30 November 2017
EMA/811808/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: estradiol / norethisterone

Procedure no.: PSUSA/00001278/201703

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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
Activelle 1 mg + 0,5 mg, tabletki powlekane	not available	4512	NOVO NORDISK A/S	PL
Activelle 1 mg/0.5 mg film-coated tablets	not available	MA 104/00501	NOVO NORDISK A/S	MT
Activelle 1 mg/0,5 mg, Filmtabletten	SE/H/0150/001	BE196515	NOVO NORDISK PHARMA SA	BE
Activelle 1 mg/0,5 mg, Filmtabletten	SE/H/0150/001	2009020162	NOVO NORDISK PHARMA SA	LU
Activelle filmtabletta	SE/H/0150/001	OGYI-T-7031/02	NOVO NORDISK A/S	HU
Активел 1 mg/0,5 mg филмирани таблетки	SE/H/0150/001	II-26801	NOVO NORDISK A/S	BG
Activelle 1 mg/0,5 mg, comprimés pelliculés	SE/H/0150/001	2009020162	NOVO NORDISK PHARMA SA	LU
Activelle 1 mg/0,5 mg filmom obložene tablete	HR-H-673855616	UP/I-530-09/14-01/257	NOVO NORDISK A/S	HR
Activelle 1 mg/0,5 mg filmdragerade tabletter	SE/H/0150/001	13621	NOVO NORDISK A/S	FI
Kliovance 1 mg/0.5 mg film-coated tablets	SE/H/0150/001	PL 03132/0125	NOVO NORDISK LIMITED	UK
Activelle 1 mg/0,5 mg comprimidos recubiertos con película	SE/H/0150/001	62.465	ISDIN, S.A.	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
Activelle 1 mg/0.5 mg film-coated tablets	SE/H/0150/001	PA 218/52/1	NOVO NORDISK A/S	IE
Activelle 1 mg/0,5 mg Filmtabletten	SE/H/0150/001	1-22699	NOVO NORDISK PHARMA GMBH	AT
Activelle 1 mg/0,5 mg filmomhulde tabletten	SE/H/0150/001	BE196515	NOVO NORDISK PHARMA SA	BE
Activelle 1 mg/0,5 mg, comprimés pelliculés	SE/H/0150/001	BE196515	NOVO NORDISK PHARMA SA	BE
Activelle 1 mg/0,5 mg potahované tablety	SE/H/0150/001	56/827/99-C	NOVO NORDISK A/S	CZ
Activelle 1 mg/0,5 mg Filmtabletten Estradiol / Norethisteronacetat	SE/H/0150/001	43555.00.00	NOVO NORDISK PHARMA GMBH	DE
Activelle 1 mg/0,5 mg plėvele dengtos tabletės	SE/H/0150/001	LT/1/05/0298/002	NOVO NORDISK A/S	LT
Activelle 1 mg/0,5mg compresse rivestite con film	SE/H/0150/001	034117022/M	NOVO NORDISK SPA	IT
Activelle, 1 mg/0,5 mg õhukese polümeerikattega tabletid	SE/H/0150/001	291799	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
Activelle 1 mg/0,5 mg kalvopäällysteiset tabletit	SE/H/0150/001	13621	NOVO NORDISK A/S	FI
ACTIVELLE, comprimé pelliculé	SE/H/0150/001	NL 23753	NOVO NORDISK	FR
Activelle filmtabletta	SE/H/0150/001	OGYI-T-7031/01	NOVO NORDISK A/S	HU
Activelle 1 mg/0,5 mg plėvele dengtos tabletės	SE/H/0150/001	LT/1/05/0298/001	NOVO NORDISK A/S	LT
Activelle 1 mg/0,5 mg apvalkotās tabletes	SE/H/0150/001	05-0410	NOVO NORDISK A/S	LV
Activelle®, filmomhulde tabletten	SE/H/0150/001	RVG 22819	NOVO NORDISK B.V.	NL
Activelle 1 mg/0,5 mg filmom obalenè tablety	SE/H/0150/001	56/0215/05-S	NOVO NORDISK A/S	SK
Activelle	SE/H/0150/001	30095	NOVO NORDISK A/S	DK
Activelle 1 mg/0,5 mg filmdragerade tabletter	SE/H/0150/001	14007	NOVO NORDISK A/S	SE
Activelle 1 mg/0,5 mg filmsko obložene tablete	SE/H/0150/001	H/99/00114/001	NOVO NORDISK A/S	SI
Activelle 1 mg/0,5 mg filmsko obložene tablete	SE/H/0150/001	H/99/00114/002	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
