19 May 2022
EMA/579444/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: estradiol (except cream/balm/emulsion for application in female genital area)

Procedure no.: PSUSA/00010440/202108

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(10 mg/22 cm2), dispositif transdermique				
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermique				
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
Estrofem® 2 mg film-coated tablets	not available	PA 218/50/1	NOVO NORDISK A/S	IE
Климара 50 микрограма/24 часа трансдермален пластир	not available	9900139	BAYER AG	BG
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 653 7 4	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 916 2 8	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 743 6 9	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 915 6 7	BAYER HEALTHCARE	FR
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR

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OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(5 mg/11 cm²), dispositif transdermique				
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
Estrofem® 2 mg film-coated tablets	not available	PA 218/50/1	NOVO NORDISK A/S	IE
Progynova ® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova® 2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
Progynova® 2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal spray, solution	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR

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Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI

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Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/01	GEDEON RICHTER PLC.	HU
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/02	GEDEON RICHTER PLC.	HU
Rewellfem 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	2020090260	GEDEON RICHTER PLC.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rewellfem 10 míkrógramma skeiðartöflur.	HU/H/0633/001	IS/1/20/086/01	GEDEON RICHTER PLC.	IS
Vagirux 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	RVG 124964	GEDEON RICHTER PLC.	NL
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	140403	GEDEON RICHTER PLC.	AT
Вагирукс 10 микрограма вагинални таблетки	HU/H/0633/001	20200165	GEDEON RICHTER PLC.	BG
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
OROMONE 1 mg, comprimé pelliculé	not available	34009 351 167 5 8	MYLAN MEDICAL SAS	FR
OROMONE 2 mg, comprimé pelliculé	not available	34009 342 436 7 7	MYLAN MEDICAL SAS	FR
Fematab 1 mg film-coated tablet	not available	PA 2010/11/1	MYLAN IRE HEALTHCARE LIMITED	IE
Fematab 2 mg Film-coated tablet	not available	PA 2010/11/2	MYLAN IRE HEALTHCARE LIMITED	IE
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal spray, solution	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku,	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermalni sprej, otopina				
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek,	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermalno pršilo, raztopina				
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
Estrofem 2 mg - Filmtabletten	not available	16 889	NOVO NORDISK PHARMA GMBH	AT
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Estrofem® 2 mg film-coated tablets	not available	PA 218/50/1	NOVO NORDISK A/S	IE
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Estrofem 1 mg Filmtabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem 1 mg - Filmtabletten	DK/H/0117/001	1-22071	NOVO NORDISK PHARMA GMBH	AT
Estrofem 2 mg - Filmtabletten	not available	16 889	NOVO NORDISK PHARMA GMBH	AT
Estrofem 1 mg, filmomhulde tabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrifam 2 mg Filmtabletten	DK/H/0117/002	40226.01.00	NOVO NORDISK PHARMA GMBH	DE
Estrifam 1 mg Filmtabletten	DK/H/0117/001	40226.00.00	NOVO NORDISK PHARMA GMBH	DE
Estrofem 1 mg kalvopäällysteiset	DK/H/0117/001	12623	NOVO NORDISK A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletit				
Estrofem 1 mg filmovertrukne tabletter	DK/H/0117/001	17461	NOVO NORDISK A/S	DK
Estrofem 2 mg filmovertrukne tabletter	DK/H/0117/002	06697	NOVO NORDISK A/S	DK
Estrofem 1 mg filmuhúðaðar töflur	DK/H/0117/001	IS/1/03/031/01	NOVO NORDISK A/S	IS
Estrofem 2 mg filmuhúðaðar töflur	DK/H/0117/002	IS/1/03/031/02	NOVO NORDISK A/S	IS
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
Estrofem 1 mg Filmtabletten	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
Climara-50, 50 µg/dobę (3,8 mg), system transdermalny	not available	4583	BAYER AG	PL
Progynova mite-Dragees	not available	14.422	BAYER AUSTRIA GMBH	AT
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Progynova 2 mg compresse rivestite	not available	021226016	BAYER SPA	IT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455384	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455483	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455582	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455681	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455889	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455780	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455988	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2456085	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 100 100microgramas/24h Sistema transdérmico	not available	2456184	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 100 100microgramas/24h Sistema transdérmico	not available	2456382	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 100 100microgramas/24h Sistema transdérmico	not available	2456383	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Dermestril 100 100microgramas/24h Sistema transdérmico	not available	2456481	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Estring 7,5 mikrog / 24 timmar vaginalinlägg	not available	11352	PFIZER OY	FI
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
Progynova mite-Dragees	not available	14.422	BAYER AUSTRIA GMBH	AT
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Климара 50 микрограма/24 часа трансдермален пластир	not available	9900139	BAYER AG	BG
Progynova mite-Dragees	not available	14.422	BAYER AUSTRIA GMBH	AT
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
ESTRADERM MX 100 mcg/die cerotto transdermico	not available	031773031	MERUS LABS LUXCO II S.À R.L.	IT
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Estrofem 2 mg filmdragerade tabletter	not available	11466	NOVO NORDISK A/S	FI
Estrofem 2 mg kalvopäällysteiset tabletit	not available	11466	NOVO NORDISK A/S	FI
Estrofem 2 mg filmsko obložene tablete	not available	H/97/00590/001	NOVO NORDISK A/S	SI
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
Estradot 100 mikrograma/24 sata, transdermalni flaster	SE/H/1421/005	HR-H-660461417-02	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 100 mikrograma/24 sata, transdermalni flaster	SE/H/1421/005	HR-H-660461417-03	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 100 mikrograma/24 sata,	SE/H/1421/005	HR-H-660461417-04	NOVARTIS HRVATSKA D.O.O.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermalni flaster				
Estradot 25 mikrograma/24 sata, transdermalni flaster	SE/H/1421/001	HR-H-259673389-02	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 25 mikrograma/24 sata, transdermalni flaster	SE/H/1421/001	HR-H-259673389-03	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 25 mikrograma/24 sata, transdermalni flaster	SE/H/1421/001	HR-H-259673389-04	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-02	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-03	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-04	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 100 mikrogram/24 timmar, depotplåster	SE/H/1421/005	17559	NOVARTIS SVERIGE AB	SE
Estradot 25 mikrogram/24 timmar, depotplåster	SE/H/1421/001	20333	NOVARTIS SVERIGE AB	SE
Estradot 37,5 mikrogram/24 timmar, depotplåster	SE/H/1421/002	17556	NOVARTIS SVERIGE AB	SE
Estradot 50 mikrogram/24 timmar, depotplåster	SE/H/1421/003	17557	NOVARTIS SVERIGE AB	SE
Estradot 75 mikrogram/24 timmar, depotplåster	SE/H/1421/004	17558	NOVARTIS SVERIGE AB	SE
Estradot 100 microgramas/24 horas, adesivo transdérmico	SE/H/1421/005	3818986	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 100 microgramas/24 horas, adesivo transdérmico	SE/H/1421/005	3819083	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 100 microgramas/24 horas, adesivo transdérmico	SE/H/1421/005	3819182	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 25 microgramas/24 horas, adesivo transdérmico	SE/H/1421/001	5068986	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 25 microgramas/24 horas, adesivo transdérmico	SE/H/1421/001	5069083	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			S.A.	
