1 December 2022
EMA/934662/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: estradiol / norethisterone

Procedure no.: PSUSA/00001278/202203

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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 0,5 mg/0,1 mg compresse rivestite con film	SE/H/0150/002	034117046	NOVO NORDISK A/S	IT
Activelle 0,5 mg/0,1 mg compresse rivestite con film	SE/H/0150/002	034117046	NOVO NORDISK A/S	IT
Activelle 0,5 mg/0,1 mg compresse rivestite con film.	SE/H/0150/002	034117034	NOVO NORDISK A/S	IT
Activelle 0,5 mg/0,1 mg compresse rivestite con film.	SE/H/0150/002	034117034	NOVO NORDISK A/S	IT
Activelle 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	1-27819	NOVO NORDISK PHARMA GMBH	AT
Activelle 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	1-27819	NOVO NORDISK PHARMA GMBH	AT
Activelle 1 mg + 0,5 mg, tabletki powlekane	not available	4512	NOVO NORDISK A/S	PL
Activelle 1 mg + 0,5 mg, tabletki powlekane	not available	4512	NOVO NORDISK A/S	PL
Activelle 1 mg/0,5 mg apvalkotās tabletes	SE/H/0150/001	05-0410	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
Activelle 1 mg/0,5 mg filmdragerade tabletter	SE/H/0150/001	13621	NOVO NORDISK A/S	FI
Activelle 1 mg/0,5 mg filmdragerade tabletter	SE/H/0150/001	14007	NOVO NORDISK A/S	SE
Activelle 1 mg/0,5 mg filmom obalenè tablety	SE/H/0150/001	56/0215/05-S	NOVO NORDISK A/S	SK
Activelle 1 mg/0,5 mg filmom obložene tablete	SE/H/0150/001	HR-H-673855616	NOVO NORDISK A/S	HR
Activelle 1 mg/0,5 mg filmomhulde tabletten	SE/H/0150/001	BE196515	NOVO NORDISK PHARMA SA	BE
Activelle 1 mg/0,5 mg filmovertrukne tabletter	SE/H/0150/001	30095	NOVO NORDISK A/S	DK
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
Activelle 1 mg/0,5 mg Filmtabletten	SE/H/0150/001	1-22699	NOVO NORDISK 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
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 filmuhúðaðar töflur	SE/H/0150/001	980218	NOVO NORDISK A/S	IS
Activelle 1 mg/0,5 mg kalvopäällysteiset tabletit	SE/H/0150/001	13621	NOVO NORDISK A/S	FI
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/0,5 mg plėvele dengtos tabletės	SE/H/0150/001	LT/1/05/0298/001	NOVO NORDISK A/S	LT
Activelle 1 mg/0,5 mg potahované tablety	SE/H/0150/001	56/827/99-C	NOVO NORDISK A/S	CZ
Activelle 1 mg/0,5 mg tablett, filmdrasjert	SE/H/0150/001	05-3235	NOVO NORDISK A/S	NO
Activelle 1 mg/0,5 mg, comprimés pelliculés	SE/H/0150/001	BE196515	NOVO NORDISK PHARMA SA	BE
Activelle 1 mg/0,5 mg, comprimés pelliculés	SE/H/0150/001	2009020162	NOVO NORDISK PHARMA SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 1 mg/0,5mg compresse rivestite con film	SE/H/0150/001	034117022	NOVO NORDISK SPA	IT
Activelle 1 mg/0,5mg compresse rivestite con film.	SE/H/0150/001	034117010	NOVO NORDISK SPA	IT
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 film-coated tablets	not available	MA 104/00501	NOVO NORDISK A/S	MT
Activelle 1mg/0,5mg comprimido revestido por película	SE/H/0150/001	2819282	ISDIN LDA	PT
Activelle 1mg/0,5mg comprimido revestido por película	SE/H/0150/001	2819381	ISDIN LDA	PT
Activelle filmtabletta	SE/H/0150/001	OGYI-T-7031/02	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
Activelle filmtabletta	SE/H/0150/001	OGYI-T-7031/01	NOVO NORDISK A/S	HU
Activelle low	SE/H/0150/002	IS/1/08/002/01	NOVO NORDISK A/S	IS
Activelle low	SE/H/0150/002	IS/1/08/002/01	NOVO NORDISK A/S	IS
Activelle low 0,5 mg/0,1 mg filmovertrukne tabletter	SE/H/0150/002	42023	NOVO NORDISK A/S	DK
