5 May 2017
EMA/269084/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: estradiol (without cream/balm/emulsion for application in the female genital area)

Procedure no.: PSUSA/00010440/201608

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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
Estrofem 1 mg filmom obložene tablete	DK/H/0117/001	UP/I-530-09/14-01/21	NOVO NORDISK A/S	HR
Estrofem 2 mg filmom obložene tablete	DK/H/0117/002	UP/I-530-09/14-01/22	NOVO NORDISK A/S	HR
Estrofem 1 mg Filmtabletten	DK/H/0117/001	BE187171	NOVO NORDISK PHARMA SA	BE
Estrofem 1 mg Filmtabletten	DK/H/0117-001	0642/09020176	NOVO NORDISK PHARMA SA	LU
Estrofem 1 mg filmdragerade tabletter	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 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

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 kalvopäällysteiset tabletit	DK/H/0117/001	12623	NOVO NORDISK A/S	FI
Estrofem, filmovertrukne tabletter	DK/H/0117/001	17461	NOVO NORDISK A/S	DK
Estrofem, filmovertrukne tabletter	DK/H/0117/002	06697	NOVO NORDISK A/S	DK
Estrofem 1 mg, comprimés pelliculés	DK/H/0117/001	0642/09020176	NOVO NORDISK PHARMA SA	LU
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, comprimidos revestidos por película	DK/H/0117/001	2717486	ISDIN LDA	PT
Estrofem 1 mg, comprimidos revestidos por película	DK/H/0117/001	4315388	ISDIN LDA	PT
Estrofem 2 mg, comprimidos revestidos por película	DK/H/0117/002	5122486	ISDIN 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
Estrofem 2 mg, comprimidos revestidos por película	DK/H/0117/002	5122585	ISDIN LDA	PT
Gynokadin® Dosiergel	not available	52958.00.00	DR. KADE / BESINS PHARMA GMBH	DE
Vagifem 25 μg κολπικά δισκία επικαλυμμένα με υμένιο	not available	2440/19-01-2015	NOVO NORDISK HELLAS LTD.	GR
Vagifem 25 Mikrogramm Vaginaltabletten	not available	17354.00.00	NOVO NORDISK PHARMA GMBH	DE
Vagifem, 25 mikrogrammi vaginaaltabletid	not available	247199	NOVO NORDISK A/S	EE
Vagifem 25 mikrogramm hüvelytabletta	not available	OGYI-T-4465/01	NOVO NORDISK A/S	HU
Vagifem 25 microgrammi compresse vaginali film-rivestite	not available	028894018	NOVO NORDISK A/S	IT
Vagifem	not available	13048	NOVO NORDISK A/S	DK
Vagifem, κολπικά δισκία επικαλυμμένα με υμένιο	not available	16409	NOVO NORDISK HELLAS LTD.	CY

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 25 míkrógrömm skeiðartöflur	not available	920192	NOVO NORDISK A/S	IS
Vagifem 0,025 mg comprimidos vaginais	not available	2243285	ISDIN LDA	PT
DERMESTRIL-Septem 25 Mikrogramm/24 Stunden, transdermales Pflaster	UK/H/302/001	46060.00.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL-Septem 50 Mikrogramm/24 Stunden, transdermales Pflaster	UK/H/0302/002	46060.01.00	MEDA PHARMA GMBH & CO. KG	DE
DERMESTRIL-Septem 75 Mikrogramm/24 Stunden, transdermales Pflaster	UK/H/0302/003	46060.02.00	MEDA PHARMA GMBH & CO. KG	DE
Sisare 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
Divigel 0,1 % gel	not available	11445	ORION OYJ	FI
Divigel 0,1 % geeli	not available	11445	ORION OYJ	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
Oestrogel 0,06% gel	not available	BE111544	BESINS HEALTHCARE BENELUX	BE
Oestrogel 0,06% gel	not available	BE154831	BESINS HEALTHCARE BENELUX	BE
Естрогел 0,06 % гел	not available	20020604	LABORATOIRES BESINS INTERNATIONAL	BG
OESTROGEL	not available	56/058/97-C	LABORATOIRES BESINS INTERNATIONAL	CZ
OESTROGEL 0,06 POUR CENT, gel pour application cutanée en tube	not available	316 892-9	LABORATOIRES BESINS INTERNATIONAL	FR
ESTRODOSE 0,75 mg gel	not available	029542026	LABORATOIRES BESINS INTERNATIONAL	IT
Oestrogel 0,6 mg/g gel	not available	2465/2010/01	LABORATOIRES BESINS INTERNATIONAL	RO
Oestrogel 0,06% gel	not available	BE154831	BESINS HEALTHCARE BENELUX	BE
Oestrogel 0,06% gel	not available	BE111544	BESINS HEALTHCARE BENELUX	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
OESTRODOSE 0,06 POUR CENT, gel pour application cutanée en flacon avec pompe doseuse	not available	338 292-4	LABORATOIRES BESINS INTERNATIONAL	FR
OESTRODOSE 0,06 POUR CENT, gel pour application cutanée en flacon avec pompe doseuse	not available	364 346-0	LABORATOIRES BESINS INTERNATIONAL	FR
Oestrogel 750 micrograms Gel Pump-Pack	not available	PA 1341/2/1	BESINS HEALTHCARE	IE
Oestrogel Pump-Pack	not available	PL 28397/0002	BESINS HEALTHCARE	UK
Estrofem 2 mg, Filmtabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, Filmtabletten	not available	0642/03/12/7920	NOVO NORDISK PHARMA SA	LU
Estrofem 2 mg filmdragerade tabletter	not available	11466	NOVO NORDISK A/S	FI
Estrofem 2 mg film-coated tablets	not available	PA 218/50/1	NOVO NORDISK A/S	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
Estrofem 2 mg Filmtabletten	not available	16 889	NOVO NORDISK PHARMA GMBH	AT
Estrofem 2 mg, filmomhulde tabletten	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Estrofem 2 mg, comprimés pelliculés	not available	BE156545	NOVO NORDISK PHARMA SA	BE
Естрофем® 2 mg филмирани таблетки	not available	9900365	NOVO NORDISK A/S	BG
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
Estrofem, 1 mg õhukese polümeerikattega tabletid	not available	204898	NOVO NORDISK A/S	EE
Estrofem 1 mg plėvele dengtos tabletės	not available	LT/1/94/0156/001	NOVO NORDISK A/S	LT
Estrofem 1 mg apvalkotās tabletes	not available	00-0579	NOVO NORDISK A/S	LV

