30 January 2020
EMA/117021/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: levonorgestrel

Procedure no.: PSUSA/00001856/201905

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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
28 mini 30 Mikrogramm überzogene Tabletten	not available	3000361.00.00	JENAPHARM GMBH & CO KG	DE
Benilexa 20 microgrammi/24 ore sistema a rilascio intrauterino	UK/H/5593/001	043233016	GEDEON RICHTER PLC.	IT
Benilexa 20 microgrammi/24 ore sistema a rilascio intrauterino.	UK/H/5593/001	043233028	GEDEON RICHTER PLC.	IT
Benilexa 20 micrograms/24 hours Intrauterine Delivery System	UK/H/5593/001	PL 04854/0157	GEDEON RICHTER PLC.	UK
DONASERT 20 microgramе/24 ore sistem cu cedare intrauterină	UK/H/3030/001	11491/2019/01	GEDEON RICHTER ROMÂNIA S.A.	RO
DONASERT 20 microgramе/24 ore sistem cu cedare intrauterină	UK/H/3030/001	5672/2013/01	GEDEON RICHTER ROMÂNIA S.A.	RO
DONASERT 52 mg (20 microgrammes/24 heures), dispositif intra-utérin	not available	34009 301 487 5 4	GEDEON RICHTER PLC.	FR
Emergency Contraceptive Consilient 1500 microgram tablet	UK/H/4569/001	PL 04854/0105	GEDEON RICHTER PLC.	UK
ESCAPELLE 1,5 mg comprimate	not available	8697/2016/01	GEDEON RICHTER PLC.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Escapelle 1,5 mg tableta	not available	UP/I-530-09/13-02/351	GEDEON RICHTER PLC.	HR
Escapelle 1,5 mg tableta	not available	H/08/00568/001	GEDEON RICHTER PLC.	SI
ESCAPELLE 1,5 mg tabletė	CZ/H/0939/001	LT/1/06/0511/001	GEDEON RICHTER PLC.	LT
Escapelle 1,5 mg tabletes	not available	04-0318	GEDEON RICHTER PLC.	LV
Escapelle 1,5 mg tablett	not available	443104	GEDEON RICHTER PLC.	EE
Escapelle 1,5 mg tabletta	not available	OGYI-T-9330/01	GEDEON RICHTER PLC.	HU
Escapelle 1500 microgrammi, compressa	UK/H/0803/001	038802017	GEDEON RICHTER PLC.	IT
ESCAPELLE 1500 mikrogramov tableta	not available	17/0187/04-S	GEDEON RICHTER PLC.	SK
Escapelle 1500 mikrogramů tableta	CZ/H/0939/001	17/168/07-C	GEDEON RICHTER PLC.	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
Escapelle tabletka 1500 mikrogramów, 1500 mikrogramów, tabletka	UK/H/0803/001	12048	GEDEON RICHTER PLC.	PL
Ezinelle 1.5 mg tablet	not available	PL 04569/1486	GENERICS [UK] LIMITED	UK
Fleree 13,5 mg intrauterīnā sistēma	SE/H/1186/001	13-0048	BAYER AG	LV
Fleree 13,5 mg vartojimo į gimdos ertmę sistema	SE/H/1186/001	LT/1/13/3219/001	BAYER AG	LT
Fleree 13,5 mg vartojimo į gimdos ertmę sistema	SE/H/1186/001	LT/1/13/3219/002	BAYER AG	LT
Fleree, 13,5 mg intrauteriinne ravivahend	SE/H/1186/001	805513	BAYER AG	EE
Frivelle, tabletter	DK/H/2357/001	53303	ORIFARM GENERICS A/S	DK
JADELLE 2 x 75 mg implante	FI/H/0128/001	5161781	BAYER PORTUGAL LDA	PT
Jadelle® 2 x 75 mg - implantaatit	FI/H/0128/001	12512	BAYER 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
Jadelle® 2 x 75 mg implantat	FI/H/0128/001	12512	BAYER OY	FI
Jadelle® sine inserter 2 x 75 mg implantaatit	FI/H/0189/001	17098	BAYER OY	FI