Estradot 25 microgramas/24 horas, adesivo transdérmico	SE/H/1421/001	5069182	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 37,5 microgramas/24 horas, adesivo transdérmico	SE/H/1421/002	3818085	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 37,5 microgramas/24 horas, adesivo transdérmico	SE/H/1421/002	3818184	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 37,5 microgramas/24 horas, adesivo transdérmico	SE/H/1421/002	3818283	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 50 microgramas/24 horas, adesivo transdérmico	SE/H/1421/003	3818382	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 50 microgramas/24 horas, adesivo transdérmico	SE/H/1421/003	3818481	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 50 microgramas/24 horas, adesivo transdérmico	SE/H/1421/003	3818580	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 75 microgramas/24 horas, adesivo transdérmico	SE/H/1421/004	3818689	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 75 microgramas/24 horas, adesivo transdérmico	SE/H/1421/004	3818788	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 75 microgramas/24 horas, adesivo transdérmico	SE/H/1421/004	3818887	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 37,5 mikrog/24 tuntia depotlaastari	SE/H/1421/002	16729	NOVARTIS FINLAND OY	FI
Estradot 100 mikrog/24 timmar depotplåster	SE/H/1421/005	16732	NOVARTIS FINLAND OY	FI
Estradot 100 mikrog/24 tuntia depotlaastari	SE/H/1421/005	16732	NOVARTIS FINLAND OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estradot 25 microgramos/24 horas, parche transdérmico	SE/H/1421/001	66.103	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot 25 mikrog/24 tuntia depotlaastari	SE/H/1421/001	19049	NOVARTIS FINLAND OY	FI
Estradot 25 mikrog/24 timmar depotplåster	SE/H/1421/001	19049	NOVARTIS FINLAND OY	FI
Estradot 37,5 mikrog/24 timmar depotplåster	SE/H/1421/002	16729	NOVARTIS FINLAND OY	FI
Estradot 37,5 microgramos/24 horas, parche transdérmico	SE/H/1421/002	64.707	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot 50 microgramos/24 horas, parche transdérmico	SE/H/1421/003	64.704	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot 50 mikrog/24 tuntia depotlaastari	SE/H/1421/003	16730	NOVARTIS FINLAND OY	FI
Estradot 50 mikrog/24 timmar depotplåster	SE/H/1421/003	16730	NOVARTIS FINLAND OY	FI
Estradot 75 mikrog/24 tuntia depotlaastari	SE/H/1421/004	16731	NOVARTIS FINLAND OY	FI
Estradot 75 mikrog/24 timmar depotplåster	SE/H/1421/004	16731	NOVARTIS FINLAND OY	FI
Estradot 75 microgramos/24 horas, parche transdérmico	SE/H/1421/004	64.705	NOVARTIS FARMACÉUTICA S.A.	ES
Vivelle Dot 37,5 mikrogram/24 timer, depotplaster	SE/H/1421/002	32852	NOVARTIS HEALTHCARE A/S	DK
Vivelle Dot 50 mikrogram/24 timer, depotplaster	SE/H/1421/003	32853	NOVARTIS HEALTHCARE A/S	DK
Vivelle Dot 75 mikrogram/24 timer, depotplaster	SE/H/1421/004	32854	NOVARTIS HEALTHCARE A/S	DK
Vivelle Dot 100 mikrogram/24 timer, depotplaster	SE/H/1421/005	32855	NOVARTIS HEALTHCARE A/S	DK
Vivelle Dot 25 mikrogram/24 timer, depotplaster	SE/H/1421/001	36054	NOVARTIS HEALTHCARE A/S	DK
Estradot® 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PL 23860/0011	NOVARTIS IRELAND LIMITED	XI
Estradot® 100 Mikrogramm /24 Stunden - transdermale Pflaster	SE/H/1421/005	1-24343	NOVARTIS PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estradot® 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PL 23860/0007	NOVARTIS IRELAND LIMITED	XI
Estradot® 25 Mikrogramm/24 Stunden - transdermale Pflaster	SE/H/1421/001	1-27019	NOVARTIS PHARMA GMBH	AT
Estradot® 37,5 Mikrogramm/24 Stunden - transdermale Pflaster	SE/H/1421/002	1-24340	NOVARTIS PHARMA GMBH	AT
Estradot® 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PL 23860/0008	NOVARTIS IRELAND LIMITED	XI
Estradot® 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PL 23860/0009	NOVARTIS IRELAND LIMITED	XI
Estradot® 50 Mikrogramm/24 Stunden - transdermale Pflaster	SE/H/1421/003	1-24341	NOVARTIS PHARMA GMBH	AT
Estradot® 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PL 23860/0010	NOVARTIS IRELAND LIMITED	XI
Estradot® 75 Mikrogramm /24 Stunden - transdermale Pflaster	SE/H/1421/004	1-24342	NOVARTIS PHARMA GMBH	AT
VIVELLEDOT 100 microgrammes/24 heures, dispositif transdermique	SE/H/1421/005	34009 358 589 2 4	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 100 microgrammes/24 heures, dispositif transdermique	SE/H/1421/005	34009 358 590 0 6	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 100 microgrammes/24 heures, dispositif transdermique	SE/H/1421/005	34009 358 591 7 4	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	34009 365 573 0 7	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	34009 365 574 7 5	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	34009 365 575 3 6	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	34009 358 579 7 2	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	34009 358 580 5 4	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	34009 358 581 1 5	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	34009 358 582 8 3	NOVARTIS PHARMA S.A.S.	FR

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VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	34009 358 583 4 4	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	34009 358 584 0 5	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	34009 358 585 7 3	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	34009 358 586 3 4	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	34009 358 588 6 3	NOVARTIS PHARMA S.A.S.	FR
Vivelle dot 100 míkrógrömm/24 klst., forðaplástur	SE/H/1421/005	IS/1/01/043/04	NOVARTIS HEALTHCARE A/S	IS
Vivelle dot 25 míkrógrömm/24 klst., forðaplástur	SE/H/1421/001	IS/1/04/005/01	NOVARTIS HEALTHCARE A/S	IS
Vivelle dot 37,5 míkrógrömm/24 klst., forðaplástur	SE/H/1421/002	IS/1/01/043/01	NOVARTIS HEALTHCARE A/S	IS
Vivelle dot 50 míkrógrömm/24 klst., forðaplástur	SE/H/1421/003	IS/1/01/043/02	NOVARTIS HEALTHCARE A/S	IS
Vivelle dot 75 míkrógrömm/24 klst., forðaplástur	SE/H/1421/004	IS/1/01/043/03	NOVARTIS HEALTHCARE A/S	IS
Estradot 100 mikrog/24 timer depotplaster	SE/H/1421/005	01-7396	NOVARTIS NORGE AS	NO
Estradot 25 mikrog/24 timer depotplaster	SE/H/1421/001	03-2344	NOVARTIS NORGE AS	NO
Estradot 37,5 mikrog/24 timer depotplaster	SE/H/1421/002	01-7393	NOVARTIS NORGE AS	NO
Estradot 50 mikrog/24 timer depotplaster	SE/H/1421/003	01-7394	NOVARTIS NORGE AS	NO
Estradot 75 mikrog/24 timer depotplaster	SE/H/1421/004	01-7395	NOVARTIS NORGE AS	NO
Estradot 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PA0896/010/004	NOVARTIS IRELAND LIMITED	IE
Estradot 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PA0896/010/005	NOVARTIS IRELAND LIMITED	IE
Estradot 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PA0896/010/001	NOVARTIS IRELAND LIMITED	IE

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Estradot 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PA0896/010/002	NOVARTIS IRELAND LIMITED	IE
Estradot 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PA0896/010/003	NOVARTIS IRELAND LIMITED	IE
Estradot 100 mikrograma/24 sata, transdermalni flaster	SE/H/1421/005	HR-H-660461417-01	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 25 mikrograma/24 sata, transdermalni flaster	SE/H/1421/001	HR-H-259673389-01	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-01	NOVARTIS HRVATSKA D.O.O.	HR
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT

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tabletes				
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Estrofem® 2 mg film-coated tablets	not available	PA 218/50/1	NOVO NORDISK A/S	IE