Activelle low 0,5 mg/0,1 mg filmovertrukne tabletter	SE/H/0150/002	42023	NOVO NORDISK A/S	DK
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 filmomhulde tabletten	SE/H/0150/002	BE329682	NOVO NORDISK PHARMA SA	BE
Activelle minor 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	BE329682	NOVO NORDISK PHARMA SA	BE
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
Activelle minor 0,5 mg/0,1 mg Filmtabletten	SE/H/0150/002	2009120001	NOVO NORDISK PHARMA SA	LU
Activelle minor 0,5 mg/0,1 mg, comprimés pelliculés	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 minor 0,5 mg/0,1 mg, comprimés pelliculés	SE/H/0150/002	2009120001	NOVO NORDISK PHARMA SA	LU
Activelle minor 0,5 mg/0,1 mg, comprimés pelliculés	SE/H/0150/002	2009120001	NOVO NORDISK PHARMA SA	LU
Activelle 0,5 mg/0,1 mg Comprimidos revestidos por película	SE/H/0150/002	5192703	ISDIN LDA	PT
Activelle 0,5 mg/0,1 mg Comprimidos revestidos por película	SE/H/0150/002	5192679	ISDIN LDA	PT
Activelle 0,5 mg/0,1 mg Comprimidos revestidos por película	SE/H/0150/002	5192703	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
Activelle 0,5 mg/0,1 mg Comprimidos revestidos por película	SE/H/0150/002	5192679	ISDIN LDA	PT
Activelle 1 mg/0,5 mg Comprimidos recubiertos con película	SE/H/0150/001	62.465	ISDIN, S.A.	ES
Activelle, 1 mg/0,5 mg õhukese polümeerikattega tabletid	SE/H/0150/001	291799	NOVO NORDISK A/S	EE
ACTIVELLE, comprimé pelliculé	SE/H/0150/001	NL23753	NOVO NORDISK	FR
Activelle, filmomhulde tabletten	SE/H/0150/001	RVG 22819	NOVO NORDISK B.V.	NL
Duofemme comprimidos recubiertos con película	DE/H/1/0304/01	64.718	ISDIN, S.A.	ES
Duofemme comprimidos recubiertos con película	DE/H/1/0304/01	64.718	ISDIN, S.A.	ES
Elleste Duet Conti Tablets 2 mg/1 mg film-coated tablets	not available	PL 46302/0166	MYLAN PRODUCTS LIMITED	XI
Elleste Duet Conti Tablets 2 mg/1 mg film-coated tablets	not available	PL 46302/0166	MYLAN PRODUCTS LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Estalis 50 microgram/250 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
Estalis 50 microgram/250 microgram/24 uur, pleister voor transdermaal gebruik	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
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 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

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	4800587	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Estalis 50 microgrammes/250 microgrammes/24 heures, dispositif transdermique	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
Estalis 50 microgrammes/250 microgrammes/24 heures, dispositif transdermique	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
Estalis 50 mikrog /250 mikrog per 24 timer "Novartis"	not available	98-3106	NOVARTIS NORGE AS	NO
Estalis 50 mikrog /250 mikrog per 24 timer "Novartis"	not available	98-3106	NOVARTIS NORGE AS	NO
Estalis 50 mikrog/250 mikrog per 24 tuntia depotlaastari	SE/H/0148/001	13620	NOVARTIS FINLAND OY	FI
Estalis 50 mikrog/250 mikrog per 24 tuntia depotlaastari	SE/H/0148/001	13620	NOVARTIS FINLAND OY	FI
Estalis 50 mikrogram/250 mikrogram/24 timmar, depotplåster	SE/H/0148/001	13620	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
Estalis 50 mikrogram/250 mikrogram/24 timmar, depotplåster	SE/H/0148/001	13620	NOVARTIS FINLAND OY	FI
Estalis 50 mikrogram/250 mikrogram/24 timmar, depotplåster	SE/H/0148/001	13868	NOVARTIS SVERIGE AB	SE