Activelle 1 mg/0,5 mg tablett, filmdrasjert	SE/H/0150/001	05/3235	NOVO NORDISK A/S	NO
Activelle	SE/H/0150/001	980218	NOVO NORDISK A/S	IS
Activelle 1 mg/0,5mg compresse rivestite con film	SE/H/0150/001	034117010/M	NOVO NORDISK SPA	IT
Activelle 1mg/0,5mg comprimido revestido	SE/H/0150/001	2819282	ISDIN LDA	PT
Activelle 1mg/0,5mg comprimido revestido	SE/H/0150/001	2819381	ISDIN LDA	PT
Estramon conti® 40/130 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/002	70879.00.00	HEXAL AG	DE
Estramon conti® 30/95 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/001	70878.00.00	HEXAL AG	DE
Kliogest 2 mg/1 mg, Filmtabletten	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg, Filmtabletten	not available	0642/03/12/7922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg filmdragerade tabletter	not available	10575	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
Kliovance 1 mg/0.5 mg film-coated tablets	SE/H/0150/001	PL 03132/0125	NOVO NORDISK LIMITED	UK
Kliogest 2 mg/1 mg film-coated tablets	not available	PA 218/22/1	NOVO NORDISK A/S	IE
Kliogest 2mg/1mg Filmtabletten	not available	1-19017	NOVO NORDISK PHARMA GMBH	AT
Kliogest 2 mg/1 mg, filmomhulde tabletten	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg comprimés pelliculés	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2mg/1mg potahované tablety	DK/H/0102/001	56/302/91-C	NOVO NORDISK A/S	CZ
Kliogest N 2 mg/1 mg Filmtabletten	not available	26118.00.00	NOVO NORDISK PHARMA GMBH	DE
Kliogest 2 mg/1 mg plėvele dengtos tabletės	DK/H/0102/001	LT/1/05/0306/002	NOVO NORDISK A/S	LT
Kliogest filmtabletta	DK/H/0102/001	OGYI-T-1877/02	NOVO NORDISK A/S	HU
Kliogest, 2 mg/1 mg õhukese polümeerikattega tabletid	DK/H/0102/001	204998	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
Kliogest kalvopäällysteiset tabletit	not available	10575	NOVO NORDISK A/S	FI
KLIOGEST, comprimé pelliculé	not available	NL 15139	NOVO NORDISK	FR
Kliogest filmtabletta	DK/H/0102/001	OGYI-T-1877/01	NOVO NORDISK A/S	HU
Kliogest 2 mg/1 mg plėvele dengtos tabletės	DK/H/0102/001	LT/1/05/0306/001	NOVO NORDISK A/S	LT
Kliogest 2 mg/1 mg apvalkotās tabletes	DK/H/0102/001	05-0411	NOVO NORDISK A/S	LV
Kliogest, filmomhulde tabletten	not available	RVG 14942	NOVO NORDISK B.V.	NL
Kliogest, 2 mg + 1 mg, tabletki powlekane	not available	R/3297	NOVO NORDISK A/S	PL
Kliogest	DK/H/0102/001	11768	NOVO NORDISK A/S	DK
Kliogest 2 mg/1 mg, comprimés pelliculés	not available	0642/03/12/7922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg filmsko obložene tablete	DK/H/0102/001	H/92/00845/001	NOVO NORDISK A/S	SI
Kliogest 2 mg/1 mg filmsko obložene tablete	DK/H/0102/001	H/92/00845/002	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
Kliogest filmuhúðaðar töflur	DK/H/0102/001	843355	NOVO NORDISK A/S	IS
Kliogest®, 2 mg + 1 mg, Comprimido revestido por película	not available	2280782	ISDIN LDA	PT
Trisequens, Filmtabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, Filmtabletten	not available	0642/08039743	NOVO NORDISK PHARMA SA	LU
Trisequens filmsko obložene tablete	not available	H/97/01568/001	NOVO NORDISK A/S	SI