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 õhukese polümeerikattega tabletid	not available	204798	NOVO NORDISK A/S	EE
Estrofem 2 mg kalvopäällysteiset tabletit	not available	11466	NOVO NORDISK A/S	FI
ESTROFEM 2 mg, comprimé pelliculé	not available	NL 12376	NOVO NORDISK	FR
ESTROFEM 1 mg, comprimé pelliculé	not available	NL 20525	NOVO NORDISK	FR
Estrofem 2 mg filmtabletta	not available	OGYI-T-5849/01	NOVO NORDISK A/S	HU
Estrofem 2 mg plėvele dengtos tabletės	not available	LT/1/94/0156/002	NOVO NORDISK A/S	LT
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
Estrofem mite, 1 mg, tabletki powlekane	not available	8232	NOVO NORDISK A/S	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
Estrofem, 2 mg, tabletki powlekane	not available	R/3307	NOVO NORDISK A/S	PL
Estrofem 1mg filmom obalené tablety	not available	56/0303/91-C/S	NOVO NORDISK A/S	SK
Estrofem 2 mg, comprimés pelliculés	not available	0642/03/12/7920	NOVO NORDISK PHARMA SA	LU
Estrofem 2 mg filmsko obložene tablete	not available	H/97/00590/001	NOVO NORDISK A/S	SI
Gynokadin® Gel	not available	25825.00.00	DR. KADE / BESINS PHARMA GMBH	DE
ESTRAMON Gel 1,0 mg	not available	39910.01.00	HEXAL AG	DE
ESTRAMON Gel 0,5 mg	not available	39910.00.00	HEXAL AG	DE
OESTROGEL	not available	30723 / 03-05-2012	ANGELINI PHARMA HELLAS S.A.	GR
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	8708164	MERUS LABS LUXCO S.À R.L	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
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	8708107	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	3273786	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	3071289	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	8708172	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	8708123	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 25; 0,75 mg; 25 µg/24 h; system transdermalny	not available	7028	MERUS LABS LUXCO S.À R.L	PL
ESTRADERM MX 50; 1,5 mg; 50 µg/24 h; system transdermalny	not available	7029	MERUS LABS LUXCO S.À R.L	PL
ESTRADERM MX 100; 3 mg; 100 µg/24 h; system transdermalny	not available	7030	MERUS LABS LUXCO S.À R.L	PL
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	3273885	MERUS LABS LUXCO S.À R.L	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
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	3152683	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 50 mcg/die cerotto transdermico	not available	031773029	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	8708180	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 100 mcg/die cerotto transdermico	not available	031773031	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	8708149	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	3273984	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	3152782	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX® 25	not available	PL 44374/0003	MERUS LABS LUXCO S.À R.L	UK

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
Estraderm MX® 50	not available	PL 44374/0004	MERUS LABS LUXCO S.À R.L	UK
Estraderm MX ®75	not available	PL 44374/0005	MERUS LABS LUXCO S.À R.L	UK
Estraderm MX ®100	not available	PL 44374/0006	MERUS LABS LUXCO S.À R.L	UK
ESTRADERM® MATRIX 25 microgramos/24 horas parches transdérmicos	not available	59.153	MERUS LABS LUXCO S.À R.L	ES
ESTRADERM® MATRIX 50 microgramos/24 horas parches transdérmicos	not available	59.154	MERUS LABS LUXCO S.À R.L	ES
ESTRADERM® MATRIX 100 microgramos/24 horas parches transdérmicos	not available	59.155	MERUS LABS LUXCO S.À R.L	ES
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 665 0 6	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 664 4 5	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 666 7 4	MERUS LABS LUXCO S.À R.L	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® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova® 1 mg dragerade tabletter	not available	5836	BAYER OY	FI
Progynova® 2 mg dragerade tabletter	not available	4881	BAYER OY	FI
Progynova®1 mg dragerade tabletter	not available	5836	BAYER OY	FI
Progynova 2 mg tabletter, drasjerte	not available	5183	BAYER PHARMA AG	NO
Progynova 1 mg tabletter, drasjerte	not available	5474	BAYER PHARMA AG	NO
PROGYNOVA 1 mg, comprimé enrobé	not available	351 653-7	BAYER HEALTHCARE	FR
PROGYNOVA 2 mg, comprimé enrobé	not available	318 922-2	BAYER HEALTHCARE	FR
PROGYNOVA 2 mg, comprimé enrobé	not available	318 923-9	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	318 916-2	BAYER HEALTHCARE	FR
PROGYNOVA 2 mg, comprimé enrobé	not available	351 652-0	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	351 743-6	BAYER HEALTHCARE	FR
PROGYNOVA 1 mg, comprimé enrobé	not available	318 915-6	BAYER HEALTHCARE	FR
PROGYNOVA 2 mg, comprimé enrobé	not available	351 651-4	BAYER HEALTHCARE	FR
Progynova 2 mg compresse rivestite	not available	021226016	BAYER SPA	IT
Progynon 2 mg dragerade tabletter	not available	8424	BAYER PHARMA AG	SE
Progynon 2 mg dragerade tabletter	not available	8424	BAYER PHARMA AG	SE
Progynon 1 mg dragerade tabletter	not available	8423	BAYER PHARMA AG	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
Progynova 2 mg, omhulde tabletten	not available	RVG 05311	BAYER BV	NL
Progynova 1 mg, omhulde tabletten	not available	RVG 05861	BAYER BV	NL
Progynova 1 mg comprimidos recubiertos	not available	47.814	BAYER HISPANIA SL	ES
Estradiol fem JENAPHARM® 1,53 mg Tabletten	not available	41162.00.00	JENAPHARM GMBH & CO KG	DE
Progynova-21, 2 mg, tabletki powlekane	not available	R/3041	BAYER PHARMA AG	PL
Estradiol fem JENAPHARM® 1,53 mg Tabletten	not available	3000328.00.00	JENAPHARM GMBH & CO KG	DE
Progynova®21 mite 1 mg, überzogene Tablette	not available	6930615.00.00	JENAPHARM GMBH & CO KG	DE
Progynon 1 mg dragerade tabletter	not available	8423	BAYER PHARMA AG	SE
Progynova®21 2 mg, überzogene Tablette	not available	6930615.01.00	JENAPHARM GMBH & CO KG	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
Progynova® 2mg	not available	PL 00010/0557	BAYER PLC	UK
Progynova® 1mg	not available	PL 00010/0556	BAYER PLC	UK
Progynova® 2mg	not available	MA513/03302	BAYER PLC	MT
Progynova mite-Dragees	not available	14.422	BAYER AUSTRIA GMBH	AT
Progynova®2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
Progynova ®1 mg päällystetyt tabletit	not available	5836	BAYER OY	FI
Progynova® 2 mg päällystetyt tabletit	not available	4881	BAYER OY	FI
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

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	PL 04854/0130	GEDEON RICHTER PLC.	UK
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, kutanspray, opløsning	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

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/doză spray transdermic, soluţie	HU/H/0361/001	8083/2015/01	GEDEON RICHTER ROMÂNIA S.A.	RO
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/dose, Solution pour pulvérisation transdermique	HU/H/0361/001	BE478426	GEDEON RICHTER PLC.	BE
Lenzetto 1,53 mg/dosis, transdermale spray, oplossing	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	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