Estalis 50 mikrogram/250 mikrogram/24 timmar, depotplåster	SE/H/0148/001	13868	NOVARTIS SVERIGE AB	SE
ESTALIS 50 Mikrogramm /250 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
ESTALIS 50 Mikrogramm /250 Mikrogramm/24 Stunden transdermales Pflaster	SE/H/0148/001	BE199245	NOVARTIS PHARMA N.V.	BE
Estalis 50/250 - transdermale Pflaster	SE/H/0148/001	1-22840	NOVARTIS PHARMA GMBH	AT
Estalis 50/250 - transdermale Pflaster	SE/H/0148/001	1-22840	NOVARTIS PHARMA GMBH	AT
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

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	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
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
ESTALIS SEQUIDOT parche transdérmico	SE/H/0149/002	69.746	NOVARTIS FARMACÉUTICA S.A.	ES
ESTALIS SEQUIDOT parche transdérmico	SE/H/0149/002	69.746	NOVARTIS FARMACÉUTICA S.A.	ES
ESTALIS SEQUIDOT, διαδερμικό έμπλαστρο	SE/H/0149/002	273100101	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
ESTALIS SEQUIDOT, διαδερμικό έμπλαστρο	SE/H/0149/002	273100102	NOVARTIS (HELLAS) S.A.C.I.	GR
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
ESTRAMON conti® 30/95 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/001	70878.00.00	HEXAL AG	DE
ESTRAMON conti® 30/95 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/001	70878.00.00	HEXAL AG	DE
Estramon conti® 40/130 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/002	70879.00.00	HEXAL AG	DE
Estramon conti® 40/130 Mikrogramm/24 h Transdermales Pflaster	DE/H/3299/002	70879.00.00	HEXAL AG	DE
Eviana 0,5 mg/0,1 mg comprimidos recubiertos	SE/H/0150/002	70.576	ISDIN, S.A.	ES
Eviana 0,5 mg/0,1 mg comprimidos recubiertos	SE/H/0150/002	70.576	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
Eviana 0,5 mg/0,1 mg filmdragerade tabletter	SE/H/0150/002	23381	NOVO NORDISK A/S	SE
Eviana 0,5 mg/0,1 mg filmdragerade tabletter	SE/H/0150/002	23381	NOVO NORDISK A/S	SE
Eviana 0,5 mg/0,1 mg filmomhulde tabletten	SE/H/0150/002	RVG 100983	NOVO NORDISK B.V.	NL
Eviana 0,5 mg/0,1 mg filmomhulde tabletten	SE/H/0150/002	RVG 100983	NOVO NORDISK B.V.	NL
Eviana 0,5 mg/0,1 mg tablett, filmdrasjert	SE/H/0150/002	07-5213	NOVO NORDISK A/S	NO
Eviana 0,5 mg/0,1 mg tablett, filmdrasjert	SE/H/0150/002	07-5213	NOVO NORDISK A/S	NO
Eviana, 0,5 mg/0,1 mg õhukese polümeerikattega tabletid	SE/H/0150/002	606708	NOVO NORDISK A/S	EE
Eviana, 0,5 mg/0,1 mg õhukese polümeerikattega tabletid	SE/H/0150/002	606708	NOVO NORDISK A/S	EE
Evo-Conti, depotplaster	not available	18600	THERAMEX IRELAND LIMITED	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
Evo-Conti, depotplaster	not available	18600	THERAMEX IRELAND LIMITED	DK
Evorel Conti 50/170 micrograms per 24 hours Transdermal Patch	not available	PA22668/009/001	THERAMEX IRELAND LIMITED	IE
Evorel Conti 50/170 micrograms per 24 hours Transdermal Patch	not available	PA22668/009/001	THERAMEX IRELAND LIMITED	IE
EVOREL CONTI depotlaastari	not available	12674	THERAMEX IRELAND LIMITED	FI
EVOREL CONTI depotlaastari	not available	12674	THERAMEX IRELAND LIMITED	FI
EVOREL SEQUI depotlaastari	not available	12676	THERAMEX IRELAND LIMITED	FI
EVOREL SEQUI depotlaastari	not available	12676	THERAMEX IRELAND LIMITED	FI
Evo-Sequi, depotplaster	not available	18601	THERAMEX IRELAND LIMITED	DK
Evo-Sequi, depotplaster	not available	18601	THERAMEX IRELAND LIMITED	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
Kliogest 2 mg/1 mg apvalkotās tabletes	DK/H/0102/001	05-0411	NOVO NORDISK A/S	LV