Jadelle® sine inserter 2 x 75 mg implantat	FI/H/0189/001	17098	BAYER OY	FI
Jaydess 13,5 mg depotlääkevalmiste, kohtuun	SE/H/1186/001	30387	BAYER OY	FI
Jaydess 13,5 mg dispositivo de libertação intrauterino	SE/H/1186/001	5566617	BERLEX ESPECIALIDADES FARMACEUTICAS LDA	PT
Jaydess 13,5 mg dispositivo de libertação intrauterino	SE/H/1186/001	5566625	BERLEX ESPECIALIDADES FARMACEUTICAS LDA	PT
Jaydess 13,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1186/001	1-31748	BAYER AUSTRIA GMBH	AT
Jaydess 13,5 mg intrauterini dostavni sistem	SE/H/1186/001	H/13/00810/001	BAYER D.O.O	SI
Jaydess 13,5 mg intrauterini dostavni sistem	SE/H/1186/001	H/13/00810/002	BAYER D.O.O	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
Jaydess 13,5 mg intrauterinní inzert	SE/H/1186/001	17/049/13-C	BAYER AG	CZ
Jaydess 13,5 mg intrauterinný inzert	SE/H/1186/001	17/0024/13-S	BAYER SPOL SRO	SK
Jaydess 13,5 mg intrauterint inlägg	SE/H/1186/001	47317	BAYER AB	SE
Jaydess 13,5 mg intrauterint innlegg	SE/H/1186/001	11-8775	BAYER AB	NO
Jaydess 13,5 mg leginnlegg	SE/H/1186/001	IS/1/13/001/01	BAYER AB	IS
Jaydess 13,5 mg méhen belüli gyógyszerleadó rendszer	SE/H/1186/001	OGYI-T-22404/01	BAYER AG	HU
Jaydess 13,5 mg sistem cu cedare intrauterină	SE/H/1186/001	11477/2019/02	BAYER AG	RO
Jaydess 13,5 mg sistem cu cedare intrauterină	SE/H/1186/001	11477/2019/01	BAYER AG	RO
Jaydess 13,5 mg sistema a rilascio intrauterino	SE/H/1186/001	042522019	BAYER SPA	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
Jaydess 13,5 mg sistema a rilascio intrauterino	SE/H/1186/001	042522021	BAYER SPA	IT
Jaydess 13,5 mg sistema de liberación intrauterino	SE/H/1186/001	77.169	BAYER HISPANIA SL	ES
JAYDESS 13,5 mg, système de diffusion intra-utérin	SE/H/1186/001	34009 274 194 7 8	BAYER HEALTHCARE	FR
JAYDESS 13,5 mg, système de diffusion intra-utérin	SE/H/1186/001	34009 585 070 8 3	BAYER HEALTHCARE	FR
Jaydess 13,5 mg afleveringsysteem voor intra-uterien gebruik	SE/H/1186/001	BE433544	BAYER SA NV	BE
Jaydess 13,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1186/001	BE433544	BAYER SA NV	BE
Jaydess 13,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1186/001	2013040105	BAYER SA NV	LU
Jaydess 13,5 mg, système de diffusion intra-utérin	SE/H/1186/001	BE433544	BAYER SA NV	BE
Jaydess 13,5 mg, système de diffusion intra-utérin	SE/H/1186/001	2013040105	BAYER SA NV	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
Jaydess 13.5 mg intrauterine delivery system	SE/H/1186/001	PA 1410/068/001	BAYER LTD	IE
Jaydess 13.5 mg intrauterine delivery system	SE/H/1186/001	MA513/05301	BAYER LTD	MT
Jaydess 13.5 mg intrauterine delivery system	SE/H/1186/001	PL 00010/0587	BAYER PLC	UK
Jaydess 13.5 mg intrauterine delivery system.	SE/H/1186/001	MA639/02101	BAYER LTD	MT
Jaydess, 13,5 mg, system terapeutyczny domaciczny	SE/H/1186/001	21101	BAYER AG	PL
Jaydess, intrauterint indlæg	SE/H/1186/001	49987	BAYER AB	DK