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/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/doză spray transdermic, soluţie	HU/H/0361/001	8083/2015/02	GEDEON RICHTER ROMÂNIA S.A.	RO
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	711046-0	GEDEON RICHTER PLC.	ES
Lenzetto 1,53 mg/razpršek, transdermalno pršilo, raztopina	HU/H/0361/001	H/16/02233/001	GEDEON RICHTER PLC.	SI
oestraclin 0,6 mg/g gel	not available	59.577	SEID, S.A.	ES
ESTRAPATCH 40 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 651 2 6	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
ESTRADIOL PIERRE FABRE SANTE 40 microgrammes/ 24 heures, dispositif transdermique	FR/H/237/01	356 649-8	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 60 microgrammes/ 24 heures, dispositif transdermique	not available	34009 356 650 6 5	PIERRE FABRE MEDICAMENT	FR
ESTRADIOL PIERRE FABRE SANTE 80 microgrammes/ 24 heures, dispositif transdermique	FR/H/237/03	359.545.9	PIERRE FABRE MEDICAMENT	FR
ESTRADIOL PIERRE FABRE SANTE 60 microgrammes/ 24 heures, dispositif transdermique	FR/H/237/02	356 648-1	PIERRE FABRE MEDICAMENT	FR
ESTRAPATCH 80 microgrammes/ 24 heures, dispositif transdermique	not available	34009 359 546 5 7	PIERRE FABRE MEDICAMENT	FR
FEMSEVEN 75 75 microgrammi/24 ore, cerotto transdermico	UK/H/145/002	029966049	RATIOPHARM ITALIA S.R.L.	IT
FEMSEVEN 75 75 microgrammi/24 ore,	UK/H/145/002	029966037	RATIOPHARM 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
cerotto transdermico				
ESTREVA® gel 0,1%	not available	5148/2005/01	LABORATOIRE THERAMEX S.A.M.	RO
FemSeven 100, 100 microgram/24 hours, transdermal patch	UK/H/0145/003	PL 00289/1801	TEVA UK LIMITED	UK
FemSeven 75, 75 microgram/24 hours, transdermal patch	UK/H/0145/002	PL 00289/1800	TEVA UK LIMITED	UK
FemSete 75,75 microgramas / 24 horas, Sistema transdérmico	UK/H/0145/002	2818888	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
ESTREVA® 0.1% gel	not available	MA 017/00101	LABORATOIRE THERAMEX S.A.M.	MT
FemSete 75, 75 microgramas / 24 horas, Sistema transdérmico	UK/H/0145/002	2818987	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
FEMINOVA 75, 75 microgrammes par 24 heures, dispositifs transdermiques	UK/H/0145/002	BE 203332	TEVA PHARMA BELGIUM N.V./S.A	BE
FEMINOVA 75, 75 microgram per 24 uur, pleisters voor transdermaal gebruik	UK/H/0145/002	BE 203332	TEVA PHARMA BELGIUM N.V./S.A	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
Feminova® 75, 75 Mikrogramm/24 Stunden, transdermales Pflaster Estradiol	UK/H/0145/002	BE 203332	TEVA PHARMA BELGIUM N.V./S.A	BE
Estreva; 0,1%, żel	not available	9606	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Sandrena 0.5 mg gel	DK/H/0105/001	PL 27925/0015	ORION CORPORATION	UK
Sandrena 1 mg gel	DK/H/0105/002	PL 27925/0016	ORION CORPORATION	UK
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
DERMESTRIL®-Septem 25 microgrammes/24 heures, dispositif transdermique	UK/H/0302/001	BE207751	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL® 25 – 0,025 mg / 24uur, transdermale pleister	not available	BE171114	BESINS HEALTHCARE BENELUX	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
DERMESTRIL® 50 – 0,050 mg / 24uur, transdermale pleister	not available	BE171123	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL® 100 – 0,100 mg / 24uur, transdermale pleister	not available	BE171132	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL®-Septem 25 microgram/24uur Transdermale Pleister	UK/H/0302/001	BE207751	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL®-Septem 50 microgrammes/24 heures Dispositif transdermique	UK/H/0302/002	BE207767	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL®-Septem 75 microgrammes/24 heures, dispositif transdermique	UK/H/0302/003	BE207776	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL®-Septem 50 microgram/24uur Transdermale Pleister	UK/H/0302/002	BE207767	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL®-Septem 75 microgram/24uur Transdermale Pleister	UK/H/0302/003	BE207776	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL® 25 – 0,025 mg / 24h, dispositifs transdermiques	not available	BE171114	BESINS HEALTHCARE BENELUX	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
DERMESTRIL® 50 – 0,050 mg / 24h, dispositifs transdermiques	not available	BE171123	BESINS HEALTHCARE BENELUX	BE
DERMESTRIL® 100 – 0,100 mg / 24h, dispositifs transdermiques	not available	BE171132	BESINS HEALTHCARE BENELUX	BE
THAISSEPT 25 microgrammes/24 heures, dispositif transdermique	UK/H/0303/001	352 397-4	LABORATOIRES BESINS INTERNATIONAL	FR
THAIS SEPT 25 microgrammes/24 heures, dispositif transdermique	UK/H/0303/001	352 398-0	LABORATOIRES BESINS INTERNATIONAL	FR
THAISSEPT 50 microgrammes/24 heures, dispositif transdermique	UK/H/0303/002	352 399-7	LABORATOIRES BESINS INTERNATIONAL	FR
THAISSEPT 50 microgrammes/24 heures, dispositif transdermique	UK/H/0303/002	352 400-5	LABORATOIRES BESINS INTERNATIONAL	FR
THAISSEPT 75 microgrammes/24 heures, dispositif transdermique	UK/H/0303/003	352 401-1	LABORATOIRES BESINS INTERNATIONAL	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
THAISSEPT 75 microgrammes/24 heures, dispositif transdermique	UK/H/0303/003	352 402-8	LABORATOIRES BESINS INTERNATIONAL	FR
THAIS 25 microgrammes/24 heures, dispositif transdermique	not available	340 909-5	LABORATOIRES BESINS INTERNATIONAL	FR
THAIS 50 microgrammes/24 heures, dispositif transdermique	not available	340 908-9	LABORATOIRES BESINS INTERNATIONAL	FR
THAIS 100 microgrammes/24 heures, dispositif transdermique	not available	340 907-2	LABORATOIRES BESINS INTERNATIONAL	FR
Divigel 0.1% w/w Gel, 1 mg/dose	not available	PA 1327/2/1	ORION CORPORATION	IE
Divigel	not available	18714	ORION CORPORATION	DK
Divigel	not available	18715	ORION CORPORATION	DK
Vagifem 10 Mikrogramm Vaginaltabletten	UK/H/2176/001	BE369031	NOVO NORDISK PHARMA SA	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
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	UK
Vagifem 10 micrograms vaginal tablets	UK/H/2176/001	PA 218/30/2	NOVO NORDISK A/S	IE
Vagifem 10 microgramos comprimidos vaginales recubiertos	UK/H/2176/001	72171	ISDIN, S.A.	ES
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

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 μg 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

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 mikrogramów, tabletki dopochwowe	UK/H/2176/001	17259	NOVO NORDISK A/S	PL
Vagifem	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	MA 104/00302	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 0,010 mg comprimido vaginal	UK/H/2176/01/DC	5270319	ISDIN 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
Vagifem 0,010 mg comprimido vaginal	UK/H/2176/01/DC	5270327	ISDIN LDA	PT
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
Gynokadin® Gel	not available	52949.00.00	DR. KADE / BESINS 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

