Nr.p.k.	Reģ. numurs	Zāļu nosaukums, zāļu forma, stiprums/ koncentrācija	Aktīvās vielas nosaukums	Reģistrācijas apliecības īpašnieks, valsts	Izmaiņu procedūras numurs	Procedūras numurs
1	2	3	4	5	6	7
1	12-0114	Gemcitabine Accord 100 mg/ml concentrate for solution for infusion, Concentrate for solution for infusion, 100 mg/ml	Gemcitabinum	Accord Healthcare Limited, Lielbritānija		NL/H/2136/001/IA/016/G
2	09-0487	Gemcitabine Hydrochloride Accord 1 g powder for solution for infusion, Powder for solution for infusion, 1 g	Gemcitabinum	Accord Healthcare Limited, Lielbritānija		UK/H/1124/002/IA/030/G
3	11-0321	Gemcitabine hydrochloride Accord 2 g powder for solution for infusion, Powder for solution for infusion, 2 g	Gemcitabinum	Accord Healthcare Limited, Lielbritānija		UK/H/1124/003/IA/030/G
4	09-0486	Gemcitabine Hydrochloride Accord 200 mg powder for solution for infusion, Powder for solution for infusion, 200 mg	Gemcitabinum	Accord Healthcare Limited, Lielbritānija		UK/H/1124/001/IA/030/G
5	15-0294	Botox 100 Allergan Units powder for solution for injection, Powder for solution for injection, 100 Allergan Units	Toxinum botulinicum A	Allergan Pharmaceuticals Ireland, Īrija		IE/H/0113/001/IA/101
6	15-0295	Botox 200 Allergan Units powder for solution for injection, Powder for solution for injection, 200 Allergan Units	Toxinum botulinicum A	Allergan Pharmaceuticals Ireland, Īrija		IE/H/0113/003/IA/101
7	15-0293	Botox 50 Allergan Units powder for solution for injection, Powder for solution for injection, 50 Allergan Units	Toxinum botulinicum A	Allergan Pharmaceuticals Ireland, Īrija		IE/H/0113/002/IA/101
8	12-0129	VONILLE 60 micrograms/15 micrograms film-coated tablets, Film-coated tablets, 60 micrograms/15 micrograms	Gestodenum, Ethinylestradiolum	Exeltis Baltics UAB, Lietuva		NL/H/1902/001/IB/006
9	08-0320	Vidotin 4 mg tablets, Tablets, 4 mg	Tert-butylamini perindoprilum	Gedeon Richter Plc., Ungārija		DK/H/1128/001/IA/011/G
10	08-0321	Vidotin 8 mg tablets, Tablets, 8 mg	Tert-butylamini perindoprilum	Gedeon Richter Plc., Ungārija		DK/H/1128/002/IA/011/G
11	08-0130	Concerta 18 mg prolonged-release tablets, Prolonged-release tablets, 18 mg	Methylphenidati hydrochloridum	Johnson & Johnson UAB, Lietuva		UK/H/0544/001/IA/080
12	10-0529	Targin 10 mg/5 mg prolonged release tablets, Prolonged-release tablets, 10 mg/5 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/001/IA/036/G
13	14-0017	Targin 15 mg/7.5 mg prolonged-release tablets, Prolonged-release tablets, 15 mg/7.5 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/006/IA/036/G