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 653 7 4	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 916 2 8	BAYER HEALTHCARE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 743 6 9	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 915 6 7	BAYER HEALTHCARE	FR
Divigel 0,1%, 0,5 mg/0,5 g, żel	not available	4476	ORION CORPORATION	PL
Divigel 0,1%, 1,0 mg/1,0 g, żel	not available	4477	ORION CORPORATION	PL
Progynova® TS 100 micrograms/24 hours Transdermal Patch	IE/H/0843/002	PL 00010/0561	BAYER PLC	XI
Climara Forte 100 micrograms/24 hours Transdermal Patch	IE/H/0843/002	PA 1410/15/2	BAYER LTD	IE
Climara 100 microgrammi/24 h cerotti transdermici	IE/H/0843/002	030183040	BAYER SPA	IT
Climara 100 microgrammi/24 h cerotti transdermici	IE/H/0843/002	030183038	BAYER SPA	IT
Progynova® TS 50 micrograms/24 hours Transdermal Patch	IE/H/0843/001	PL 00010/0560	BAYER PLC	XI
Climara 50 micrograms/24 hours Transdermal Patch	IE/H/0843/001	PA 1410/15/1	BAYER LTD	IE
Climara 50 microgrammi/24 h cerotti transdermici	IE/H/0843/001	030183014	BAYER SPA	IT
Climara 50 microgrammi/24 h cerotti transdermici	IE/H/0843/001	030183026	BAYER SPA	IT
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
heures (7,5 mg/16,5 cm2), dispositif transdermique				
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermique				
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
Progynova®21 2 mg, überzogene Tablette	not available	6930615.01.00	JENAPHARM GMBH & CO KG	DE
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
DERMESTRIL 25 microgrammes/24 heures, dispositif transdermique	not available	34009 340 906 6 0	ROTTAPHARM SAS	FR
DERMESTRIL 50 microgrammes/24 heures, dispositif transdermique	not available	34009 340 904 3 1	ROTTAPHARM SAS	FR
DERMESTRIL 100 microgrammes/24 heures, dispositif transdermique	not available	34009 340 903 7 0	ROTTAPHARM SAS	FR
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
Estrofem 1 mg Filmtabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem 1 mg - Filmtabletten	DK/H/0117/001	1-22071	NOVO NORDISK PHARMA GMBH	AT
Estrofem 2 mg - Filmtabletten	not available	16 889	NOVO NORDISK PHARMA GMBH	AT
Estrofem 1 mg, filmomhulde tabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrifam 2 mg Filmtabletten	DK/H/0117/002	40226.01.00	NOVO NORDISK PHARMA GMBH	DE
Estrifam 1 mg Filmtabletten	DK/H/0117/001	40226.00.00	NOVO NORDISK PHARMA GMBH	DE
Estrofem 1 mg kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem 1 mg filmovertrukne tabletter	DK/H/0117/001	17461	NOVO NORDISK A/S	DK
Estrofem 2 mg filmovertrukne tabletter	DK/H/0117/002	06697	NOVO NORDISK A/S	DK
Estrofem 1 mg filmuhúðaðar töflur	DK/H/0117/001	IS/1/03/031/01	NOVO NORDISK A/S	IS
Estrofem 2 mg filmuhúðaðar töflur	DK/H/0117/002	IS/1/03/031/02	NOVO NORDISK A/S	IS
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
Estrofem 1 mg Filmtabletten	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
Progynova 1 mg, omhulde tabletten	not available	RVG 05861	BAYER BV	NL
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
DIVIGEL 0,5 mg geel DIVIGEL 1 mg	not available	195998	ORION CORPORATION	EE

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geel				
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Progynova®21 mite 1 mg, überzogene Tablette	not available	6930615.00.00	JENAPHARM GMBH & CO KG	DE
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Progynova 2 mg tabletter, drasjerte	not available	5183	BAYER AB	NO
Estrofem 2 mg filmomhulde tabletten	not available	RVG 09810	NOVO NORDISK B.V.	NL
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Climara-50, 50 µg/dobę (3,8 mg), system transdermalny	not available	4583	BAYER AG	PL
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
SYSTEN 50; 3,2 mg, system transdermalny, plaster	not available	R/1692	THERAMEX IRELAND LIMITED	PL
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Gynokadin Dosiergel 0,6 mg/g Gel	not available	52958.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Estrofem 1 mg Filmtabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem 1 mg - Filmtabletten	DK/H/0117/001	1-22071	NOVO NORDISK PHARMA GMBH	AT
Estrofem 2 mg - Filmtabletten	not available	16 889	NOVO NORDISK PHARMA GMBH	AT
Estrofem 1 mg, filmomhulde tabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrifam 2 mg Filmtabletten	DK/H/0117/002	40226.01.00	NOVO NORDISK PHARMA GMBH	DE
Estrifam 1 mg Filmtabletten	DK/H/0117/001	40226.00.00	NOVO NORDISK PHARMA GMBH	DE
Estrofem 1 mg kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem 1 mg filmovertrukne tabletter	DK/H/0117/001	17461	NOVO NORDISK A/S	DK
Estrofem 2 mg filmovertrukne tabletter	DK/H/0117/002	06697	NOVO NORDISK A/S	DK
Estrofem 1 mg filmuhúðaðar töflur	DK/H/0117/001	IS/1/03/031/01	NOVO NORDISK A/S	IS
Estrofem 2 mg filmuhúðaðar töflur	DK/H/0117/002	IS/1/03/031/02	NOVO NORDISK A/S	IS
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
Estrofem 1 mg Filmtabletten	DK/H/0117/001	2009020176	NOVO NORDISK PHARMA SA	LU
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Estrofem, 1 mg õhukese polümeerikattega tabletid	not available	204898	NOVO NORDISK A/S	EE
Estrofem, 2 mg õhukese polümeerikattega tabletid	not available	204798	NOVO NORDISK A/S	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagirux 10 microgram vaginal tablets	HU/H/0632/001	MA1031/01001	GEDEON RICHTER PLC.	MT
Vagirux 10 micrograms vaginal tablets	HU/H/0632/001	PL 04854/0184	GEDEON RICHTER PLC.	XI
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/01	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/02	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogrami vaginālās tabletes	HU/H/0632/001	20-0097	GEDEON RICHTER PLC.	LV
Vagirux 10 mikrogramov vaginálne tablety	HU/H/0632/001	56/0184/20-S	GEDEON RICHTER PLC.	SK
Rewellfem	HU/H/0632/001	62712	GEDEON RICHTER PLC.	DK
Vagirux, 10 mikrogrammi vaginaaltabletid	HU/H/0632/001	1010320	GEDEON RICHTER PLC.	EE
Vagirux 10 mikrogramů vaginální tablety	HU/H/0632/001	54/192/19-C	GEDEON RICHTER PLC.	CZ
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/001	GEDEON RICHTER PLC.	LT
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/002	GEDEON RICHTER PLC.	LT
Vagirux 10 mikrograma tablete za rodnicu	HU/H/0632/001	HR-H-853360202	GEDEON RICHTER PLC.	HR
Вагирукс 10 микрограма вагинални таблетки	HU/H/0632/001	20200165	GEDEON RICHTER PLC.	BG
ESTRADIOL RICHTER 10 microgame comprimate vaginale	HU/H/0632/001	13484/2020/01	GEDEON RICHTER PLC.	RO
ESTRADIOL RICHTER 10 microgame comprimate vaginale	HU/H/0632/001	13484/2020/02	GEDEON RICHTER PLC.	RO
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 0 3	GEDEON RICHTER PLC.	FR
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 1 0	GEDEON RICHTER PLC.	FR
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801816	GEDEON RICHTER PLC.	PT
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801824	GEDEON RICHTER PLC.	PT
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	19-12867	GEDEON RICHTER PLC.	NO
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856013	GEDEON RICHTER PLC.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856025	GEDEON RICHTER PLC.	IT
Vagirux 10 microgramos comprimidos vaginales	HU/H/0632/001	85352	GEDEON RICHTER PLC.	ES
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/001	GEDEON RICHTER PLC.	SI
Vagirux, 10 mikrogramów, tabletki dopochwowe	HU/H/0632/001	26164	GEDEON RICHTER PLC.	PL
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/002	GEDEON RICHTER PLC.	SI
Vagirux 10 mikrogrammaa emätinpuikko, tabletti	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 Mikrogramm Vaginaltabletten	HU/H/0632/001	2204223.00.00	GEDEON RICHTER PLC.	DE
Vagirux 10 microgram vaginal tablets.	HU/H/0632/001	PA1330/028/001	GEDEON RICHTER PLC.	IE
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	59171	GEDEON RICHTER PLC.	SE
VAGIRUX 10 μικρογραμμάρια κολπικά δισκία	HU/H/0632/001	20658/12-03-2021	GEDEON RICHTER PLC.	GR