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 1 mg/dózis gél	not available	OGYI-T-6230/01	ORION CORPORATION	HU
Systen 50 Mikrogramm/24 h transdermales Pflaster	not available	BE 156204	JANSSEN-CILAG NV	BE
Evorel 25 mikrog/24 tuntia depotplåster	not available	11554	JANSSEN-CILAG OY	FI
Systen 50 microgrammes / 24h, dispositifs transdermiques	not available	BE 156204	JANSSEN-CILAG NV	BE
Systen 50 microgram/24h, pleisters voor transdermaal gebruik	not available	BE 156204	JANSSEN-CILAG NV	BE
Evorel 50 Patch	not available	18171	JANSSEN-CILAG INTERNATIONAL NV	CY
Evorel	not available	14769	JANSSEN-CILAG A/S	DK
Evorel	not available	17507	JANSSEN-CILAG A/S	DK
Evorel	not available	15858	JANSSEN-CILAG A/S	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
Evorel	not available	15857	JANSSEN-CILAG A/S	DK
Fematab 2 mg Film-coated tablet	not available	PA 2007/7/2	BGP PRODUCTS LTD.	IE
Fematab 1 mg Film-coated tablet	not available	PA 2007/7/1	BGP PRODUCTS LTD.	IE
ZUMENON 2 mg comprimés pelliculés	not available	2009070399	MYLAN EPD SPRL	LU
ZUMENON, 2 mg filmomhulde tabletten	not available	BE198721	MYLAN EPD SPRL	BE
ZUMENON, 1 mg filmomhulde tabletten	not available	BE233317	MYLAN EPD SPRL	BE
ZUMENON 1 mg Filmtabletten	not available	BE233317	MYLAN EPD SPRL	BE
ZUMENON 2 mg Filmtabletten	not available	BE198721	MYLAN EPD SPRL	BE
Zumenon® 1mg Film-coated Tablets	not available	PL 46302/0052	MYLAN PRODUCTS LIMITED	UK

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® 2mg Film-coated Tablets	not available	PL 46302/0053	MYLAN PRODUCTS LIMITED	UK
OESCLIM 25 microgrammes/24 heures (5 mg/11 cm2), dispositif transdermique	FR/H/0109/001	338 108-9	MYLAN MEDICAL SAS	FR
Oesclim 25, 5 mg, 25 mikrogramów/24 godziny, system transdermalny	not available	7615	BGP PRODUCTS POLAND SP. Z.O.O.	PL
OESCLIM 37,5 microgrammes/24 heures (7,5 mg/16,5 cm2), dispositif transdermique	FR/H/0109/002	344 664-7	MYLAN MEDICAL SAS	FR
OESCLIM 50 microgrammes/24 heures (10 mg/22 cm2), dispositif transdermique	FR/H/0109/003	338 115-5	MYLAN MEDICAL SAS	FR
Oesclim 50, 10 mg, 50 mikrogramów/24 godziny, system transdermalny	not available	7616	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Zumenon, omhulde tabletten 2 mg	not available	RVG 15462	BGP PRODUCTS B.V.	NL
ZUMENON 1 mg comprimés pelliculés	not available	BE233317	MYLAN EPD SPRL	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
Zumenon 2 mg comprimidos revestidos por película	not available	2420289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
OROMONE 1 mg, comprimé pelliculé	not available	351 167-5	MYLAN MEDICAL SAS	FR
Femoston® mono 2 mg Filmtabletten	not available	26518.00.00	MYLAN HEALTHCARE GMBH	DE
OROMONE 2 mg, comprimé pelliculé	not available	342 436-7	MYLAN MEDICAL SAS	FR
ZUMENON 2 mg comprimés pelliculés	not available	BE198721	MYLAN EPD SPRL	BE
Zumenon 2 mg tabletti, päällystetty	not available	13398	BGP PRODUCTS B.V.	FI
Meriestra 2 mg comprimidos recubiertos	not available	60.207	NOVARTIS FARMACÉUTICA S.A.	ES
Vivelle dot 100 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/005	BE232881	NOVARTIS PHARMA N.V.	BE
Vivelle dot 25 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/001	BE265106	NOVARTIS PHARMA N.V.	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
Vivelle dot 37,5 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/002	BE232854	NOVARTIS PHARMA N.V.	BE
Vivelle dot 50 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/003	BE232863	NOVARTIS PHARMA N.V.	BE
Vivelle dot 75 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/004	BE232872	NOVARTIS PHARMA N.V.	BE
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, transdermalni flaster	SE/H/1421/005	HR-H-660461417-04	NOVARTIS HRVATSKA D.O.O.	HR
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

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 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
Sequidot transdermalni flaster	SE/H/0149/002	HR-H-674150675-01	NOVARTIS HRVATSKA D.O.O.	HR
Sequidot transdermalni flaster	SE/H/0149/002	HR-H-674150675-02	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 100 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/005	RVG 24640	NOVARTIS PHARMA B.V.	NL
Estradot 25 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/001	RVG 27888	NOVARTIS PHARMA B.V.	NL
Estradot 37,5 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/002	RVG 24637	NOVARTIS PHARMA B.V.	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
Estradot 50 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/003	RVG 24638	NOVARTIS PHARMA B.V.	NL
Estradot 75 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/004	RVG 24639	NOVARTIS PHARMA B.V.	NL
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
Meriestra 1 mg comprimidos recubiertos	not available	60.206	NOVARTIS FARMACÉUTICA S.A.	ES
ESTALIS SEQUI cerotti transdermici	SE/H/0149/002	034209039	NOVARTIS FARMA 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
ESTALIS SEQUI cerotti transdermici	SE/H/0149/002	034209041	NOVARTIS FARMA S.P.A.	IT
ESTALIS SEQUI, adesivo transdérmico	SE/H/0149/002	5019526	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
ESTALIS SEQUI, adesivo transdérmico	SE/H/0149/002	5019534	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
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 S.A.	PT
Estradot 25 microgramas/24 horas, adesivo transdérmico	SE/H/1421/001	5069182	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	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
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

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 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510501	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510502	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510503	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510504	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510401	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510402	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510403	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510404	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510101	NOVARTIS (HELLAS) S.A.C.I.	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
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510102	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510103	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510104	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510201	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510202	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510203	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510204	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510301	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510302	NOVARTIS (HELLAS) S.A.C.I.	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
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510303	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510304	NOVARTIS (HELLAS) S.A.C.I.	GR
ESTALIS SEQUIDOT parche transdérmico	SE/H/0149/002	69.746	NOVARTIS FARMACÉUTICA S.A.	ES
Estardot 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
Estradot 25 microgramos/24 horas parches transdérmicos	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

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 37,5 mikrog/24 timmar depotplåster	SE/H/1421/002	16729	NOVARTIS FINLAND OY	FI
Estradot 37.5 microgramos/24 horas parches transdérmicos	SE/H/1421/002	64.707	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot 50 microgramos/24 horas parches transdérmicos	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
EstradotT 75 microgramos/24 horas parches transdérmicos	SE/H/1421/004	64.705	NOVARTIS FARMACÉUTICA S.A.	ES
Sequidot depotlaastari	SE/H/0149/002	22301	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
Sequidot transdermal patches	SE/H/0149/002	22301	NOVARTIS FINLAND OY	FI
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
MERIMONO® 1 MG FILMTABLETTEN	UK/H/0122/001	37986.00.00	NOVARTIS PHARMA GMBH	DE
Sequidot® 50/250 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/0149/002	66641.00.00	NOVARTIS PHARMA GMBH	DE
Estradot® 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PL 00101/0645	NOVARTIS PHARMACEUTICALS UK LIMITED	UK