1	2	3	4	5	6	7
14	14-0016	Targin 2.5 mg/1.25 mg prolonged-release tablets, Prolonged-release tablets, 2.5 mg/1.25 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/005/IA/036/G
15	10-0530	Targin 20 mg/10 mg prolonged release tablets, Prolonged-release tablets, 20 mg/10 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/002/IA/036/G
16	14-0018	Targin 30 mg/15 mg prolonged-release tablets, Prolonged-release tablets, 30 mg/15 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/007/IA/036/G
17	10-0531	Targin 40 mg/20 mg prolonged release tablets, Prolonged-release tablets, 40 mg/20 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/003/IA/036/G
18	10-0528	Targin 5 mg/2.5 mg prolonged release tablets, Prolonged-release tablets, 5 mg/2.5 mg	Oxycodoni hydrochloridum, Naloxoni hydrochloridum	Mundipharma GmbH, Austrija	DE/H/xxxx/IA/805/G	DE/H/1612/004/IA/036/G
19	10-0320	Diovan 3 mg/ml oral solution, Oral solution, 3 mg/ml	Valsartanum	Novartis Finland Oy, Somija		SE/H/0406/007/P/001
20	10-0502	Latalux 50 micrograms/ml eye drops, solution, Eye drops, solution, 50 µg/ml	Latanoprostum	PharmaSwiss Česka Republika s.r.o., Čehija		NL/H/1654/001/II/009/G
21	05-0020	Stadapress 0,2 mg film-coated tablets, Film-coated tablets, 0,2 mg	Moxonidinum	Stada Arzneimittel AG, Vācija		NL/H/0406/001/IA/046
22	05-0021	Stadapress 0,3 mg film-coated tablets, Film-coated tablets, 0,3 mg	Moxonidinum	Stada Arzneimittel AG, Vācija		NL/H/0406/002/IA/046
23	05-0022	Stadapress 0,4 mg film-coated tablets, Film-coated tablets, 0,4 mg	Moxonidinum	Stada Arzneimittel AG, Vācija		NL/H/0406/003/IA/046
24	15-0168	Duloxetine Teva 30 mg gastro-resistant capsules, hard, Gastro-resistant capsules, hard, 30 mg	Duloxetinum	Teva B.V., Nīderlande		DE/H/5010/001/IB/003/G
25	15-0169	Duloxetine Teva 60 mg gastro-resistant capsules, hard, Gastro-resistant capsules, hard, 60 mg	Duloxetinum	Teva B.V., Nīderlande		DE/H/5010/002/IB/003/G
26	12-0217	Avixar 100 mg chewable tablets, Chewable tablets, 100 mg	Sildenafilum	Teva Pharma B.V., Nīderlande		NL/H/2396/003/IA/012
27	12-0216	Avixar 50 mg chewable tablets, Chewable tablets, 50 mg	Sildenafilum	Teva Pharma B.V., Nīderlande		NL/H/2396/002/IA/012
28	08-0131	Concerta 36 mg prolonged-release tablets, Prolonged-release tablets, 36 mg	Methylphenidati hydrochloridum	UAB Johnson & Johnson, Lietuva		UK/H/0544/002/IA/080
29	08-0132	Concerta 54 mg prolonged-release tablets, Prolonged-release tablets, 54 mg	Methylphenidati hydrochloridum	UAB Johnson & Johnson, Lietuva		UK/H/0544/003/IA/080
30	98-0099	Rispolept 1 mg film-coated tablets, Film-coated tablets, 1 mg	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/003/IA/057

1	2	3	4	5	6	7
31	01-0372	Rispolept 1 mg/ml oral solution, Oral solution, 1 mg/ml	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/008/IA/057
32	98-0100	Rispolept 2 mg film-coated tablets, Film-coated tablets, 2 mg	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/004/IA/057
33	98-0101	Rispolept 3 mg film-coated tablets, Film-coated tablets, 3 mg	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/005/IA/057
34	98-0102	Rispolept 4 mg film-coated tablets, Film-coated tablets, 4 mg	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/006/IA/057
35	03-0096	Rispolept Consta 25 mg powder and solvent for suspension for prolonged-release injection, Powder and solvent for suspension for prolonged-release injection, 25 mg/2 ml	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/013/IA/057
36	03-0097	Rispolept Consta 37.5 mg powder and solvent for suspension for prolonged-release injection, Powder and solvent for suspension for prolonged-release injection, 37.5 mg/2 ml	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/014/IA/057
37	03-0098	Rispolept Consta 50 mg powder and solvent for suspension for prolonged-release injection, Powder and solvent for suspension for prolonged-release injection, 50 mg/2 ml	Risperidonum	UAB Johnson & Johnson, Lietuva		DE/H/2184/015/IA/057
38	11-0313	Osagrand 3 mg/3 ml solution for injection, Solution for injection, 3 mg/3 ml	Acidum ibandronicum	Zentiva, k.s., Čehija		CZ/H/0255/001/IB/009/G

Zāļu reģistrācijas
departamenta vadītāja
M.Emersone