Vagirux 10 microgram vaginal tablets	HU/H/0632/001	MA1031/01001	GEDEON RICHTER PLC.	MT
Vagirux 10 micrograms vaginal tablets	HU/H/0632/001	PL 04854/0184	GEDEON RICHTER PLC.	XI
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/01	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/02	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogrami vaginālās tabletes	HU/H/0632/001	20-0097	GEDEON RICHTER PLC.	LV
Vagirux 10 mikrogramov vaginálne tablety	HU/H/0632/001	56/0184/20-S	GEDEON RICHTER PLC.	SK
Rewellfem	HU/H/0632/001	62712	GEDEON RICHTER PLC.	DK
Vagirux, 10 mikrogrammi vaginaaltabletid	HU/H/0632/001	1010320	GEDEON RICHTER PLC.	EE
Vagirux 10 mikrogramů vaginální tablety	HU/H/0632/001	54/192/19-C	GEDEON RICHTER PLC.	CZ
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/001	GEDEON RICHTER PLC.	LT
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/002	GEDEON RICHTER PLC.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagirux 10 mikrograma tablete za rodnicu	HU/H/0632/001	HR-H-853360202	GEDEON RICHTER PLC.	HR
Вагирукс 10 микрограма вагинални таблетки	HU/H/0632/001	20200165	GEDEON RICHTER PLC.	BG
ESTRADIOL RICHTER 10 microgrames comprimate vaginale	HU/H/0632/001	13484/2020/01	GEDEON RICHTER PLC.	RO
ESTRADIOL RICHTER 10 microgrames comprimate vaginale	HU/H/0632/001	13484/2020/02	GEDEON RICHTER PLC.	RO
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 0 3	GEDEON RICHTER PLC.	FR
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 1 0	GEDEON RICHTER PLC.	FR
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801816	GEDEON RICHTER PLC.	PT
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801824	GEDEON RICHTER PLC.	PT
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	19-12867	GEDEON RICHTER PLC.	NO
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856013	GEDEON RICHTER PLC.	IT
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856025	GEDEON RICHTER PLC.	IT
Vagirux 10 microgramos comprimidos vaginales	HU/H/0632/001	85352	GEDEON RICHTER PLC.	ES
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/001	GEDEON RICHTER PLC.	SI
Vagirux, 10 mikrogramów, tabletki dopochwowe	HU/H/0632/001	26164	GEDEON RICHTER PLC.	PL
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/002	GEDEON RICHTER PLC.	SI
Vagirux 10 mikrogrammaa emätinpuikko, tabletti	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 Mikrogramm Vaginaltabletten	HU/H/0632/001	2204223.00.00	GEDEON RICHTER PLC.	DE
Vagirux 10 microgram vaginal tablets.	HU/H/0632/001	PA1330/028/001	GEDEON RICHTER PLC.	IE
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	59171	GEDEON RICHTER PLC.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VAGIRUX 10 μικρογραμμάρια κολπικά δισκία	HU/H/0632/001	20658/12-03-2021	GEDEON RICHTER PLC.	GR
Estrofem 2 mg filmsko obložene tablete	not available	H/97/00590/001	NOVO NORDISK A/S	SI
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
vaginaux				
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel	not available	18714	ORION CORPORATION	DK
Divigel	not available	18715	ORION CORPORATION	DK
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
spray, solution				
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Estrofem 1mg filmom obalené tablety	not available	56/0303/91-C/S	NOVO NORDISK A/S	SK
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 653 7 4	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 916 2 8	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 351 743 6 9	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	34009 318 915 6 7	BAYER HEALTHCARE	FR
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
Progynova ® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova® 2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
Progynova® 2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
Femoston® mono 2 mg Filmtabletten	not available	26518.00.00	MYLAN HEALTHCARE GMBH	DE
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Estrofem 2 mg filmdragerade tabletter	not available	11466	NOVO NORDISK A/S	FI
Estrofem 2 mg kalvopäällysteiset	not available	11466	NOVO NORDISK A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletit				
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem, 2 mg, tabletki powlekane	not available	R/3307	NOVO NORDISK A/S	PL
DERMESTRIL 25 microgrammes/24 heures, dispositif transdermique	not available	34009 340 906 6 0	ROTTAPHARM SAS	FR
DERMESTRIL 50 microgrammes/24 heures, dispositif transdermique	not available	34009 340 904 3 1	ROTTAPHARM SAS	FR
DERMESTRIL 100 microgrammes/24 heures, dispositif transdermique	not available	34009 340 903 7 0	ROTTAPHARM SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(5 mg/11 cm²), dispositif transdermique				
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
Estraderm MX ®75	not available	PL 20011/0066	NORGINE PHARMACEUTICALS LIMITED	XI
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(5 mg/11 cm²), dispositif transdermique				
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Progynova mite-Dragees	not available	14.422	BAYER AUSTRIA GMBH	AT
Vagirux 10 microgram vaginal tablets	HU/H/0632/001	MA1031/01001	GEDEON RICHTER PLC.	MT
Vagirux 10 micrograms vaginal tablets	HU/H/0632/001	PL 04854/0184	GEDEON RICHTER PLC.	XI
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/01	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/02	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogrami vaginālās tabletes	HU/H/0632/001	20-0097	GEDEON RICHTER PLC.	LV
Vagirux 10 mikrogramov vaginálne tablety	HU/H/0632/001	56/0184/20-S	GEDEON RICHTER PLC.	SK
Rewellfem	HU/H/0632/001	62712	GEDEON RICHTER PLC.	DK
Vagirux, 10 mikrogrammi vaginaaltabletid	HU/H/0632/001	1010320	GEDEON RICHTER PLC.	EE
Vagirux 10 mikrogramů vaginální tablety	HU/H/0632/001	54/192/19-C	GEDEON RICHTER PLC.	CZ
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/001	GEDEON RICHTER PLC.	LT
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/002	GEDEON RICHTER PLC.	LT
Vagirux 10 mikrograma tablete za rodnicu	HU/H/0632/001	HR-H-853360202	GEDEON RICHTER PLC.	HR
Вагирукс 10 микрограма вагинални таблетки	HU/H/0632/001	20200165	GEDEON RICHTER PLC.	BG
ESTRADIOL RICHTER 10 microgram comprimate vaginale	HU/H/0632/001	13484/2020/01	GEDEON RICHTER PLC.	RO
ESTRADIOL RICHTER 10 microgram comprimate vaginale	HU/H/0632/001	13484/2020/02	GEDEON RICHTER PLC.	RO
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 0 3	GEDEON RICHTER PLC.	FR
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 1 0	GEDEON RICHTER PLC.	FR
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801816	GEDEON RICHTER PLC.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801824	GEDEON RICHTER PLC.	PT
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	19-12867	GEDEON RICHTER PLC.	NO
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856013	GEDEON RICHTER PLC.	IT
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856025	GEDEON RICHTER PLC.	IT
Vagirux 10 microgramos comprimidos vaginales	HU/H/0632/001	85352	GEDEON RICHTER PLC.	ES
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/001	GEDEON RICHTER PLC.	SI
Vagirux, 10 mikrogramów, tabletki dopochwowe	HU/H/0632/001	26164	GEDEON RICHTER PLC.	PL
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/002	GEDEON RICHTER PLC.	SI
Vagirux 10 mikrogrammaa emätinpuikko, tabletti	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 Mikrogramm Vaginaltabletten	HU/H/0632/001	2204223.00.00	GEDEON RICHTER PLC.	DE
Vagirux 10 microgram vaginal tablets.	HU/H/0632/001	PA1330/028/001	GEDEON RICHTER PLC.	IE
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	59171	GEDEON RICHTER PLC.	SE
VAGIRUX 10 μικρογραμμάρια κολπικά δισκία	HU/H/0632/001	20658/12-03-2021	GEDEON RICHTER PLC.	GR