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 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PA 13/110/4	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT® 100 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/005	51816.03.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 100 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/005	0135/11041060	NOVARTIS PHARMA GMBH	LU
ESTRADOT 100 MIKROGRAMM /24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/005	1-24343	NOVARTIS PHARMA GMBH	AT
ESTRADOT 100; 1,56 mg; 100 µg/24 h; system transdermalny	not available	9395	NOVARTIS POLAND SP. Z O. O.	PL
Estradot® 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PL 00101/0703	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PA 13/110/5	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT 25 MIKROGRAMM/24 STUNDEN -	SE/H/1421/001	1-27019	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
TRANSDERMALE PFLASTER				
ESTRADOT® 25 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/001	51816.04.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 25 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/001	0135/11041056	NOVARTIS PHARMA GMBH	LU
ESTRADOT® 37,5 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/002	51816.00.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 37,5 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/002	0135/11041057	NOVARTIS PHARMA GMBH	LU
ESTRADOT 37,5 MIKROGRAMM/24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/002	1-24340	NOVARTIS PHARMA GMBH	AT
Estradot 37,5; 0,585 mg; 37,5 µg/24 h; system transdermalny	not available	9392	NOVARTIS POLAND SP. Z O. O.	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
Estradot® 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PL 00101/0642	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PA 13/110/1	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
Estradot® 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PL 00101/0643	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PA 13/110/2	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT® 50 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/003	51816.01.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 50 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/003	0135/11041058	NOVARTIS PHARMA GMBH	LU
ESTRADOT 50 MIKROGRAMM/24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/003	1-24341	NOVARTIS PHARMA GMBH	AT
Estradot 50; 0,78 mg; 50 µg/24 h; system transdermalny	not available	9393	NOVARTIS POLAND SP. Z O. O.	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
Estradot® 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PL 00101/0644	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PA 13/110/3	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT® 75 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/004	51816.02.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 75 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/004	0135/11041059	NOVARTIS PHARMA GMBH	LU
ESTRADOT 75 MIKROGRAMM /24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/004	1-24342	NOVARTIS PHARMA GMBH	AT
Estradot 75; 1,17 mg; 75 µg/24 h; system transdermalny	not available	9394	NOVARTIS POLAND SP. Z O. O.	PL
VIVELLEDOT 100 microgrammes/24 heures, dispositif transdermique	SE/H/1421/005	3400935858924	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 100 microgrammes/24	SE/H/1421/005	3400935859006	NOVARTIS PHARMA S.A.S.	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				
VIVELLEDOT 100 microgrammes/24 heures, dispositif transdermique	SE/H/1421/005	3400935859174	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	3400936557307	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	3400936557475	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 25 microgrammes/24 heures, dispositif transdermique	SE/H/1421/001	3400936557536	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	3400935857972	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	3400935858054	NOVARTIS PHARMA S.A.S.	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
VIVELLEDOT 37,5 microgrammes/24 heures, dispositif transdermique	SE/H/1421/002	3400935858115	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	3400935858283	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	3400935858344	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 50 microgrammes/24 heures, dispositif transdermique	SE/H/1421/003	3400935858405	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	3400935858573	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	3400935858634	NOVARTIS PHARMA S.A.S.	FR
VIVELLEDOT 75 microgrammes/24 heures, dispositif transdermique	SE/H/1421/004	3400935858863	NOVARTIS PHARMA S.A.S.	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
CLIMAGEST 2mg film-coated tablet	UK/H/0127/001	PL 00101/0366	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Climaval® 1 mg film-coated tablet	UK/H/0122/001	PL 00101/0307	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Climaval® 2 mg film-coated tablet	UK/H/0122/002	PL 00101/0308	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Sequidot 2-Phasen transdermale Matrixpflaster	SE/H/0149/002	1-26854	NOVARTIS PHARMA GMBH	AT
Sequidot transdermal patches	SE/H/0149/002	21846	NOVARTIS SVERIGE AB	SE
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

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
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
Sequidot depotplaster	not available	04-3137	NOVARTIS NORGE AS	NO
MERIMONO® 2 MG FILMTABLETTEN	UK/H/0122/002	37986.01.00	NOVARTIS PHARMA GMBH	DE
VIVELLE DOT 100 MICROGRAMMES/24 HEURES, DISPOSITIF TRANSDERMIQUE	SE/H/1421/005	BE232881	NOVARTIS PHARMA N.V.	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
Vivelle dot 100 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/005	BE232881	NOVARTIS PHARMA N.V.	BE
VIVELLE DOT 25 MICROGRAMMES/24 HEURES, DISPOSITIF TRANSDERMIQUE	SE/H/1421/001	BE265106	NOVARTIS PHARMA N.V.	BE
Vivelle dot 25 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/001	BE265106	NOVARTIS PHARMA N.V.	BE
VIVELLE DOT 37,5 MICROGRAMMES/24 HEURES, DISPOSITIF TRANSDERMIQUE	SE/H/1421/002	BE232854	NOVARTIS PHARMA N.V.	BE
Vivelle dot 37,5 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/002	BE232854	NOVARTIS PHARMA N.V.	BE
VIVELLE DOT 50 MICROGRAMMES/24 HEURES, DISPOSITIF TRANSDERMIQUE	SE/H/1421/003	BE232863	NOVARTIS PHARMA N.V.	BE
Vivelle dot 50 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/003	BE232863	NOVARTIS PHARMA N.V.	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
VIVELLE DOT 75 MICROGRAMMES/24 HEURES, DISPOSITIF TRANSDERMIQUE	SE/H/1421/004	BE232872	NOVARTIS PHARMA N.V.	BE
Vivelle dot 75 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/004	BE232872	NOVARTIS PHARMA N.V.	BE
ESTALIS SEQUIDOT, διαδερμικό έμπλαστρο	SE/H/0149/002	273100101	NOVARTIS (HELLAS) S.A.C.I.	GR
ESTALIS SEQUIDOT, διαδερμικό έμπλαστρο	SE/H/0149/002	273100102	NOVARTIS (HELLAS) S.A.C.I.	GR
Mericomb 1 mg Filmtabletten	UK/H/0127/002	39490.00.00	NOVARTIS PHARMA GMBH	DE
Mericomb 2 mg Filmtabletten	UK/H/0127/001	37970.00.00	NOVARTIS PHARMA GMBH	DE
Climagest 1mg film-coated tablet	UK/H/0127/002	PL 00101/0328	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
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

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 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-01	NOVARTIS HRVATSKA D.O.O.	HR
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, transdermalni flaster	SE/H/1421/005	HR-H-660461417-04	NOVARTIS HRVATSKA D.O.O.	HR
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

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 50 mikrograma/24 sata, transdermalni flaster	SE/H/1421/003	HR-H-207838847-04	NOVARTIS HRVATSKA D.O.O.	HR
Estradot 100 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/005	RVG 24640	NOVARTIS PHARMA B.V.	NL
Estradot 25 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/001	RVG 27888	NOVARTIS PHARMA B.V.	NL
Estradot 37,5 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/002	RVG 24637	NOVARTIS PHARMA B.V.	NL
Estradot 50 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/003	RVG 24638	NOVARTIS PHARMA B.V.	NL
Estradot 75 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/1421/004	RVG 24639	NOVARTIS PHARMA B.V.	NL
Estradot 100 mikrogram/24 timmar, depotplåster	SE/H/1421/005	17559	NOVARTIS SVERIGE AB	SE
Estradot 25 mikrogram/24 timmar,	SE/H/1421/001	20333	NOVARTIS SVERIGE AB	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
depotplåster				
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 S.A.	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
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

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 75 microgramas/24 horas, adesivo transdérmico	SE/H/1421/004	3818887	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510501	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510502	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510503	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 25 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/001	253510504	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510401	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510402	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510403	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 100 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/005	253510404	NOVARTIS (HELLAS) S.A.C.I.	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
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510101	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510102	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510103	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 37,5 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/002	253510104	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510201	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510202	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510203	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 50 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/003	253510204	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510301	NOVARTIS (HELLAS) S.A.C.I.	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
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510302	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510303	NOVARTIS (HELLAS) S.A.C.I.	GR
Estradot 75 micrograms/24 hrs, διαδερμικό έμπλαστρο	SE/H/1421/004	253510304	NOVARTIS (HELLAS) S.A.C.I.	GR
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
Estradot 25 microgramos/24 horas parches transdérmicos	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

