

pharmaceutical market



Market data

Marketing authorisation information

2016	EMA	OGYI	2016 - Q4	EMA	OGYI	January 2017	EMA	OGYI
New brands	71	173	New brands	10	45	New brands	2	9
New SKUs	625	1 765	New SKUs	123	472	New SKUs	6	55
Source: Healthware analysis based on OGYI's and EMA's data								

TOP10 **DISTRIBUTOR** by all reimbursement paid in January 2017



TOP10 BRAND by all reimbursement paid in January 2017



TOP10 ATC by all reimbursement paid in January 2017



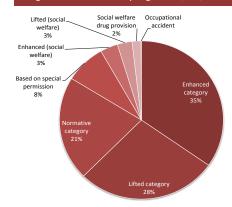
Source: Healthware analysis based on the sales turnover that pharmacies produced from POM

Average number of medical sales reps; 01/2017 1 283 Medical aids + nutritions Medicinal products Nutritions

Medicinal products + nutritions 10 All Medical aids 262 Source: Healthware analysis based on OGYI's

Drug reimbursement by legal title; 01/2017

Medicinal products + aids



TOP10 ATC by number of patients in January 2017

TOP 10 - ATC	International non-proprietary name (INN)	Patients
B01AC06	acetilszalicilsav	351 509
C09BA04	perindopril and diuretics	298 320
C07AB12	nebivolol	256 661
C08CA01	amlodipin	254 852
A02BC02	pantoprazol	237 718
J01CR02	amoxicillin - laktamázgátló kombinációk	236 803
C10AA07	rosuvastatin	225 004
A11CC05	kolekalciferol	224 263
M04AA01	allopurinol	214 466
C10AA05	atorvastatin	212 821

Source: Healthware analysis based on the sales turnover that pharmacies produced from POM

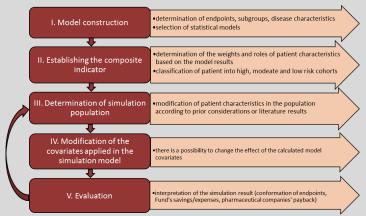
A possible implementation of performance-based funding — Case study

From the Health Fund's perspective, reimbursements should not be allocated only by the outcomes of clinical trials, but significant real life data about the examined patient population is also needed, especially in case of high-value therapies. It can provide the basis for the paybacks of pharmaceutical companies in the case of confirmed inefficiency. The validation and efficient use of widely accepted financing logic require investments and complex, observational study protocol level approach, since the therapeutic effectiveness should be hold by all actors in the health care system. The outcome based financing system provides a framework for the payer and the pharmaceutical company to agree on a price and payback system connected to the clinical or intermediate endpoints, measured in the future, in relation to the patients' quality of life [1]. For these agreements, it is expedient that the parties can measure and simulate the behaviour and expected results of the particular system in advance. Hereinafter, a possible implementation of this process will be presented.

First of all, the relevant disease-specific endpoints, subgroups and characteristics have to be chosen according to expert opinions or information available in the professional literature. Since the volume of disease characteristics can be huge, the aim is to create a complex indicator (composite indicator), in which these variables can be concentrated and interpreted more easily (through dimension reduction). Being aware of the composite indicator's distribution, patients can be distinguished by their different risk profiles (e.g. high, moderate and low risk cohorts). The modeling of the divergent endpoints can necessitate various statistical methods. Consequently, in the determination of the endpoints, the modeling methodology of effectiveness should also be settled. The disease characteristics have different weights, which are defined by the proportion of their significant effects on the examined endpoints, and whether these effects were mostly positive or negative.

During the simulation, different scenarios can be taken into consideration, the initial population can be modified or the estimated covariates from the modeling can be altered. With the modification of the population we can investigate the possible alterations of the different endpoints if patients with certain characteristics were present in smaller or larger proportion (e.g. we reduce the presence of the high-risk patients) within the patient populaThe revision of the covariates can be justified by the outcomes of other trials, expert opinions or even the presumed influence of the different interventions (e.g. higher expenditure on the medical provision of the high risk patients could improve the measured covariates' value). Subsequently, as a result of the simulation performed with the changed parameters, beside the expected improvement or deterioration of endpoints, the payer can model the expected value of future savings or expenses, while the pharmaceutical companies can model the value of payback.

In order for the outcome based financing system to become efficient for all actors, strategic planning and successful preparations are sufficient, for which the above mentioned simulation framework can provide a reliable support.



[1] Carlson J, Sullivana S, Garrisona L, at al. Linking payment to health outcomes: A taxonomy and examination of perform eimbursement schemes between healthcare payers and and manufacturers. Health Policy 2010; doi: 10.1016/j. healthpol. 2010.02.05