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/01	GEDEON RICHTER PLC.	HU
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/02	GEDEON RICHTER PLC.	HU
Rewellfem 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	2020090260	GEDEON RICHTER PLC.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rewellfem 10 míkrógramma skeiðartöflur.	HU/H/0633/001	IS/1/20/086/01	GEDEON RICHTER PLC.	IS
Vagirux 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	RVG 124964	GEDEON RICHTER PLC.	NL
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	140403	GEDEON RICHTER PLC.	AT
Вагирукс 10 микрограма вагинални таблетки	HU/H/0633/001	20200165	GEDEON RICHTER PLC.	BG
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
oestraclin 0,6 mg/g gel	not available	59.577	SEID, S.A.	ES
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
DERMESTRIL-Septem 25 Mikrogramm/24 Stunden, transdermales Pflaster	DE/H/5763/001	46060.00.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL-Septem 50 Mikrogramm/24 Stunden, transdermales Pflaster	DE/H/5763/002	46060.01.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL-Septem 75 Mikrogramm/24 Stunden, transdermales Pflaster	DE/H/5763/003	46060.02.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL –Septem 50 microgrammi/24 ore, cerotto transdermico	DE/H/5763/002	029001068	ROTTAPHARM S.P.A.	IT
DERMESTRIL – Septem 25 microgrammi/24 ore, cerotto transdermico	DE/H/5763/001	029001043	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 75 microgrammi/24 ore, cerotto transdermico	DE/H/5763/003	029001082	ROTTAPHARM S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DERMESTRIL – Septem 25 microgrammi/24 ore, cerotto transdermico	DE/H/5763/001	029001056	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 50 microgrammi/24 ore, cerotto transdermico	DE/H/5763/002	029001070	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 75 microgrammi/24 ore, cerotto transdermico	DE/H/5763/003	029001094	ROTTAPHARM S.P.A.	IT
DERMESTRIL SEPTEM 25 microgrammes/24 heures, dispositif transdermique	DE/H/5763/001	34009 352 393 9 6	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL SEPTEM 25 microgrammes/24 heures, dispositif transdermique	DE/H/5763/001	34009 352 394 5 7	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL SEPTEM 50 microgrammes/24 heures, dispositif transdermique	DE/H/5763/002	34009 352 391 6 7	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL SEPTEM 50 microgrammes/24 heures, dispositif transdermique	DE/H/5763/002	34009 352 392 2 8	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL SEPTEM 75 microgrammes/24 heures, dispositif transdermique	DE/H/5763/003	34009 352 395 1 8	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL SEPTEM 75 microgrammes/24 heures, dispositif transdermique	DE/H/5763/003	34009 352 396 8 6	LABORATOIRES ROTTAPHARM SARL	FR
DERMESTRIL-Septem 25 microgramas/24 horas sistema transdérmico	DE/H/5763/001	2970382	BGP PRODUCTS UNIPESSOAL, LDA.	PT
DERMESTRIL-Septem 25 microgramas/24 horas sistema transdérmico	DE/H/5763/001	2970481	BGP PRODUCTS UNIPESSOAL, LDA.	PT
DERMESTRIL-Septem 50 microgramas/24 horas sistema transdérmico	DE/H/5763/002	2970580	BGP PRODUCTS UNIPESSOAL, LDA.	PT
DERMESTRIL-Septem 50	DE/H/5763/002	2970689	BGP PRODUCTS UNIPESSOAL,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
microgramas/24 horas sistema transdérmico			LDA.	
DERMESTRIL-Septem 75 microgramas/24 horas sistema transdérmico	DE/H/5763/003	2970788	BGP PRODUCTS UNIPESSOAL, LDA.	PT
DERMESTRIL-Septem 75 microgramas/24 horas sistema transdérmico	DE/H/5763/003	2970887	BGP PRODUCTS UNIPESSOAL, LDA.	PT
SYSTEN 50; 3,2 mg, system transdermalny, plaster	not available	R/1692	THERAMEX IRELAND LIMITED	PL
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko,	UK/H/2176/001	26990	NOVO NORDISK A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletti				
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
oestraclin 0,6 mg/g gel	not available	59.577	SEID, S.A.	ES
ESTRADERM® MATRIX 25 microgramos/24 horas parches transdérmicos	not available	59.153	MERUS LABS LUXCO II S.À R.L.	ES
ESTRADERM® MATRIX 50 microgramos/24 horas parches transdérmicos	not available	59.154	MERUS LABS LUXCO II S.À R.L.	ES
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
Zumenon 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
ZUMENON 2 mg Filmtabletten	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
Zumenon 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
ZUMENON 2 mg Filmtabletten	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
PROGYNOVA 2 mg, comprimé enrobé	not available	34009 318 923 9 7	BAYER HEALTHCARE	FR
PROGYNOVA 2 mg, comprimé enrobé	not available	34009 351 651 4 5	BAYER HEALTHCARE	FR
OROMONE 1 mg, comprimé pelliculé	not available	34009 351 167 5 8	MYLAN MEDICAL SAS	FR
OROMONE 2 mg, comprimé pelliculé	not available	34009 342 436 7 7	MYLAN MEDICAL SAS	FR
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
ESTRADERM MX 100 mcg/die cerotto transdermico	not available	031773031	MERUS LABS LUXCO II S.À R.L.	IT
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
Estrofem, 2 mg, tabletki powlekane	not available	R/3307	NOVO NORDISK A/S	PL
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
SYSTEN 50; 3,2 mg, system	not available	R/1692	THERAMEX IRELAND LIMITED	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermalny, plaster				
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 665 0 6	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 664 4 5	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 666 7 4	MERUS LABS LUXCO II S.À R.L.	FR
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
PROVAMES 2 mg, comprimé pelliculé	not available	34009 354 794 0 2	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 2 mg, comprimé pelliculé	not available	34009 354 793 4 1	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 2 mg, comprimé pelliculé	not available	34009 337 341 1 4	MERUS LABS LUXCO II S.À R.L.	FR
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
vaginaux				
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά δισκία	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estring® 2 mg Vaginalinsert Estradiol	not available	29279.00.00	PFIZER PHARMA GMBH	DE
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
Oestring 7,5 mikrogram/24 timmar vaginalinlägg	not available	11783	PFIZER AB	SE
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
SYSTEN 50; 3,2 mg, system transdermalny, plaster	not available	R/1692	THERAMEX IRELAND LIMITED	PL
Estrofem 2 mg filmsko obložene tablete	not available	H/97/00590/001	NOVO NORDISK A/S	SI
ESTRAMON Gel 0,5 mg	not available	39910.00.00	HEXAL AG	DE
SisareÒ Gel mono 0,5 mg	not available	39909.00.00	ORION CORPORATION	DE
Sisare® Gel mono 1 mg	not available	39909.01.00	ORION CORPORATION	DE
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
Progynon 2 mg dragerade tabletter	not available	8424	BAYER AB	SE
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
Systen 100, pleisters voor transdermaal gebruik 100 microgram/24 uur	not available	RVG 19259	THERAMEX IRELAND LIMITED	NL
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lenzetto 1,53 mg/ spray, transdermal spray, solution	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermales Spray, Lösung				
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
Estraderm MX® 50	not available	PL 20011/0065	NORGINE PHARMACEUTICALS LIMITED	XI
Systen 50, pleisters voor transdermaal gebruik 50 microgram/24 uur	not available	RVG 16080	THERAMEX IRELAND LIMITED	NL
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Климара 50 микрограма/24 часа	not available	9900139	BAYER AG	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
трансдермален пластир				