Kliogest 2 mg/1 mg apvalkotās tabletes	DK/H/0102/001	05-0411	NOVO NORDISK A/S	LV
Kliogest 2 mg/1 mg comprimés pelliculés	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg comprimés pelliculés	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg film-coated tablets	not available	PA 218/22/1	NOVO NORDISK A/S	IE
Kliogest 2 mg/1 mg film-coated tablets	not available	PA 218/22/1	NOVO NORDISK A/S	IE
Kliogest 2 mg/1 mg filmdragerade tabletter	not available	10575	NOVO NORDISK A/S	FI
Kliogest 2 mg/1 mg filmdragerade tabletter	not available	10575	NOVO NORDISK A/S	FI
Kliogest 2 mg/1 mg filmsko obložene tablete	DK/H/0102/001	H/92/00845/001	NOVO NORDISK A/S	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kliogest 2 mg/1 mg filmsko obložene tablete	DK/H/0102/001	H/92/00845/002	NOVO NORDISK A/S	SI
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
Kliogest 2 mg/1 mg filmtabletta	DK/H/0102/001	OGYI-T-1877/02	NOVO NORDISK A/S	HU
Kliogest 2 mg/1 mg filmtabletta	DK/H/0102/001	OGYI-T-1877/01	NOVO NORDISK A/S	HU
Kliogest 2 mg/1 mg filmtabletta	DK/H/0102/001	OGYI-T-1877/02	NOVO NORDISK A/S	HU
Kliogest 2 mg/1 mg 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/1/05/0306/002	NOVO NORDISK A/S	LT
Kliogest 2 mg/1 mg plėvele dengtos tabletės	DK/H/0102/001	LT/1/05/0306/001	NOVO NORDISK A/S	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
Kliogest 2 mg/1 mg plėvele dengtos tabletės	DK/H/0102/001	LT/1/05/0306/002	NOVO NORDISK A/S	LT
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 potahované tablety	DK/H/0102/001	56/302/91-C	NOVO NORDISK A/S	CZ
Kliogest 2 mg/1 mg potahované tablety	DK/H/0102/001	56/302/91-C	NOVO NORDISK A/S	CZ
Kliogest 2 mg/1 mg, comprimés pelliculés	not available	2003127922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg, comprimés pelliculés	not available	2003127922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg, filmomhulde tabletten	not available	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg, filmomhulde tabletten	not available	BE165462	NOVO NORDISK PHARMA SA	BE
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	BE165462	NOVO NORDISK PHARMA SA	BE
Kliogest 2 mg/1 mg, Filmtabletten	not available	2003127922	NOVO NORDISK PHARMA SA	LU
Kliogest 2 mg/1 mg, Filmtabletten	not available	2003127922	NOVO NORDISK PHARMA SA	LU
Kliogest 2mg/1mg Filmtabletten	not available	1-19017	NOVO NORDISK PHARMA GMBH	AT
Kliogest 2mg/1mg Filmtabletten	not available	1-19017	NOVO NORDISK PHARMA GMBH	AT
Kliogest filmovertrukne tabletter	DK/H/0102/001	11768	NOVO NORDISK A/S	DK
Kliogest filmovertrukne tabletter	DK/H/0102/001	11768	NOVO NORDISK A/S	DK
Kliogest filmuhúðaðar töflur	DK/H/0102/001	843355	NOVO NORDISK A/S	IS
Kliogest filmuhúðaðar töflur	DK/H/0102/001	843355	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
Kliogest kalvopäällysteiset tabletit	not available	10575	NOVO NORDISK A/S	FI
Kliogest kalvopäällysteiset tabletit	not available	10575	NOVO NORDISK A/S	FI
Kliogest N 2 mg/1 mg Filmtabletten	not available	26118.00.00	NOVO NORDISK PHARMA GMBH	DE
Kliogest N 2 mg/1 mg Filmtabletten	not available	26118.00.00	NOVO NORDISK PHARMA GMBH	DE
Kliogest, 2 mg + 1 mg, tabletki powlekane	not available	R/3297	NOVO NORDISK A/S	PL
Kliogest, 2 mg + 1 mg, tabletki powlekane	not available	R/3297	NOVO NORDISK A/S	PL
Kliogest, 2 mg/1 mg õhukese polümeerikattega tabletid	DK/H/0102/001	204998	NOVO NORDISK A/S	EE
Kliogest, 2 mg/1 mg õhukese polümeerikattega tabletid	DK/H/0102/001	204998	NOVO NORDISK A/S	EE