Trisequens, δισκία επικαλυμμένα με λεπτό υμένιο	not available	13501	NOVO NORDISK HELLAS LTD.	CY
Trisekvens filmdragerade tabletter	not available	8424	NOVO NORDISK A/S	FI
Trisequens® comprimidos recubiertos	not available	60.706	ISDIN, S.A.	ES
Trisequens film-coated tablets	not available	PL 03132/0122	NOVO NORDISK LIMITED	UK
Trisequens film-coated tablets	not available	PA 218/8/1	NOVO NORDISK A/S	IE
Trisequens potahované tablety	not available	56/307/91-C	NOVO NORDISK A/S	CZ

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
Трисеквенс филмирана таблетки	not available	20010005	NOVO NORDISK A/S	BG
Trisequens, comprimés pelliculés	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, filmomhulde tabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens - Filmtabletten	not available	16.893	NOVO NORDISK PHARMA GMBH	AT
Trisequens Filmtabletten Estradiol / Norethisteronacetat	not available	6529829.00.00	NOVO NORDISK PHARMA GMBH	DE
Trisequens, õhukese polümeerikattega tabletid	not available	205098	NOVO NORDISK A/S	EE
Trisekvens kalvopäällysteiset tabletit	not available	8424	NOVO NORDISK A/S	FI
TRISEQUENS, comprimé pelliculé	not available	NL 12375	NOVO NORDISK	FR
Trisequens filmom obložene tablete	not available	HR-H-919057630	NOVO NORDISK HRVATSKA D.O.O.	HR
Trisequens filmtabletta	not available	OGYI-T-5851/01	NOVO NORDISK A/S	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
Trisequens, δισκία επικαλυμμένα με λεπτό υμένιο	not available	46163/06-07-2015	NOVO NORDISK HELLAS LTD.	GR
Trisequens plėvele dengtos tabletės	not available	LT/1/94/0158/001	NOVO NORDISK A/S	LT
Trisequens apvalkotās tabletes	not available	98-0705	NOVO NORDISK A/S	LV
Trisequens, filmomhulde tablet	not available	RVG 09812	NOVO NORDISK B.V.	NL
Trisequens, 2 mg (niebieskie), 2 mg + 1 mg (białe), 1 mg (czerwone), tabletki powlekane	not available	R/3299	NOVO NORDISK A/S	PL
Trisekvens, filmovertrukne tabletter	not available	6699	NOVO NORDISK A/S	DK
Trisequens, comprimés pelliculés	not available	0642/08039743	NOVO NORDISK PHARMA SA	LU
Trisekvens tablett, filmdrasjert	not available	6323	NOVO NORDISK A/S	NO
Trisekvens filmuhúðaðar töflur	not available	751874	NOVO NORDISK 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
Trisequens associação comprimidos revestidos por película	not available	2480986	ISDIN LDA	PT
Novofem Filmtabletten	DE/H/0304/001	BE231682	NOVO NORDISK PHARMA SA	BE
Novofem, Filmtabletten	DE/H/0304/001	0642/07039243	NOVO NORDISK PHARMA SA	LU
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/001	NOVO NORDISK A/S	SI
Novofem filmom obložene tablete	DE/H/0304/001	HR-H-818589847	NOVO NORDISK A/S	HR
Novofem filmdragerade tabletter	DE/H/0304/001	16601	NOVO NORDISK A/S	FI
Duofemme comprimidos recubiertos con película	DE/H/1/0304/01	64.718	ISDIN, S.A.	ES
Novofem film-coated tablet	DE/H/0304/001	PL 03132/0141	NOVO NORDISK LIMITED	UK
Novofem film-coated tablets	DE/H/0304/001	PA 218/053/001	NOVO NORDISK A/S	IE
Novofem potahované tablety	DE/H/0304/001	56/005/03-C	NOVO NORDISK A/S	CZ
Novofem, comprimés pelliculés	DE/H/0304/001	BE231682	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