Jaydess® 13,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1186/001	86537.00.00	JENAPHARM GMBH & CO KG	DE
Jaydess® 13,5 mg, intrauterint inlägg	SE/H/1186/001	30387	BAYER OY	FI
Kyleena 19,5 mg afleveringsysteem voor intra-uterien gebruik	SE/H/1587/001	BE502000	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
Kyleena 19,5 mg depotlääkevalmiste, kohtuun	SE/H/1587/001	33805	BAYER OY	FI
Kyleena 19,5 mg dispositivo de libertação intrauterino	SE/H/1587/001	5694930	BAYER PORTUGAL LDA	PT
Kyleena 19,5 mg dispositivo de libertação intrauterino	SE/H/1587/001	5694948	BAYER PORTUGAL LDA	PT
Kyleena 19,5 mg intrauterīna ierīce	SE/H/1587/001	16-0237	BAYER AG	LV
Kyleena 19,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1587/001	137280	BAYER AUSTRIA GMBH	AT
Kyleena 19,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1587/001	BE502000	BAYER SA NV	BE
Kyleena 19,5 mg intrauterini dostavni sistem	SE/H/1587/001	H/17/02281/001	BAYER D.O.O	SI
Kyleena 19,5 mg intrauterini dostavni sistem	SE/H/1587/001	H/17/02281/002	BAYER D.O.O	SI
Kyleena 19,5 mg intrauterini sustav	SE/H/1587/001	HR-H-476936442-01	BAYER DOO	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
Kyleena 19,5 mg intrauterini sustav	SE/H/1587/001	HR-H-476936442-02	BAYER DOO	HR
Kyleena 19,5 mg intrauterinní inzert	SE/H/1587/001	17/460/16-C	BAYER AG	CZ
Kyleena 19,5 mg intrauterinný inzert	SE/H/1587/001	17/0500/16-S	BAYER SPOL SRO	SK
Kyleena 19,5 mg intrauterint inlägg	SE/H/1587/001	53746	BAYER AB	SE
Kyleena 19,5 mg intrauterint innlegg	SE/H/1587/001	15-10974	BAYER AB	NO
Kyleena 19,5 mg sistema de liberación intrauterino	SE/H/1587/001	81.418	BAYER HISPANIA SL	ES
Kyleena 19,5 mg vartojimo į gimdos ertmę sistema	SE/H/1587/001	LT/1/17/4028/001	BAYER AG	LT
Kyleena 19,5 mg vartojimo į gimdos ertmę sistema	SE/H/1587/001	LT/1/17/4028/002	BAYER AG	LT
Kyleena 19,5 mg, afleveringsysteem voor intra-uterien gebruik	SE/H/1587/001	RVG 118462	BAYER BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kyleena 19,5 mg, intrauterint inlägg	SE/H/1587/001	33805	BAYER OY	FI
Kyleena 19,5 mg, système de diffusion intra-utérin	SE/H/1587/001	BE502000	BAYER SA NV	BE
KYLEENA 19,5 mg, système de diffusion intra-utérin	SE/H/1587/001	34009 300 947 5 4	BAYER HEALTHCARE	FR
KYLEENA 19,5 mg, système de diffusion intra-utérin	SE/H/1587/001	34009 550 334 6 2	BAYER HEALTHCARE	FR
Kyleena 19.5 mg intrauterine delivery system	SE/H/1587/001	PA1410/081/001	BAYER LTD	IE
Kyleena 19.5 mg intrauterine delivery system.	SE/H/1587/001	PL 00010/0664	BAYER PLC	UK
Kyleena, 19,5 mg intrauteriinne ravivahend	SE/H/1587/001	922616	BAYER AG	EE
Kyleena, 19,5 mg, system terapeutyczny domaciczny	SE/H/1587/001	23637	BAYER AG	PL
Kyleena, intrauterint indlæg	SE/H/1587/001	56994	BAYER AB	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
Kyleena19,5 mg sistema a rilascio intrauterino	SE/H/1587/001	044756017	BAYER SPA	IT