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 37.5 microgramos/24 horas parches transdérmicos	SE/H/1421/002	64.707	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot 50 microgramos/24 horas parches transdérmicos	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
EstradotT 75 microgramos/24 horas parches transdérmicos	SE/H/1421/004	64.705	NOVARTIS FARMACÉUTICA S.A.	ES
Estradot® 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PL 00101/0645	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 100 micrograms/24 hours, transdermal patch	SE/H/1421/005	PA 13/110/4	NOVARTIS PHARMACEUTICALS UK LIMITED	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
ESTRADOT® 100 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/005	51816.03.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 100 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/005	0135/11041060	NOVARTIS PHARMA GMBH	LU
ESTRADOT 100 MIKROGRAMM /24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/005	1-24343	NOVARTIS PHARMA GMBH	AT
ESTRADOT 100; 1,56 mg; 100 µg/24 h; system transdermalny	not available	9395	NOVARTIS POLAND SP. Z O. O.	PL
Estradot® 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PL 00101/0703	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 25 micrograms/24 hours, transdermal patch	SE/H/1421/001	PA 13/110/5	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT 25 MIKROGRAMM/24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/001	1-27019	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 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/001	51816.04.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 25 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/001	0135/11041056	NOVARTIS PHARMA GMBH	LU
ESTRADOT® 37,5 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/002	51816.00.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 37,5 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/002	0135/11041057	NOVARTIS PHARMA GMBH	LU
ESTRADOT 37,5 MIKROGRAMM/24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/002	1-24340	NOVARTIS PHARMA GMBH	AT
Estradot 37,5; 0,585 mg; 37,5 µg/24 h; system transdermalny	not available	9392	NOVARTIS POLAND SP. Z O. O.	PL
Estradot® 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PL 00101/0642	NOVARTIS PHARMACEUTICALS UK LIMITED	UK

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 37.5 micrograms/24 hours, transdermal patch	SE/H/1421/002	PA 13/110/1	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
Estradot® 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PL 00101/0643	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estradot 50 micrograms/24 hours, transdermal patch	SE/H/1421/003	PA 13/110/2	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT® 50 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/003	51816.01.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 50 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/003	0135/11041058	NOVARTIS PHARMA GMBH	LU
ESTRADOT 50 MIKROGRAMM/24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/003	1-24341	NOVARTIS PHARMA GMBH	AT
Estradot 50; 0,78 mg; 50 µg/24 h; system transdermalny	not available	9393	NOVARTIS POLAND SP. Z O. O.	PL
Estradot® 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PL 00101/0644	NOVARTIS PHARMACEUTICALS UK LIMITED	UK

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 75 micrograms/24 hours, transdermal patch	SE/H/1421/004	PA 13/110/3	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
ESTRADOT® 75 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/1421/004	51816.02.00	NOVARTIS PHARMA GMBH	DE
ESTRADOT® 75 MIKROGRAMM/24 STUNDEN TRANSDERMALES PFLASTER	SE/H/1421/004	0135/11041059	NOVARTIS PHARMA GMBH	LU
ESTRADOT 75 MIKROGRAMM /24 STUNDEN - TRANSDERMALE PFLASTER	SE/H/1421/004	1-24342	NOVARTIS PHARMA GMBH	AT
Estradot 75; 1,17 mg; 75 µg/24 h; system transdermalny	not available	9394	NOVARTIS POLAND SP. Z O. O.	PL
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

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 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 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
EPIESTROL 25 microgrammi/24 ore cerotti transdermici	not available	029000015	ROTTAPHARM S.P.A.	IT
EPIESTROL 50 microgrammi/24 ore cerotti transdermici	not available	029000027	ROTTAPHARM S.P.A.	IT
EPIESTROL 100 microgrammi/24 ore cerotti transdermici	not available	029000039	ROTTAPHARM S.P.A.	IT
Elleste Solo MX 40 micrograms Transdermal Patch	not available	PL 15142/0060	MEDA PHARMACEUTICALS LTD	UK

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
Elleste Solo MX 80 micrograms Transdermal Patch	not available	PL 15142/0059	MEDA PHARMACEUTICALS LTD	UK
EPIESTROL –Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0303/001	029000041	ROTTAPHARM S.P.A.	IT
EPIESTROL –Septem 50 microgrammi/24 ore, cerotto transdermico	UK/H/0303/002	029000066	ROTTAPHARM S.P.A.	IT
EPIESTROL –Septem 75 microgrammi/24 ore, cerotto transdermico	UK/H/0303/003	029000080	ROTTAPHARM S.P.A.	IT
DERMESTRIL25 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
DERMESTRIL®-Septem 25 micrograms/24hours Transdermal Patch	UK/H/0302/001	PL 18219/0001	ROTTAPHARM LIMITED	UK
DERMESTRIL®-Septem 75 micrograms/24hours Transdermal Patch	UK/H/0302/003	PL 18219/0003	ROTTAPHARM LIMITED	UK
DERMESTRIL®-Septem 50 micrograms/24hours Transdermal Patch	UK/H/0302/002	PL 18219/0002	ROTTAPHARM LIMITED	UK

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
EPIESTROL®-Septem 25 micrograms/24hours Transdermal Patch	UK/H/0303/001	PL 18219/0007	ROTTAPHARM LIMITED	UK
EPIESTROL®-Septem 50 micrograms/24hours Transdermal Patch	UK/H/0303/002	PL 18219/0008	ROTTAPHARM LIMITED	UK
EPIESTROL®-Septem 75 micrograms/24hours Transdermal Patch	UK/H/0303/003	PL 18219/0009	ROTTAPHARM LIMITED	UK
Dermestril 50 mikrogramm/24 óra transzdermális tapasz	not available	OGYI-T-5116/02	ROTTAPHARM S.P.A.	HU
Dermestril 25 mikrogramm/24 óra transzdermális tapasz	not available	OGYI-T-5116/01	ROTTAPHARM S.P.A.	HU
Dermestril 100 mikrogramm/24 óra transzdermális tapasz	not available	OGYI-T-5116/03	ROTTAPHARM S.P.A.	HU
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