Novofem, filmomhulde tabletten	DE/H/0304/001	BE231682	NOVO NORDISK PHARMA SA	BE
Novofem Filmtabletten Estradiol / Norethisteronacetat	DE/H/0304/001	47567.00.00	NOVO NORDISK PHARMA GMBH	DE
Novofem, õhukese polümeerikattega tabletid	DE/H/0304/001	407903	NOVO NORDISK A/S	EE
Novofem kalvopäällysteiset tabletit	DE/H/0304/001	16601	NOVO NORDISK A/S	FI
NOVOFEMME, comprimé pelliculé	DE/H/0304/001	NL 26893	NOVO NORDISK	FR
Novofem filmtabletta	DE/H/0304/001	OGYI-T-10 607/01	NOVO NORDISK A/S	HU
Novofem plėvele dengtos tabletės	DE/H/0304/001	LT/1/05/0316/002	NOVO NORDISK A/S	LT
Novofem plėvele dengtos tabletės	DE/H/0304/001	LT/1/05/0316/001	NOVO NORDISK A/S	LT
Novofem apvalkotās tabletes	DE/H/0304/001	05-0412	NOVO NORDISK A/S	LV
Novofem, filmomhulde tabletten	DE/H/0304/001	RVG 26864	NOVO NORDISK 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
Novofem, tabletki powlekane	DE/H/0304/001	11888	NOVO NORDISK A/S	PL
Novofem filmom obalené tablety	DE/H/0304/001	56/0291/05-S	NOVO NORDISK A/S	SK
Novofem filmovertrukne tabletter	DE/H/0304/001	32745	NOVO NORDISK A/S	DK
Novofem filmdragerade tabletter	DE/H/0304/001	17439	NOVO NORDISK A/S	SE
Novofem, comprimés pelliculés	DE/H/0304/001	0642/07039243	NOVO NORDISK PHARMA SA	LU
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/002	NOVO NORDISK A/S	SI
Novofem tablett, filmdrasjert	DE/H/0304/001	01-5767	NOVO NORDISK A/S	NO
Novofem filmuhúðaðar töflur	DE/H/0304/001	IS/1/01/040/01	NOVO NORDISK A/S	IS
Novofem comprimidos revestidos por película	DE/H/0304/001	3804986	ISDIN LDA	PT
Novofem comprimidos revestidos por película	DE/H/0304/001	3805082	ISDIN LDA	PT
Activelle minor 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	BE329682	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
Activelle minor 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	2009120001	NOVO NORDISK PHARMA SA	LU
Eviana 0,5 mg/0,1 mg Filmtabletta	SE/H/0150/002	OGYI-T-22398/02	NOVO NORDISK A/S	HU
Noviana 0,5 mg/0,1 mg filmdragerade tabletter	SE/H/0150/002	24071	NOVO NORDISK A/S	FI
Eviana 0,5 mg/0,1 mg comprimidos recubiertos	SE/H/0150/002	70.576	ISDIN, S.A.	ES
Activelle 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	1-27819	NOVO NORDISK PHARMA GMBH	AT
Activelle minor 0,5 mg/0,1 mg filmomhulde tabletten	SE/H/0150/002	BE329682	NOVO NORDISK PHARMA SA	BE
Activelle minor 0,5 mg/0,1 mg, comprimés pelliculés	SE/H/0150/002	BE329682	NOVO NORDISK PHARMA SA	BE
Activelle 0,5 mg/0,1 mg compresse rivestite con film	SE/H/0150/002	034117046/M	NOVO NORDISK A/S	IT
Eviana, 0,5 mg/0,1 mg õhukese polümeerikattega tabletid	SE/H/0150/002	606708	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
Noviana 0,5 mg/0,1 mg kalvopäällysteiset tabletit	SE/H/0150/002	24071	NOVO NORDISK A/S	FI
Activelle 0,5 mg/0,1 mg compresse rivestite con film	SE/H/0150/002	034117034/M	NOVO NORDISK A/S	IT
Eviana filmomhulde tabletten	SE/H/0150/002	RVG 100983	NOVO NORDISK B.V.	NL
Activelle low	SE/H/0150/002	42023	NOVO NORDISK A/S	DK
Eviana 0,5 mg/0,1 mg filmdragerade tabletter	SE/H/0150/002	23381	NOVO NORDISK A/S	SE