Kyleena19,5 mg sistema a rilascio intrauterino	SE/H/1587/001	044756029	BAYER SPA	IT
KyleenaTM 19,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1587/001	96796.00.00	JENAPHARM GMBH & CO KG	DE
Levonelle One Step 1500 microgram tablet	not available	PL 04854/0151	GEDEON RICHTER PLC.	UK
Levonelle® 1500 microgram tablet	UK/H/0803/001	PL 04854/0150	GEDEON RICHTER PLC.	UK
Levonorgestrel 1.5 mg tablet	not available	PL 20416/0510	CRESCENT PHARMA LIMITED	UK
Levonorgestrel 1.5 mg tablet	FI/H/1041/001	PL 04854/0136	GEDEON RICHTER PLC.	UK
Levonorgestrel 1.5 mg tablet P.	not available	PL 39352/0394	KOSEI PHARMA UK LIMITED	UK
LEVONORGESTREL BIOGARAN 1,5 mg, comprimé	UK/H/0803/001	373 075-6	GEDEON RICHTER PLC.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levonorgestrel Richter 20 micrograms/24 hours Intrauterine Delivery System	UK/H/6755/001	PL 04854/0159	GEDEON RICHTER PLC.	UK
Levonortis 20 microgram/24 uur, afleveringsysteem voor intra-uterien gebruik	UK/H/5666/01/DC	BE463626	EXELTIS GERMANY GMBH	BE
Levonortis 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5666/01/DC	BE463626	EXELTIS GERMANY GMBH	BE
Levonortis 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5666/01/DC	2015040082	EXELTIS GERMANY GMBH	LU
Levonortis 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5666/01/DC	2015040082	EXELTIS GERMANY GMBH	LU
Levonortis 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5666/01/DC	2015040082	EXELTIS GERMANY GMBH	LU
Levonortis 20 Mikrogramm/24 Stunden, Intrauterines Wirkstofffreisetzungssystem	UK/H/5666/01/DC	BE463626	EXELTIS GERMANY GMBH	BE
Levosert 0,02 mg cada 24 horas sistema de liberación intrauterino	UK/H/5593/001	80021	GEDEON RICHTER PLC.	ES
Levosert 20 microgram/24 hours Intrauterine Delivery System	UK/H/3030/001	PL 04854/0158	GEDEON RICHTER PLC.	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
Levosert 20 microgram/24 uur, afleveringsysteem voor intra-uterien gebruik	UK/H/5491/001	BE446586	MITHRA PHARMACEUTICALS SA	BE
Levosert 20 microgramas/24 horas Dispositivo de Libertação Intrauterino	UK/H/6755/001	5762836	GEDEON RICHTER PLC.	PT
Levosert 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5491/001	BE446586	MITHRA PHARMACEUTICALS SA	BE
Levosert 20 microgrammes/24 heures, système de diffusion intra-utérin	UK/H/5491/01/DC	2014060143	MITHRA PHARMACEUTICALS SA	LU
Levosert 20 micrograms/24 hours Intrauterine Delivery System	UK/H/5593/001	MA 1031/00401	GEDEON RICHTER PLC.	MT
Levosert 20 mikrogram/24 timer intrauterint innlegg	UK/H/5593/001	13-9909	GEDEON RICHTER PLC.	NO
Levosert 20 mikrogram/24 timmar intrauterint inlägg	DK/H/3066/001	50571	GEDEON RICHTER PLC.	SE
Levosert 20 mikrograma/24 sata intrauterini sustav za isporuku	PT/H/2373/001	HR-H-896425945	GEDEON RICHTER PLC.	HR
Levosert 20 mikrogramm/24 óra méhen belüli gyógyszerleadó rendszer	UK/H/3030/001	OGYI-T-22505/01	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