KLIOGEST, comprimé pelliculé	not available	NL 15139	NOVO NORDISK	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
KLIOGEST, comprimé pelliculé	not available	NL 15139	NOVO NORDISK	FR
Kliogest, filmomhulde tabletten	not available	RVG 14942	NOVO NORDISK B.V.	NL
Kliogest, filmomhulde tabletten	not available	RVG 14942	NOVO NORDISK B.V.	NL
Kliogest®, 2 mg + 1 mg, Comprimido revestido por película	not available	2280782	ISDIN LDA	PT
Kliogest®, 2 mg + 1 mg, Comprimido revestido por película	not available	2280782	ISDIN LDA	PT
Kliovance 1 mg/0.5 mg film-coated tablets	SE/H/0150/001	PL 03132/0125	NOVO NORDISK LIMITED	XI
Novofem apvalkotās tabletes	DE/H/0304/001	05-0412	NOVO NORDISK A/S	LV
Novofem apvalkotās tabletes	DE/H/0304/001	05-0412	NOVO NORDISK A/S	LV
Novofem comprimidos revestidos por película	DE/H/0304/001	3804986	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
Novofem comprimidos revestidos por película	DE/H/0304/001	3805082	ISDIN LDA	PT
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
Novofem film-coated tablet	DE/H/0304/001	PL 03132/0141	NOVO NORDISK LIMITED	XI
Novofem film-coated tablet	DE/H/0304/001	PL 03132/0141	NOVO NORDISK LIMITED	XI
Novofem film-coated tablets	DE/H/0304/001	PA 0218/053/001	NOVO NORDISK A/S	IE
Novofem film-coated tablets	DE/H/0304/001	PA 0218/053/001	NOVO NORDISK A/S	IE
Novofem filmdragerade tabletter	DE/H/0304/001	16601	NOVO NORDISK A/S	FI
Novofem filmdragerade tabletter	DE/H/0304/001	16601	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
Novofem filmdragerade tabletter	DE/H/0304/001	17439	NOVO NORDISK A/S	SE
Novofem filmdragerade tabletter	DE/H/0304/001	17439	NOVO NORDISK A/S	SE
Novofem filmom obalené tablety	DE/H/0304/001	56/0291/05-S	NOVO NORDISK A/S	SK
Novofem filmom obalené tablety	DE/H/0304/001	56/0291/05-S	NOVO NORDISK A/S	SK
Novofem filmom obložene tablete	DE/H/0304/001	HR-H-818589847	NOVO NORDISK A/S	HR
Novofem filmom obložene tablete	DE/H/0304/001	HR-H-818589847	NOVO NORDISK A/S	HR
Novofem filmomhulde tabletten 1 mg/1 mg	DE/H/0304/001	RVG 26864	NOVO NORDISK B.V.	NL
Novofem filmomhulde tabletten 1 mg/1 mg	DE/H/0304/001	RVG 26864	NOVO NORDISK B.V.	NL
Novofem filmovertrukne tabletter	DE/H/0304/001	32745	NOVO NORDISK 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
Novofem filmovertrukne tabletter	DE/H/0304/001	32745	NOVO NORDISK A/S	DK
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/001	NOVO NORDISK A/S	SI
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/002	NOVO NORDISK A/S	SI
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/001	NOVO NORDISK A/S	SI
Novofem filmsko obložene tablete	DE/H/0304/001	H/03/01133/002	NOVO NORDISK A/S	SI
Novofem Filmtabletten	DE/H/0304/001	BE231682	NOVO NORDISK PHARMA SA	BE
Novofem Filmtabletten	DE/H/0304/001	2007039243	NOVO NORDISK PHARMA SA	LU
Novofem Filmtabletten Estradiol / Norethisteronacetat	DE/H/0304/001	47567.00.00	NOVO NORDISK PHARMA GMBH	DE
Novofem Filmtabletten Estradiol / Norethisteronacetat	DE/H/0304/001	47567.00.00	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
Novofem filmuhúðaðar töflur	DE/H/0304/001	IS/1/01/040/01	NOVO NORDISK A/S	IS
Novofem filmuhúðaðar töflur	DE/H/0304/001	IS/1/01/040/01	NOVO NORDISK A/S	IS
Novofem kalvopäällysteiset tabletit	DE/H/0304/001	16601	NOVO NORDISK A/S	FI
Novofem kalvopäällysteiset tabletit	DE/H/0304/001	16601	NOVO NORDISK A/S	FI
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 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 potahované tablety	DE/H/0304/001	56/005/03-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