Activelle minor 0,5 mg/0,1 mg, comprimés pelliculés	SE/H/0150/002	2009120001	NOVO NORDISK PHARMA SA	LU
Eviana 0,5 mg/0,1 mg tablett, filmdrasjert	SE/H/0150/002	07-5213	NOVO NORDISK A/S	NO
Activelle low	SE/H/0150/002	IS/1/08/002/01	NOVO NORDISK A/S	IS
Activelle 0,5 mg/0,1 mg Comprimido revestido por película	SE/H/0150/002	5192703	ISDIN LDA	PT
Activelle 0,5 mg/0,1 mg Comprimido revestido por película	SE/H/0150/002	5192679	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
Eviana 0,5 mg/0,1 mg filmtabletta	SE/H/0150/002	OGYI-T-22398/01	NOVO NORDISK A/S	HU
ESTALIS 50 Mikrogramm /250 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
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
ESTALIS 0,6 mg; 50 µg/24h + 2,7 mg; 140 µg/24h, system transdermalny	not available	8166	NOVARTIS POLAND SP. Z O. O.	PL
Estalis; 0,5mg; 50 µg/24h + 4,8 mg; 250 µg/24h, system transdermalny	not available	8165	NOVARTIS POLAND SP. Z O. O.	PL
MERIGEST® 2 MG/0,7 MG FILMTABLETTEN	UK/H/0143/001	39491.00.00	NOVARTIS PHARMA GMBH	DE
Climesse® 2mg/0.7 film-coated tablets	UK/H/0143/001	PL 00101/0396	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
Estalis 50 microgramas/250 microgramas/24 horas, adesivo transdérmico	SE/H/0148/001	2777084	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estalis 50 microgramas/250 microgramas/24 horas, adesivo transdérmico	SE/H/0148/001	2777183	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estalis 50 microgramas/250 microgramas/24 horas, adesivo transdérmico	SE/H/0148/001	4800587	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
ESTALIS SEQUI cerotti transdermici	SE/H/0149/002	034209039	NOVARTIS FARMA S.P.A.	IT
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

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 50 μικρογραμμάρια/250 μικρογραμμάρια/24 ώρες, διαδερμικό έμπλαστρο	SE/H/0148/001	250630101	NOVARTIS (HELLAS) S.A.C.I.	GR
Estalis 50 μικρογραμμάρια/250 μικρογραμμάρια/24 ώρες, διαδερμικό έμπλαστρο	SE/H/0148/001	250630102	NOVARTIS (HELLAS) S.A.C.I.	GR
Estalis 50 mikrog/250 mikrog per 24 tuntia depotlaastari	SE/H/0148/001	13620	NOVARTIS FINLAND OY	FI
Estalis 50 mikrog/250 mikrog/24 timmar, depotplåster	SE/H/0148/001	13620	NOVARTIS FINLAND OY	FI
ESTALIS 50 microgramos/250 microgramos/24 horas, parche transdérmico	SE/H/0148/001	62.463	NOVARTIS FARMACÉUTICA S.A.	ES
ESTALIS SEQUIDOT parche transdérmico	SE/H/0149/002	69.746	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
Sequidot® 50/250 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/0149/002	66641.00.00	NOVARTIS PHARMA GMBH	DE
CLIMAGEST 2mg film-coated tablet	UK/H/0127/001	PL 00101/0366	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Estalis 50/250 - transdermale Pflaster	SE/H/0148/001	1-22840	NOVARTIS PHARMA GMBH	AT
Estalis 50/250 mikrogram/ 24 timmar, depotplåster	SE/H/0148/001	13868	NOVARTIS SVERIGE AB	SE
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
Estalis 50 mikrog /250 mikrog per 24 timer “Novartis”	not available	98-3106	NOVARTIS NORGE AS	NO
Sequidot depotplaster	not available	04-3137	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