Levosert 20 Mikrogramm/24 Stunden intrauterines Wirkstofffreisetzungssystem	DK/H/3066/001	136091	GEDEON RICHTER PLC.	AT
Levosert 20 Mikrogramm/24 Stunden intrauterines Wirkstofffreisetzungssystem	UK/H/5593/001	91463.00.00	GEDEON RICHTER PLC.	DE
Levosert 20 Mikrogramm/24 Stunden, Intrauterines Wirkstofffreisetzungssystem	UK/H/5491/001	BE446586	MITHRA PHARMACEUTICALS SA	BE
Levosert 20 Mikrogramm/24 Stunden, Intrauterines Wirkstofffreisetzungssystem	UK/H/5491/001	2014060143	MITHRA PHARMACEUTICALS SA	LU
Levosert 20 mikrogrammi/24 tunnis intrauteriinne ravivahend	PT/H/2373/001	979519	GEDEON RICHTER PLC.	EE
Levosert 20 mikrogramov/ 24 ur intrauterini dostavni sistem	UK/H/5593/001	H/15/02074/001	GEDEON RICHTER PLC.	SI
Levosert 20 mikrogramov/ 24 ur intrauterini dostavni sistem	UK/H/5593/001	H/15/02074/002	GEDEON RICHTER PLC.	SI
Levosert 20 mikrogramov/24 hodín, intrauterinný inzert	UK/H/3030/001	17/0236/13-S	GEDEON RICHTER PLC.	SK

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
LEVOSERT 20 mikrogramų / 24 valandų vartojimo į gimdos ertmę sistema	UK/H/3030/001	LT/1/13/3319/001	GEDEON RICHTER PLC.	LT
Levosert 20 mikrogramů/24 hodin, intrauterinní inzert	UK/H/3030/001	17/285/13-C	GEDEON RICHTER PLC.	CZ
Levosert 20 mikrogramu/24 stundās intrauterīna sistēma	UK/H/3030/001	13-0032	GEDEON RICHTER PLC.	LV
Levosert 20 míkrógrömm/24 klst. leginnlegg	UK/H/5593/001	IS/1/15/008/01	GEDEON RICHTER PLC.	IS
Levosert 20 μικρογραμμάρια/24 ώρες Ενδομήτρια Συσκευή Αποδέσμευσης	UK/H/5593/001	022402	GEDEON RICHTER PLC.	CY
Levosert 52 mg Intrauterine Delivery System	UK/H/5593/001	PA 1330/022/001	GEDEON RICHTER PLC.	IE
Levosert, 52 mg (20 mikrogramów/24 h), system terapeutyczny domaciczny	UK/H/3030/001	21336	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
Levosert, intrauterint indlæg	UK/H/5593/001	53445	GEDEON RICHTER PLC.	DK
Luadei 13.5 mg intrauterine delivery system	SE/H/1187/001	47573	BAYER 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
Luadei® 13,5 mg intrauterines Wirkstofffreisetzungssystem	SE/H/1187/001	86538.00.00	JENAPHARM GMBH & CO KG	DE
Microlut 30 Mikrogramm überzogene Tabletten	not available	6929492.00.00	JENAPHARM GMBH & CO KG	DE
Microluton® 30 mikrog -tabletti, päällystetty	not available	6424	BAYER OY	FI
Microluton® 30 mikrog, dragerad tablett	not available	6424	BAYER OY	FI
MICROVAL, comprimé enrobé	not available	34009 322 006 7 2	PFIZER HOLDING FRANCE	FR
MICROVAL, comprimé enrobé	not available	34009 322 007 3 3	PFIZER HOLDING FRANCE	FR
Mirena 0,02 mg cada 24 horas sistema de liberación intrauterino	PT/H/0100/001	63.158	BAYER HISPANIA SL	ES
Mirena 20 microgram / 24 uur, afleveringsysteem voor intra-uterien gebruik (IUS)	not available	BE170737	BAYER SA NV	BE
Mirena 20 microgramas/24 horas dispositivo de libertação intrauterino	PT/H/0100/001	2477784	BAYER PORTUGAL 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