Novofem potahované tablety	DE/H/0304/001	56/005/03-C	NOVO NORDISK A/S	CZ
Novofem tablett, filmdrasjert	DE/H/0304/001	01-5767	NOVO NORDISK A/S	NO
Novofem tablett, filmdrasjert	DE/H/0304/001	01-5767	NOVO NORDISK A/S	NO
Novofem, comprimés pelliculés	DE/H/0304/001	BE231682	NOVO NORDISK PHARMA SA	BE
Novofem, comprimés pelliculés	DE/H/0304/001	2007039243	NOVO NORDISK PHARMA SA	LU
Novofem, filmomhulde tabletten	DE/H/0304/001	BE231682	NOVO NORDISK PHARMA SA	BE
Novofem, õhukese polümeerikattega tabletid	DE/H/0304/001	407903	NOVO NORDISK A/S	EE
Novofem, õhukese polümeerikattega tabletid	DE/H/0304/001	407903	NOVO NORDISK A/S	EE
Novofem, tabletki powlekane	DE/H/0304/001	11888	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
Novofem, tabletki powlekane	DE/H/0304/001	11888	NOVO NORDISK A/S	PL
NOVOFEMME, comprimé pelliculé	DE/H/0304/001	NL 26893	NOVO NORDISK	FR
NOVOFEMME, comprimé pelliculé	DE/H/0304/001	NL 26893	NOVO NORDISK	FR
Sequidot 2-Phasen transdermale Matrixpflaster	SE/H/0149/002	1-26854	NOVARTIS PHARMA GMBH	AT
Sequidot 2-Phasen transdermale Matrixpflaster	SE/H/0149/002	1-26854	NOVARTIS PHARMA GMBH	AT
Sequidot depotlaastari	SE/H/0149/002	22301	NOVARTIS FINLAND OY	FI
Sequidot depotlaastari	SE/H/0149/002	22301	NOVARTIS FINLAND OY	FI
Sequidot depotplaster	not available	04-3137	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
SEQUIDOT depotplåster	SE/H/0149/002	22301	NOVARTIS FINLAND OY	FI
SEQUIDOT depotplåster	SE/H/0149/002	22301	NOVARTIS FINLAND OY	FI
SEQUIDOT, depotplåster	SE/H/0149/002	21846	NOVARTIS SVERIGE AB	SE
SEQUIDOT, depotplåster	SE/H/0149/002	21846	NOVARTIS SVERIGE AB	SE
SYSTEN CONTI; 3,2 mg + 11,2 mg, system transdermalny, plaster	not available	4447	THERAMEX IRELAND LIMITED	PL
SYSTEN CONTI; 3,2 mg + 11,2 mg, system transdermalny, plaster	not available	4447	THERAMEX IRELAND LIMITED	PL
SYSTEN SEQUI	not available	4448	THERAMEX IRELAND LIMITED	PL
SYSTEN SEQUI	not available	4448	THERAMEX IRELAND LIMITED	PL
Trisekvens filmdragerade tabletter	not available	8424	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
Trisekvens filmdragerade tabletter	not available	8424	NOVO NORDISK A/S	FI
Trisekvens filmovertrukne tabletter	not available	6699	NOVO NORDISK A/S	DK
Trisekvens filmovertrukne tabletter	not available	6699	NOVO NORDISK A/S	DK
Trisekvens filmuhúðaðar töflur	not available	751874	NOVO NORDISK A/S	IS
Trisekvens filmuhúðaðar töflur	not available	751874	NOVO NORDISK A/S	IS
Trisekvens kalvopäällysteiset tabletit	not available	8424	NOVO NORDISK A/S	FI
Trisekvens kalvopäällysteiset tabletit	not available	8424	NOVO NORDISK A/S	FI
Trisekvens tablett, filmdrasjert	not available	6323	NOVO NORDISK A/S	NO
Trisekvens tablett, filmdrasjert	not available	6323	NOVO NORDISK A/S	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
Trisequens - Filmtabletten	not available	16.893	NOVO NORDISK PHARMA GMBH	AT
Trisequens - Filmtabletten	not available	16.893	NOVO NORDISK PHARMA GMBH	AT
Trisequens apvalkotās tabletes	not available	98-0705	NOVO NORDISK A/S	LV
Trisequens apvalkotās tabletes	not available	98-0705	NOVO NORDISK A/S	LV
Trisequens associação comprimidos revestidos por película	not available	2480986	ISDIN LDA	PT
Trisequens associação comprimidos revestidos por película	not available	2480986	ISDIN LDA	PT
Trisequens comprimidos recubiertos	not available	60.706	ISDIN, S.A.	ES