Estalis 50 microgram/250 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
ESTALIS 50/250 dispositifs transdermiques	SE/H/0148/001	BE199245	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
CLIMAGEST 2mg film-coated tablet	UK/H/0127/001	PL 00101/0366	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
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
Kliane 1 mg/2 mg plėvele dengtos tabletės	not available	LT/1/2000/0892/001	BAYER PHARMA AG	LT
Kliane 1 mg/2 mg apvalkotās tabletes	not available	00-1227	BAYER PHARMA AG	LV
KLIANE Potahované tablety	not available	56/177/00-C	BAYER AG	CZ
Nuvelle™ Continuous	not available	PL 00010/0552	BAYER PLC	UK
Evo-Sequi	not available	18601	JANSSEN-CILAG A/S	DK
Evo-Conti	not available	18600	JANSSEN-CILAG A/S	DK
EVOREL CONTI	not available	PL 00242/0319	JANSSEN-CILAG LIMITED	UK
EVOREL SEQUI	not available	PL 00242/0320	JANSSEN-CILAG LIMITED	UK
Evorel Conti forðaplástur.	not available	960226	JANSSEN-CILAG AB	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
SYSTEN SEQUI: Systen 50; 3,2 mg + Systen Conti; 3,2 mg +11,2 mg, system transdermalny	not available	4448	JANSSEN-CILAG INTERNATIONAL NV	PL
Evorel Sequi forðaplástur.	not available	960227	JANSSEN-CILAG AB	IS
SYSTEN CONTI; 3,2 mg + 11,2 mg, system transdennalny	not available	4447	JANSSEN-CILAG INTERNATIONAL NV	PL
Evorel Conti 50/170 micrograms per 24 hours Transdermal patch	not available	PA 748/008/001	JANSSEN-CILAG LIMITED	IE
EVOREL SEQUI depotlaastari	not available	12676	JANSSEN-CILAG OY	FI
EVOREL CONTI depotlaastari	not available	12674	JANSSEN-CILAG OY	FI
MERIGEST® 2 MG/0,7 MG FILMTABLETTEN	UK/H/0143/001	39491.00.00	NOVARTIS PHARMA GMBH	DE
Climesse® 2mg/0.7 film-coated tablets	UK/H/0143/001	PL 00101/0396	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Kliogest 2 mg/1 mg, Filmtabletten	not available	BE165462	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
Kliogest 2 mg/1 mg, Filmtabletten	not available	0642/03/12/7922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg filmdragerade tabletter	not available	10575	NOVO NORDISK A/S	FI
Kliofem® film-coated tablets	not available	PL 3132/0080	NOVO NORDISK LIMITED	UK
Kliogest 2 mg/1 mg film-coated tablets	not available	PA 218/22/1	NOVO NORDISK A/S	IE
Kliogest 2mg/1mg Filmtabletten	not available	1-19017	NOVO NORDISK PHARMA GMBH	AT
Kliogest 2 mg/1 mg, filmomhulde tabletten	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg comprimés pelliculés	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest N 2 mg/1 mg Filmtabletten	not available	26118.00.00	NOVO NORDISK PHARMA GMBH	DE
Kliogest kalvopäällysteiset tabletit	not available	10575	NOVO NORDISK A/S	FI
KLIOGEST, comprimé pelliculé	not available	NL 15139	NOVO NORDISK	FR
Kliogest, filmomhulde tabletten	not available	RVG 14942	NOVO NORDISK 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
Kliogest, 2 mg + 1 mg, tabletki powlekane	not available	R/3297	NOVO NORDISK A/S	PL
Kliogest 2 mg/1 mg, comprimés pelliculés	not available	0642/03/12/7922	NOVO NORDISK PHARMA SA	LU
Kliogest®, 2 mg + 1 mg, Comprimido revestido por película	not available	2280782	ISDIN LDA	PT