Zumenon 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
ZUMENON 2 mg Filmtabletten	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
Zumenon 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
ZUMENON 2 mg Filmtabletten	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
Estradiol 10 micrograms vaginal tablets	not available	PL 12762/0619	MERCURY PHARMACEUTICALS LTD.	XI
Zumenon, omhulde tabletten 2 mg	not available	RVG 15462	MYLAN HEALTHCARE B.V.	NL
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
Estrofem, 1 mg õhukese polümeerikattega tabletid	not available	204898	NOVO NORDISK A/S	EE
Estrofem, 2 mg õhukese polümeerikattega tabletid	not available	204798	NOVO NORDISK A/S	EE
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
heures, dispositif transdermique				
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Естрофем 2 mg филмирани таблетки	not available	9900365	NOVO NORDISK A/S	BG
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
Progynova 2 mg, omhulde tabletten	not available	RVG 05311	BAYER BV	NL
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Estrofem 2 mg Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg potahované tablety	not available	56/303/91-C	NOVO NORDISK A/S	CZ
Estrofem 1 mg potahované tablety	not available	56/499/99-C	NOVO NORDISK A/S	CZ
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO II S.À R.L.	IT
Progynova 1 mg, omhulde tabletten	not available	RVG 05861	BAYER BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ESTRING 7,5 mikrograma na 24 ur vaginalni dostavni sistem	FR/H/0689/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	SI
ESTRING 2 mg, système de diffusion vaginal	FR/H/0689/001	34009 275 661 8 9	PFIZER HOLDING FRANCE	FR
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	FR/H/0689/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	FR/H/0689/001	042840013	PFIZER ITALIA S.R.L.	IT
Vagirux 10 microgram vaginal tablets	HU/H/0632/001	MA1031/01001	GEDEON RICHTER PLC.	MT
Vagirux 10 micrograms vaginal tablets	HU/H/0632/001	PL 04854/0184	GEDEON RICHTER PLC.	XI
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/01	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogramm hüvelytabletta	HU/H/0632/001	OGYI-T-23727/02	GEDEON RICHTER PLC.	HU
Vagirux 10 mikrogrami vaginālās tabletes	HU/H/0632/001	20-0097	GEDEON RICHTER PLC.	LV
Vagirux 10 mikrogramov vaginálne tablety	HU/H/0632/001	56/0184/20-S	GEDEON RICHTER PLC.	SK
Rewellfem	HU/H/0632/001	62712	GEDEON RICHTER PLC.	DK
Vagirux, 10 mikrogrammi vaginaaltabletid	HU/H/0632/001	1010320	GEDEON RICHTER PLC.	EE
Vagirux 10 mikrogramů vaginální tablety	HU/H/0632/001	54/192/19-C	GEDEON RICHTER PLC.	CZ
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/001	GEDEON RICHTER PLC.	LT
VAGIRUX 10 mikrogramų makšties tabletės	HU/H/0632/001	LT/1/20/4609/002	GEDEON RICHTER PLC.	LT
Vagirux 10 mikrograma tablete za rodnicu	HU/H/0632/001	HR-H-853360202	GEDEON RICHTER PLC.	HR
Вагирукс 10 микрограма вагинални таблетки	HU/H/0632/001	20200165	GEDEON RICHTER PLC.	BG
ESTRADIOL RICHTER 10 microgram comprimate vaginale	HU/H/0632/001	13484/2020/01	GEDEON RICHTER PLC.	RO
ESTRADIOL RICHTER 10 microgram comprimate vaginale	HU/H/0632/001	13484/2020/02	GEDEON RICHTER PLC.	RO
VAGIRUX 10 microgrammes, comprimé vaginal	HU/H/0632/001	34009 302 121 0 3	GEDEON RICHTER PLC.	FR
VAGIRUX 10 microgrammes,	HU/H/0632/001	34009 302 121 1 0	GEDEON RICHTER PLC.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimé vaginal				
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801816	GEDEON RICHTER PLC.	PT
Formyra 10 microgramas comprimidos vaginais	HU/H/0632/001	5801824	GEDEON RICHTER PLC.	PT
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	19-12867	GEDEON RICHTER PLC.	NO
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856013	GEDEON RICHTER PLC.	IT
Vagirux 10 microgrammi compresse vaginali	HU/H/0632/001	048856025	GEDEON RICHTER PLC.	IT
Vagirux 10 microgramos comprimidos vaginales	HU/H/0632/001	85352	GEDEON RICHTER PLC.	ES
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/001	GEDEON RICHTER PLC.	SI
Vagirux, 10 mikrogramów, tabletki dopochwowe	HU/H/0632/001	26164	GEDEON RICHTER PLC.	PL
Vagirux 10 mikrogramov vaginalne tablete	HU/H/0632/001	H/20/02764/002	GEDEON RICHTER PLC.	SI
Vagirux 10 mikrogrammaa emätinpuikko, tabletti	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	37077	GEDEON RICHTER PLC.	FI
Vagirux 10 Mikrogramm Vaginaltabletten	HU/H/0632/001	2204223.00.00	GEDEON RICHTER PLC.	DE
Vagirux 10 microgram vaginal tablets.	HU/H/0632/001	PA1330/028/001	GEDEON RICHTER PLC.	IE
Vagirux 10 mikrogram vaginaltabletter	HU/H/0632/001	59171	GEDEON RICHTER PLC.	SE
VAGIRUX 10 μικρογραμμάρια κολπικά δισκία	HU/H/0632/001	20658/12-03-2021	GEDEON RICHTER PLC.	GR
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel	not available	18714	ORION CORPORATION	DK
Divigel	not available	18715	ORION CORPORATION	DK
Elleste Solo MX 80 micrograms Transdermal Patch	not available	PL 46302/0168	MYLAN PRODUCTS LIMITED	XI
Elleste Solo MX 40 micrograms Transdermal Patch	not available	PL 46302/0167	MYLAN PRODUCTS LIMITED	XI
Oesclim 25, 5 mg, 25	not available	7615	MYLAN HEALTHCARE SP.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mikrogramów/24 godziny, system transdermalny			Z.O.O.	
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermique				
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 678 8 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 665 3 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 667 6 2	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 668 2 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 669 9 1	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 679 4 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 670 7 3	MYLAN MEDICAL SAS	FR
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	34009 344 664 7 2	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 508-98	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 116-17	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 117-85	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures	FR/H/0109/003	34009 338 118-46	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(10 mg/22 cm2), dispositif transdermique				
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 119-07	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 345 509-59	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 120-96	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	34009 338 115-56	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 109 5 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 110 3 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 113 2 7	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 512 6 0	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 114 9 5	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 338 108 9 4	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif transdermique	FR/H/0109/001	34009 345 510 3 1	MYLAN MEDICAL SAS	FR
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm²), dispositif	FR/H/0109/001	34009 338 112 6 6	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermique				
ESTRAMON Gel 0,5 mg	not available	39910.00.00	HEXAL AG	DE
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Progynova®21 mite 1 mg, überzogene Tablette	not available	6930615.00.00	JENAPHARM GMBH & CO KG	DE
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
DERMESTRIL 25 microgrammi/24 ore cerotti transdermici	not available	029001017	ROTTAPHARM S.P.A.	IT
DERMESTRIL 50 microgrammi/24 ore cerotti transdermici	not available	029001029	ROTTAPHARM S.P.A.	IT
DERMESTRIL 100 microgrammi/24 ore cerotti transdermici	not available	029001031	ROTTAPHARM S.P.A.	IT
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