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 50 microgrammi/24 ore, cerotto transdermico	UK/H/0302/002	029001068	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0302/001	029001043	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 75 microgrammi/24 ore, cerotto transdermico	UK/H/0302/003	029001082	ROTTAPHARM S.P.A.	IT
DERMESTRIL®-Septem 25 micrograms/24hours Transdermal Patch	UK/H/0302/001	PA 868/2/4	ROTTAPHARM LIMITED	IE
DERMESTRIL®-Septem 50 micrograms/24hours Transdermal Patch	UK/H/0302/002	PA 868/2/5	ROTTAPHARM LIMITED	IE
DERMESTRIL®-Septem 75 micrograms/24hours Transdermal Patch	UK/H/0302/003	PA 868/2/6	ROTTAPHARM LIMITED	IE
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455681	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455582	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 25 25microgramas/24h Sistema transdérmico	not available	2455483	LABORATÓRIOS DELTA, S.A.	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 25 25microgramas/24h Sistema transdérmico	not available	2455384	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2456085	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455988	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455889	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 50 50microgramas/24h Sistema transdérmico	not available	2455780	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL-Septem 50 microgramas/24 horas sistema transdérmico	UK/H/0302/002	2970580	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 100 100 microgramas/24h Sistema transdérmico	not available	2456481	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 100 100 microgramas/24h Sistema transdérmico	not available	2456382	LABORATÓRIOS DELTA, S.A.	PT
Dermestril 100 100 microgramas/24h Sistema transdérmico	not available	2456383	LABORATÓRIOS DELTA, S.A.	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 100 100 microgramas/24h Sistema transdérmico	not available	2456184	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL –Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0302/001	029001056	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 50 microgrammi/24 ore, cerotto transdermico	UK/H/0302/002	029001070	ROTTAPHARM S.P.A.	IT
DERMESTRIL –Septem 75 microgrammi/24 ore, cerotto transdermico	UK/H/0302/003	029001094	ROTTAPHARM S.P.A.	IT
DERMESTRIL-Septem 25 microgramas/24 horas sistema transdérmico	UK/H/0302/001	2970481	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL-Septem 25 microgramas/24 horas sistema transdérmico	UK/H/0302/001	2970382	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL-Septem 50 microgramas/24 horas sistema transdérmico	UK/H/0302/002	2970689	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL-Septem 75 microgramas/24 horas sistema transdérmico	UK/H/0302/003	2970887	LABORATÓRIOS DELTA, S.A.	PT
DERMESTRIL-Septem 75 microgramas/24 horas sistema transdérmico	UK/H/0302/003	2970788	LABORATÓRIOS DELTA, S.A.	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	not available	56/505/95-B/C	MEDA PHARMA S.R.O.	CZ
DERMESTRIL 25	not available	56/505/95-A/C	MEDA PHARMA S.R.O.	CZ
DERMESTRIL-Septem 50	not available	56/836/99-C	MEDA PHARMA S.R.O.	CZ
DERMESTRIL-Septem 25	not available	56/835/99-C	MEDA PHARMA S.R.O.	CZ
EPIESTROL –Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0303/001	029000054	ROTTAPHARM S.P.A.	IT
EPIESTROL –Septem 50 microgrammi/24 ore, cerotto transdermico	UK/H/0303/002	029000078	ROTTAPHARM S.P.A.	IT
EPIESTROL –Septem 75 microgrammi/24 ore, cerotto transdermico	UK/H/0303/003	029000092	ROTTAPHARM S.P.A.	IT
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