Trisequens comprimidos recubiertos	not available	60.706	ISDIN, S.A.	ES
Trisequens film-coated tablets	not available	PA 218/8/1	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
Trisequens film-coated tablets	not available	PA 218/8/1	NOVO NORDISK A/S	IE
Trisequens filmom obložene tablete	not available	HR-H-919057630	NOVO NORDISK A/S	HR
Trisequens filmom obložene tablete	not available	HR-H-919057630	NOVO NORDISK A/S	HR
Trisequens filmsko obložene tablete	not available	H/97/01568/001	NOVO NORDISK A/S	SI
Trisequens filmsko obložene tablete	not available	H/97/01568/001	NOVO NORDISK A/S	SI
Trisequens filmtabletta	not available	OGYI-T-5851/01	NOVO NORDISK A/S	HU
Trisequens filmtabletta	not available	OGYI-T-5851/01	NOVO NORDISK A/S	HU
Trisequens Filmtabletten	not available	6529829.00.00	NOVO NORDISK PHARMA GMBH	DE
Trisequens Filmtabletten	not available	6529829.00.00	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
Trisequens õhukese polümeerikattega tabletid	not available	205098	NOVO NORDISK A/S	EE
Trisequens õhukese polümeerikattega tabletid	not available	205098	NOVO NORDISK A/S	EE
Trisequens plėvele dengtos tabletės	not available	LT/1/94/0158/001	NOVO NORDISK A/S	LT
Trisequens plėvele dengtos tabletės	not available	LT/1/94/0158/001	NOVO NORDISK A/S	LT
Trisequens potahované tablety	not available	56/307/91-C	NOVO NORDISK A/S	CZ
Trisequens potahované tablety	not available	56/307/91-C	NOVO NORDISK A/S	CZ
Trisequens δισκία επικαλυμμένα με λεπτό υμένιο	not available	13501	NOVO NORDISK HELLAS LTD.	CY
Trisequens δισκία επικαλυμμένα με λεπτό υμένιο	not available	13501	NOVO NORDISK HELLAS LTD.	CY
Trisequens, 2 mg (niebieskie), 2 mg + 1 mg (białe), 1 mg (czerwone), tabletki powlekane	not available	R/3299	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
Trisequens, 2 mg (niebieskie), 2 mg + 1 mg (białe), 1 mg (czerwone), tabletki powlekane	not available	R/3299	NOVO NORDISK A/S	PL
TRISEQUENS, comprimé pelliculé	not available	NL 12375	NOVO NORDISK	FR
TRISEQUENS, comprimé pelliculé	not available	NL 12375	NOVO NORDISK	FR
Trisequens, comprimés pelliculés	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, comprimés pelliculés	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, comprimés pelliculés	not available	2008039743	NOVO NORDISK PHARMA SA	LU
Trisequens, comprimés pelliculés	not available	2008039743	NOVO NORDISK PHARMA SA	LU
Trisequens, filmomhulde tablet	not available	RVG 09812	NOVO NORDISK B.V.	NL
Trisequens, filmomhulde tablet	not available	RVG 09812	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
Trisequens, filmomhulde tabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, filmomhulde tabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, Filmtabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, Filmtabletten	not available	BE156134	NOVO NORDISK PHARMA SA	BE
Trisequens, Filmtabletten	not available	2008039743	NOVO NORDISK PHARMA SA	LU
Trisequens, Filmtabletten	not available	2008039743	NOVO NORDISK PHARMA SA	LU
Trisequens, δισκία επικαλυμμένα με λεπτό υμένιο	not available	40771/10/31-05-2011	NOVO NORDISK HELLAS LTD.	GR
Trisequens, δισκία επικαλυμμένα με λεπτό υμένιο	not available	40771/10/31-05-2011	NOVO NORDISK HELLAS LTD.	GR
Активел 1 mg/0,5 mg филмирани таблетки	SE/H/0150/001	9900351	NOVO NORDISK A/S	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Трисеквенс филмирана таблетки	not available	20010005	NOVO NORDISK A/S	BG
Трисеквенс филмирана таблетки	not available	20010005	NOVO NORDISK A/S	BG