Estraderm MX® 50	not available	PL 20011/0065	NORGINE PHARMACEUTICALS LIMITED	XI
Estraderm MX ®100	not available	PL 20011/0067	NORGINE PHARMACEUTICALS LIMITED	XI
Lenzetto 1,53 mg/pulverização solução para pulverização transdérmica	not available	5734751	GEDEON RICHTER PLC.	PT
Lenzetto 1,53 mg/pulverização solução para pulverização transdérmica	not available	5734769	GEDEON RICHTER PLC.	PT
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/02	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/02	ORION CORPORATION	HU
Divigel 0,5 mg/dózis gél	not available	OGYI-T-6229/01	ORION CORPORATION	HU
Divigel 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Progynova® 1 mg dragerade tabletter	not available	5836	BAYER OY	FI
Progynova® 1 mg dragerade tabletter	not available	5836	BAYER OY	FI
Progynova® 1 mg päällystetyt tabletit	not available	5836	BAYER OY	FI
Progynova® 1 mg päällystetyt tabletit	not available	5836	BAYER OY	FI
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
ESTRADERM® MATRIX 100 microgramos/24 horas parches transdérmicos	not available	59.155	MERUS LABS LUXCO II S.À R.L.	ES
Zumenon 2 mg tabletti, kalvopäällysteinen	not available	13398	MYLAN HEALTHCARE B.V.	FI
Zumenon 2 mg filmdragerad tablett	not available	13398	MYLAN HEALTHCARE B.V.	FI
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
ESTRADERM MX 50 mcg/die cerotto transdermico	not available	031773029	MERUS LABS LUXCO II S.À R.L.	IT
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
Estrofem 2 mg filmomhulde tabletten	not available	RVG 09810	NOVO NORDISK B.V.	NL
Evorel 50 micrograms per 24 hours Transdermal Patch	not available	PA22668/008/001	THERAMEX IRELAND LIMITED	IE
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	THERAMEX IRELAND LIMITED	BE
Климара 50 микрограма/24 часа трансдермален пластир	not available	9900139	BAYER AG	BG
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
spray, solution				
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
ZUMENON 1 mg Filmtabletten	not available	BE233317	MYLAN EPD BVBA/SPRL	BE
ZUMENON 2 mg Filmtabletten	not available	BE198721	MYLAN EPD BVBA/SPRL	BE
ZUMENON 1 mg comprimés pelliculés	not available	BE233317	MYLAN EPD BVBA/SPRL	BE
ZUMENON 2 mg comprimés pelliculés	not available	BE198721	MYLAN EPD BVBA/SPRL	BE
ZUMENON, 1 mg filmomhulde tabletten	not available	BE233317	MYLAN EPD BVBA/SPRL	BE
ZUMENON, 2 mg filmomhulde tabletten	not available	BE198721	MYLAN EPD BVBA/SPRL	BE
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER AB	NO
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651-2 6	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650-6 5	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546.5 7	PIERRE FABRE MEDICAMENT	FR
Progynova 2 mg compresse rivestite	not available	021226016	BAYER SPA	IT
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Progynova 1 mg comprimidos recubiertos	not available	47.814	BAYER HISPANIA SL	ES
Progynova®21 2 mg, überzogene Tablette	not available	6930615.01.00	JENAPHARM GMBH & CO KG	DE
SisareÒ Gel mono 0,5 mg	not available	39909.00.00	ORION CORPORATION	DE
Sisare® Gel mono 1 mg	not available	39909.01.00	ORION CORPORATION	DE
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	MYLAN HEALTHCARE SP. Z.O.O.	PL
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	MYLAN HEALTHCARE SP. Z.O.O.	PL
SYSTEN 50; 3,2 mg, system transdermalny, plaster	not available	R/1692	THERAMEX IRELAND LIMITED	PL
Vagidonna 10 mikrog emätinpuikko, tabletti	SE/H/1959/001	36960	SANDOZ A/S	FI
Vagidonna 10 mikrogram vaginaltablett	SE/H/1959/001	59006	SANDOZ A/S	SE
Vagidonna 10 mikrogram vaginaltablett	SE/H/1959/001	19-12777	SANDOZ A/S	NO
Vagidonna 10 míkróg skeiðartafla	SE/H/1959/001	IS/1/20/039/01	SANDOZ A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zumenon 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
ZUMENON 2 mg Filmtabletten	not available	2009070399	MYLAN EPD BVBA/SPRL	LU
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	XI
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	XI
Sandrena 0,5 mg gel	DK/H/0105/001	032991010/M	ORION CORPORATION	IT
Sandrena 0,5 mg gel	DK/H/0105/001	032991022/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991034/M	ORION CORPORATION	IT
SANDRENA® 1 MG GEL	DK/H/0105/002	032991046/M	ORION CORPORATION	IT
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/001	15734	ORION CORPORATION	DK
Ercostrol, gel, enkeltdosisbeholder	DK/H/0105/002	15735	ORION CORPORATION	DK
Divigel 0,5 mg gel	DK/H/0105/001	13136	ORION CORPORATION	SE
Divigel 1 mg gel	DK/H/0105/002	13296	ORION CORPORATION	SE
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 401-9	ORION CORPORATION	FR
Delidose 0,5 mg, gel en sachet-dose	DK/H/0105/001	342 402-5	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 399-4	ORION CORPORATION	FR
Delidose 1 mg, gel en sachet-dose	DK/H/0105/002	342 400-2	ORION CORPORATION	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 665 0 6	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 664 4 5	MERUS LABS LUXCO II S.À R.L.	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 666 7 4	MERUS LABS LUXCO II S.À R.L.	FR
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
oestraclin 0,6 mg/g gel	not available	59.577	SEID, S.A.	ES
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
ESTREVA 0,1 %, gel	FR/H/0133/001	34009 339 130 8 3	THERAMEX IRELAND LIMITED	FR
ESTREVA 0,1 % GEL	FR/H/0133/001	BE204057	THERAMEX IRELAND LIMITED	BE
Estreva® 0,1 %, gel	FR/H/0133/001	BE204057	THERAMEX IRELAND LIMITED	BE
Estreva® 0,1% gel.	FR/H/0133/001	BE204057	THERAMEX IRELAND LIMITED	BE
Estreva® 0,1 %, gel	FR/H/0133/001	2000075881	THERAMEX IRELAND LIMITED	LU
ESTREVA 0,1 % GEL	FR/H/0133/001	2000075881	THERAMEX IRELAND LIMITED	LU
ESTREVA® 0,1% GEL.	FR/H/0133/001	038008025	THERAMEX IRELAND LIMITED	IT
ESTREVA® 0,1% GEL.	FR/H/0133/001	038008013	THERAMEX IRELAND LIMITED	IT
ESTREVA 0.1% gel	FR/H/0133/001	2902880	THERAMEX IRELAND LIMITED	PT
ESTREVA 0.1% gel	FR/H/0133/001	3419389	THERAMEX IRELAND LIMITED	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estrena 0,1 % geeli	FR/H/0133/001	13671	THERAMEX IRELAND LIMITED	FI
Estrena 0,1 % geeli	FR/H/0133/001	13671	THERAMEX IRELAND LIMITED	FI
EVOPAD 100 µg/24 h parches transdérmicos	not available	61.702	THERAMEX IRELAND LIMITED	ES
EVOPAD 75 µg/24 h parches transdérmicos	not available	61.701	THERAMEX IRELAND LIMITED	ES
EVOPAD 25 µg/24 h parches transdérmicos	not available	61.700	THERAMEX IRELAND LIMITED	ES
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
Divigel 0,5 mg gelis	not available	LT/1/98/0238/001	ORION CORPORATION	LT
Divigel 1 mg gelis	not available	LT/1/98/0238/002	ORION CORPORATION	LT
Estraderm MX® 25	not available	PL 20011/0064	NORGINE PHARMACEUTICALS LIMITED	XI
Systen 75, pleisters voor transdermaal gebruik 75 microgram/24 uur	not available	RVG 19258	THERAMEX IRELAND LIMITED	NL
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Gynokadin Dosiergel 0,6 mg/g Gel	not available	31673.00.00	BESINS HEALTHCARE GERMANY GMBH	DE
Estreva; 0,1%, żel	not available	9606	THERAMEX IRELAND LIMITED	PL
Климара 50 микрограма/24 часа трансдермален пластир	not available	9900139	BAYER AG	BG
Estradiol fem JENAPHARM® 1,53 mg Tabletten	not available	41162.00.00	JENAPHARM GMBH & CO KG	DE
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
Progynon 1 mg dragerade tabletter	not available	8423	BAYER AB	SE
OROMONE 1 mg, comprimé pelliculé	not available	34009 351 167 5 8	MYLAN MEDICAL SAS	FR
OROMONE 2 mg, comprimé pelliculé	not available	34009 342 436 7 7	MYLAN MEDICAL SAS	FR
Progynova 1 mg, omhulde tabletten	not available	RVG 05861	BAYER BV	NL
Fematab 1 mg film-coated tablet	not available	PA 2010/11/1	MYLAN IRE HEALTHCARE LIMITED	IE
Fematab 2 mg Film-coated tablet	not available	PA 2010/11/2	MYLAN IRE HEALTHCARE LIMITED	IE