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 100 microgrammes/24 heures, dispositif transdermique	not available	34009 340 903 7 0	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 25 microgrammes/24 heures, dispositif transdermique	UK/H/0302/001	34009 352 393 9 6	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 25 microgrammes/24 heures, dispositif transdermique	UK/H/0302/001	34009 352 394 5 7	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 50 microgrammes/24 heures, dispositif transdermique	UK/H/0302/002	34009 352 391 6 7	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 50 microgrammes/24 heures, dispositif transdermique	UK/H/0302/002	34009 352 392 2 8	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 75 microgrammes/24 heures, dispositif transdermique	UK/H/0302/003	34009 352 395 1 8	ROTTAPHARM SAS	FR
DERMESTRIL SEPTEM 75 microgrammes/24 heures, dispositif transdermique	UK/H/0302/003	34009 352 396 8 6	ROTTAPHARM 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
EPIESTROL-Septem 75, 75 microgramas/24 horas sistema transdérmico	UK/H/0303/003	2971380	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
EPIESTROL-Septem 75, 75 microgramas/24 horas sistema transdérmico	UK/H/0303/003	2971489	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
EPIESTROL-Septem 50, 50 microgramas/24 horas sistema transdérmico	UK/H/0303/002	2971182	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
EPIESTROL-Septem 50, 50 microgramas/24 horas sistema transdérmico	UK/H/0303/002	2971281	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
EPIESTROL-Septem 25, 25 microgramas/24 horas sistema transdérmico	UK/H/0303/001	2970986	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
EPIESTROL-Septem 25, 25 microgramas/24 horas sistema transdérmico	UK/H/0303/001	2971083	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 25 25 microgramas/24h Sistema transdérmico	not available	2458180	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 25 25 microgramas/24h Sistema transdérmico	not available	2458289	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 25 25 microgramas/24h Sistema transdérmico	not available	2458388	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	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
Epiestrol 25 25 microgramas/24h Sistema transdérmico	not available	2458487	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 50 50 microgramas/24h Sistema transdérmico	not available	2458586	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 50 50 microgramas/24h Sistema transdérmico	not available	2458685	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 50 50 microgramas/24h Sistema transdérmico	not available	2458883	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 50 50 microgramas/24h Sistema transdérmico	not available	2458784	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 100 100 microgramas/24h Sistema transdérmico	not available	2458982	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 100 100 microgramas/24h Sistema transdérmico	not available	2459089	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 100 100 microgramas/24h Sistema transdérmico	not available	2459188	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Epiestrol 100 100 microgramas/24h Sistema transdérmico	not available	2459287	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	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
ARMONIL 100 microgrammi/24 ore cerotti transdermici	not available	032926038	MEDA PHARMA S.P.A.	IT
ARMONIL 25 microgrammi/24 ore cerotti transdermici	not available	032926014	MEDA PHARMA S.P.A.	IT
ARMONIL 50 microgrammi/24 ore cerotti transdermici	not available	032926026	MEDA PHARMA S.P.A.	IT
ARMONIL–Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0302/001	032926040	MEDA PHARMA S.P.A.	IT
ARMONIL–Septem 25 microgrammi/24 ore, cerotto transdermico	UK/H/0302/001	032926053	MEDA PHARMA S.P.A.	IT
ARMONIL–Septem 50 microgrammi/24 ore, cerotto transdermico	UK/H/0302/002	032926077	MEDA PHARMA S.P.A.	IT
ARMONIL–Septem 50 microgrammi/24 ore, cerotto transdermico	UK/H/0302/002	032926065	MEDA PHARMA S.P.A.	IT
ARMONIL–Septem 75 microgrammi/24 ore, cerotto transdermico	UK/H/0302/003	032926089	MEDA PHARMA S.P.A.	IT
DIVIGEL, 0,1% geel	not available	195998	ORION CORPORATION	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
Climara 50 microgrammes/24 heures dispositif transdermique	UK/H/0114/001	BE177091	BAYER SA NV	BE
Climara 50 Mikrogramm/24 Stunden transdermales Pflaster	UK/H/0114/001	BE177091	BAYER SA NV	BE
Climara 50 Mikrogramm/24 Stunden transdermales Pflaster	UK/H/0114/001	1996060370	BAYER SA NV	LU
Climara 50 mikrogramov/24 ur transdermalni obliž	not available	H/01/00400/001	BAYER PHARMA AG	SI
Climara 50 micrograme/24 ore plasture transdennic	not available	7541/2015/01-02	BAYER PHARMA AG	RO
Климара 50 микрограма/24 часа трансдермален пластир	not available	9900139	BAYER PHARMA AG	BG
Climara 50 mikrogrammi/24 tunnis transdermaalne plaaster	not available	160197	BAYER PHARMA AG	EE
Climara-50; 50 µg/dobę (3,8 mg), system transdermalny	not available	4583	BAYER PHARMA AG	PL
Progynova® TS 50 micrograms/24 hours Transdermal Patch	not available	MA513/03301	BAYER 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
Climara 50 microgrammes/24 heures dispositif transdermique	UK/H/0114/001	1996060370	BAYER SA NV	LU
Progynova® TS 100 micrograms/24 hours Transdermal Patch	UK/H/0114/002	PL 00010/0561	BAYER PLC	UK
Climara Forte 100 micrograms/24 hours Transdermal Patch	UK/H/0114/002	PA 1410/15/2	BAYER LTD	IE
Climara 100 microgrammi/24 h cerotti transdermici	UK/H/0114/002	030183040	BAYER SPA	IT
Climara 100 microgrammi/24 h cerotti transdermici	UK/H/0114/002	030183038	BAYER SPA	IT
Climara 50 Mikrogramm/24 h transdermales Pflaster	UK/H/0114/001	1-21511	BAYER AUSTRIA GMBH	AT
Progynova® TS 50 micrograms/24 hours Transdermal Patch	UK/H/0114/001	PL 00010/0560	BAYER PLC	UK
Climara 50 microgramas/ 24 horas sistemas transdérmicos	UK/H/0114/001	2671485	BAYER PORTUGAL LDA	PT
Climara 50 microgram/24 uur pleister voor transdermaal gebruik	UK/H/0114/001	BE177091	BAYER SA NV	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
Climara 50 micrograms/24 hours Transdermal Patch	UK/H/0114/001	PA 1410/15/1	BAYER LTD	IE
Climara 50 microgramas/ 24 horas sistemas transdérmicos	UK/H/0114/001	2671386	BAYER PORTUGAL LDA	PT
Climara 50 50 µg/24 hodin	not available	56/683/96-C	BAYER PHARMA AG	CZ
Climara 50 mikrogramu/24 stundās transdermāls plāksteris	not available	99-0016	BAYER PHARMA AG	LV
Climara 50 microgrammi/24 h cerotti transdermici	UK/H/0114/001	030183014	BAYER SPA	IT
Climara 50 microgrammi/24 h cerotti transdermici	UK/H/0114/001	030183026	BAYER SPA	IT
Gynokadin® Dosiergel	not available	31673.00.00	DR. KADE / BESINS PHARMA GMBH	DE
GINAIKOS 1,5 mg gel	not available	034727014	ABIOGEN PHARMA S.P.A.	IT
Estring 7,5 mikrograma na 24 ur vaginalni dostavni sistem	UK/H/5157/001	H/13/00589/001	PFIZER LUXEMBOURG SARL	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
ESTRING 2 mg, système de diffusion vaginal	UK/H/5157/001	34009 275 661 8 9	PFIZER HOLDING FRANCE (S.C.A.)	FR
Estring 7,5 mikrog/24 timmar vaginalinlägg	not available	11352	PFIZER OY	FI
Estring 7,5 microgramos cada 24 horas sistema de liberación vaginal	UK/H/5157/001	77867	PFIZER, S.L.	ES
Estring 7,5 microgrammi/24 ore, dispositivo vaginale	UK/H/5157/001	042840013	PFIZER ITALIA S.R.L.	IT
ESTRING® 2 mg Vaginalinsert Estradiol	not available	29279.00.00	PFIZER PHARMA PFE GMBH	DE
Oestring 7,5 mikrogram/24 timmar vaginalinlägg	not available	11783	PFIZER AB	SE
Estring 7,5 míkróg/24 klst.skeiðarinnlegg	not available	920143	PFIZER APS	IS
Estring, vaginalindlæg	not available	15084	PFIZER APS	DK
Estring 7,5 mikrogram/ 24 timer vaginalinnlegg	not available	94-2203	PFIZER AS	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
Estring 7,5 mikrog / 24 tuntia depotlääkevalmiste, emättimeen	not available	11352	PFIZER OY	FI
Estring 7.5 microgram/24 hours, vaginal delivery system	UK/H/5157/001	PL 00057/1424	PFIZER LIMITED	UK
DERMESTRIL ® - Septem 25 micrograms / 24 hrs Διαδερμικό έμπλαστρο	UK/H/0303/001	57757/18-9-2013	PROTON PHARMA S.A.	GR
DERMESTRIL ® - Septem 50 micrograms / 24 hrs Διαδερμικό έμπλαστρο	UK/H/0303/002	68915/18-9-2013	PROTON PHARMA S.A.	GR
DERMESTRIL ® - Septem 75 micrograms / 24 hrs Διαδερμικό έμπλαστρο	UK/H/0303/003	68916/18-9-2013	PROTON PHARMA S.A.	GR
DERMESTRIL	not available	57765/18-9-2013	PROTON PHARMA S.A.	GR
DERMESTRIL	not available	68913/18/9/2013	PROTON PHARMA S.A.	GR
DERMESTRIL	not available	68914/18/9/2013	PROTON PHARMA S.A.	GR
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	8708164	MERUS LABS LUXCO S.À R.L	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
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	8708107	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	3273786	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 25 25 microgramas/24 h Adesivo transdérmico	not available	3071289	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	8708172	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	8708123	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 25; 0,75 mg; 25 µg/24 h; system transdermalny	not available	7028	MERUS LABS LUXCO S.À R.L	PL
ESTRADERM MX 50; 1,5 mg; 50 µg/24 h; system transdermalny	not available	7029	MERUS LABS LUXCO S.À R.L	PL
ESTRADERM MX 100; 3 mg; 100 µg/24 h; system transdermalny	not available	7030	MERUS LABS LUXCO S.À R.L	PL
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	3273885	MERUS LABS LUXCO S.À R.L	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
ESTRADERM MX 25 mcg/die cerotto transdermico	not available	031773017	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 50 50 microgramas/24 h Adesivo transdérmico	not available	3152683	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 50 mcg/die cerotto transdermico	not available	031773029	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	8708180	MERUS LABS LUXCO S.À R.L	PT
ESTRADERM MX 100 mcg/die cerotto transdermico	not available	031773031	MERUS LABS LUXCO S.À R.L	IT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	8708149	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	3273984	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX 100 100 microgramas/24 h Adesivo transdérmico	not available	3152782	MERUS LABS LUXCO S.À R.L	PT
Estraderm MX® 25	not available	PL 44374/0003	MERUS LABS LUXCO S.À R.L	UK

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
Estraderm MX® 50	not available	PL 44374/0004	MERUS LABS LUXCO S.À R.L	UK
Estraderm MX ®75	not available	PL 44374/0005	MERUS LABS LUXCO S.À R.L	UK
Estraderm MX ®100	not available	PL 44374/0006	MERUS LABS LUXCO S.À R.L	UK
ESTRADERM® MATRIX 25 microgramos/24 horas parches transdérmicos	not available	59.153	MERUS LABS LUXCO S.À R.L	ES
ESTRADERM® MATRIX 50 microgramos/24 horas parches transdérmicos	not available	59.154	MERUS LABS LUXCO S.À R.L	ES
ESTRADERM® MATRIX 100 microgramos/24 horas parches transdérmicos	not available	59.155	MERUS LABS LUXCO S.À R.L	ES
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 665 0 6	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 664 4 5	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 1 mg, comprimé pelliculé	not available	34009 346 666 7 4	MERUS LABS LUXCO S.À R.L	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
PROVAMES 2 mg, comprimé pelliculé	not available	34009 354 794 0 2	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 2 mg, comprimé pelliculé	not available	34009 354 793 4 1	MERUS LABS LUXCO S.À R.L	FR
PROVAMES 2 mg, comprimé pelliculé	not available	34009 337 341 1 4	MERUS LABS LUXCO S.À R.L	FR