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
Fematab 1 mg film-coated tablet	not available	PA 2010/11/1	MYLAN IRE HEALTHCARE	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			LIMITED	
Fematab 2 mg Film-coated tablet	not available	PA 2010/11/2	MYLAN IRE HEALTHCARE LIMITED	IE
Zumenon 2 mg comprimido revestido por película	not available	2420289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	XI
Zumenon® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	XI
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV
Estrofem 2 mg apvalkotās tabletes	not available	00-0580	NOVO NORDISK A/S	LV
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal spray, solution	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pulvérisation transdermique				
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek,	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
transdermalno pršilo, raztopina				
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/spray, transdermal spray, solution	HU/H/0361/001	PL 04854/0130	GEDEON RICHTER PLC.	XI
Lenzetto 1.53 mg/spray, transdermal spray, solution	HU/H/0361/001	PA1330/017/001	GEDEON RICHTER PLC.	IE
Lenzetto 1,53 mg/ spray, transdermal spray, solution	HU/H/0361/001	MA1031/00101	GEDEON RICHTER PLC.	MT
Lenzetto 1,53 mg/úðaskammt, úði til notkunar um húð	HU/H/0361/001	IS/1/15/098/01	GEDEON RICHTER PLC.	IS
Lenzetto, 1,53 mg/dawkę, aerozol przezskórny, roztwór	HU/H/0361/001	22626	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Lenzetto	HU/H/0361/001	53385	GEDEON RICHTER PLC.	DK
Lenzetto 1,53 mg/dávka transdermální sprej, roztok	HU/H/0361/001	56/381/15-C	GEDEON RICHTER PLC.	CZ
Lenzetto 1,53 mg/spray transdermal spray, oppløsning	HU/H/0361/001	13-9878	GEDEON RICHTER PLC.	NO
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/01	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/annuses, transdermaalne sprei, lahus	HU/H/0361/001	881715	GEDEON RICHTER PLC.	EE
Lenzetto 1,53 mg po potisku, transdermalni sprej, otopina	HU/H/0361/001	HR-H-173338597	GEDEON RICHTER PLC.	HR
Lenzetto 1,53 mg/dávka transdermálna roztoková aerodisperzia	HU/H/0361/001	56/0382/15-S	GEDEON RICHTER PLC.	SK
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermales Spray, Lösung	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/001	GEDEON RICHTER PLC.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lenzetto 1,53 mg/suihke transdermaalisumute, liuos	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	31925	GEDEON RICHTER PLC.	FI
Lenzetto 1,53 mg/izsmidzinājumā transdermāls aerosols, šķīdums	HU/H/0361/001	15-0279	GEDEON RICHTER PLC.	LV
Лензето 1,53 mg/впръскване, трансдермален спрей, разтвор	HU/H/0361/001	20150382	GEDEON RICHTER PLC.	BG
Lenzetto 1,53 mg/Sprühstoß transdermales Spray, Lösung	HU/H/0361/001	91153.00.00	GEDEON RICHTER PLC.	DE
Lenzetto 1,53 mg/dosis, spray voor transdermaal gebruik, oplossing	HU/H/0361/001	RVG 114597	GEDEON RICHTER PLC.	NL
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205018	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/sprayning, transdermal spray, lösning	HU/H/0361/001	50431	GEDEON RICHTER PLC.	SE
Lenzetto 1,53 mg/adag transzdermális oldatos spray	HU/H/0361/001	OGYI-T-22890/02	GEDEON RICHTER PLC.	HU
Lenzetto 1,53 mg/išpurškime transderminis purškalas (tirpalas)	HU/H/0361/001	LT/1/15/3791/002	GEDEON RICHTER PLC.	LT
Lenzetto 1,53 mg/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	2016040053	GEDEON RICHTER PLC.	LU
Lenzetto 1,53 mg/dosis, solución para pulverización transdérmica	HU/H/0361/001	80608	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/nebulizzazione, spray transdermico, soluzione	HU/H/0361/001	043205020	GEDEON RICHTER PLC.	IT
Lenzetto 1,53 mg/ψεκασμό, διαδερμικό εκνέφωμα, διάλυμα	HU/H/0361/001	71987	GEDEON RICHTER PLC.	GR
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/002	GEDEON RICHTER PLC.	SI
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lenzetto 1,53 mg/doză spray transdermic, soluţie	HU/H/0361/001	13913/2021/02	GEDEON RICHTER ROMÂNIA S.A.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SYSTEN 50; 3,2 mg, system transdermalny, plaster	not available	R/1692	THERAMEX IRELAND LIMITED	PL
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
Estrofem 2 mg filmsko obložene tablete	not available	H/97/00590/001	NOVO NORDISK A/S	SI
Estring 7,5 míkróg/24 klst. skeiðarinnlegg.	not available	920143	PFIZER APS	IS
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/01	GEDEON RICHTER PLC.	HU
Rewellfem 10 mikrogramm hüvelytabletta	HU/H/0633/001	OGYI-T-23728/02	GEDEON RICHTER PLC.	HU
Rewellfem 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	BE571715	GEDEON RICHTER PLC.	BE
Rewellfem 10 microgrammes comprimés vaginaux	HU/H/0633/001	2020090260	GEDEON RICHTER PLC.	LU
Rewellfem 10 míkrógramma skeiðartöflur.	HU/H/0633/001	IS/1/20/086/01	GEDEON RICHTER PLC.	IS
Vagirux 10 microgram tabletten voor vaginaal gebruik	HU/H/0633/001	RVG 124964	GEDEON RICHTER PLC.	NL
Rewellfem 10 Mikrogramm Vaginaltabletten	HU/H/0633/001	140403	GEDEON RICHTER PLC.	AT
Вагирукс 10 микрограма вагинални таблетки	HU/H/0633/001	20200165	GEDEON RICHTER PLC.	BG
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrograma tablete za rodnicu.	UK/H/2176/001	HR-H-793020012	NOVO NORDISK A/S	HR
Vagifem 10 mikrogramov vaginalne	UK/H/2176/001	H/10/01589/001	NOVO NORDISK A/S	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tablete				
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PL 04668/0237	NOVO NORDISK A/S	XI
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgram, tabletten voor vaginaal gebruik	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	BE
Vagifem10 μικρογραμμάρια κολπικά δισκία.	UK/H/2176/001	20906	NOVO NORDISK A/S	CY
Vagifem 10 mikrogramů vaginální tablety	UK/H/2176/001	54/233/10-C	NOVO NORDISK A/S	CZ
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/002	NOVO NORDISK A/S	LT
Vagifem, 10 mikrogrammi vaginaaltabletid	UK/H/2176/001	676710	NOVO NORDISK A/S	EE
Vagifem 10 mikrog emätinpuikko, tabletti	UK/H/2176/001	26990	NOVO NORDISK A/S	FI
Vagifem 10 mikrogramu makšties tabletes	UK/H/2176/001	LT/1/94/0157/003	NOVO NORDISK A/S	LT
Vagifem 10 mikrogramov vaginalne tablete	UK/H/2176/001	H/10/01589/002	NOVO NORDISK A/S	SI
Vagifem 10 mikrogrami vaginalas tabletes	UK/H/2176/001	10-0093	NOVO NORDISK A/S	LV
Vagifem 10 microgram, filmomhulde tabletten voor vaginaal gebruik	UK/H/2176/001	RVG 103834	NOVO NORDISK A/S	NL
Vagifem, 10 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem, vaginaltabletter 10 mikrogram	UK/H/2176/001	44279	NOVO NORDISK A/S	DK
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	41707	NOVO NORDISK A/S	SE
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	MA104/00302	NOVO NORDISK A/S	MT
Vagifem 10 microgrammes, comprimés vaginaux	UK/H/2176/001	0642/10120049	NOVO NORDISK PHARMA SA	LU
Vagifem 10 mikrogram vaginaltabletter	UK/H/2176/001	08-6376	NOVO NORDISK A/S	NO
Vagifem10 μικρογραμμάρια κολπικά	UK/H/2176/001	54165/6-7-2016	NOVO NORDISK HELLAS LTD.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
δισκία				
Vagifem 10 míkrógrömm skeiðartafla	UK/H/2176/001	IS/1/09/059/01	NOVO NORDISK A/S	IS
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/02	NOVO NORDISK A/S	HU
Vagifem 10 mikrogramm hüvelytabletta	UK/H/2176/001	OGYI-T-4465/03	NOVO NORDISK A/S	HU
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894020	NOVO NORDISK A/S	IT
Vagifem 10 microgrammi compresse vaginali.	UK/H/2176/001	028894032	NOVO NORDISK A/S	IT
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	PL
DERMESTRIL 25 µg/24 Stunden, transdermales Pflaster	not available	30327.00.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL 50 µg/24 Stunden, transdermales Pflaster	not available	30327.01.00	MEDA PHARMA GMBH & CO. KG	DE