Mirena 20 microgame/24 ore sistem cu cedare intrauterină	not available	7842/2015/01	BAYER AG	RO
Mirena 20 microgrammes / 24 heures, système de diffusion intra-utérin (SIU)	not available	BE170737	BAYER SA NV	BE
Mirena 20 microgrammes / 24 heures, système de diffusion intra-utérin (SIU)	not available	2011051166	BAYER SA NV	LU
Mirena 20 microgrammi/24ore sistema a rilascio intrauterino	not available	029326016	BAYER AG	IT
Mirena 20 mikg/24 timer intrauterint indlæg	not available	14881	BAYER AB	DK
Mirena 20 mikrog/24 timmar intrauterint inlägg	not available	10212	BAYER OY	FI
Mirena 20 mikrog/24 tuntia depotlääkevalmiste, kohtuun	not available	10212	BAYER OY	FI
Mirena 20 mikrogram/24 timer intrauterint innlegg	not available	7935	BAYER AB	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mirena 20 mikrogram/24 timmar intrauterint inlägg	not available	11668	BAYER AB	SE
Mirena 20 mikrograma/24 sata intrauterini sustav	not available	HR-H-494254913-01	BAYER DOO	HR
Mirena 20 Mikrogramm/24 Stunden Intrauterinpessar	not available	1-21529	BAYER AUSTRIA GMBH	AT
Mirena 20 Mikrogramm/24 Stunden, intrauterines Wirkstofffreisetzungssystem (IUS)	not available	BE170737	BAYER SA NV	BE
Mirena 20 Mikrogramm/24 Stunden, intrauterines Wirkstofffreisetzungssystem (IUS)	not available	2011051166	BAYER SA NV	LU
Mirena 20 mikrogramov/24 hodín intrauterinný inzert	not available	17/0234/99-S	BAYER SPOL SRO	SK
Mirena 20 mikrogramov/24 ur intrauterini dostavni sistem	not available	H/99/01031/001	BAYER D.O.O	SI
Mirena 20 mikrogramu/ 24 stundās intrauterīna sistēma	not available	99-0502	BAYER AG	LV
Mirena 20 mikrogramų/24 valandoms vartojimo į	not available	LT/1/98/0172/001	BAYER AG	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
gimdos ertmę sistema				
Mirena 20 míkrógrömm/24 klst. leginnlegg	not available	MTNR 920074 (IS)	BAYER AB	IS
MIRENA 52 mg (20 microgrammes/24 heures), dispositif intra-utérin	not available	34009 339 292 8 2	BAYER HEALTHCARE	FR
Mirena 52 mg Intrauterine Delivery System	not available	PA 1410/8/1	BAYER LTD	IE
Mirena 52 mg intrauterinní inzert	not available	17/372/97-C	BAYER AG	CZ
Mirena méhen belüli gyógyszerleadó rendszer	not available	OGYI-T-6210/01	BAYER AG	HU
Mirena, 20 mikrogrammi/24 tunnis intrauteriinne ravivahend	not available	127696	BAYER AG	EE
Mirena, 20 mikrogramów/24 h, system terapeutyczny domaciczny	PT/H/0100/001	11959	BAYER AG	PL
Mirena, IUD 20 microgram/24 uur	not available	RVG 16681	BAYER BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mirena® 20 micrograms/24 hours intrauterine delivery system	not available	MA513/03901	BAYER LTD	MT
Mirena® 20 micrograms/24 hours intrauterine delivery system	not available	PL 00010/0547	BAYER PLC	UK
Mirena®, 52 mg, Intrauterinpessar mit Hormonabgabe	not available	30495.00.00	JENAPHARM GMBH & CO KG	DE
Mirena®, 52mg, Intrauterinpessar mit Hormonabgabe	not available	41880.00.00	JENAPHARM GMBH & CO KG	DE
Mirena®, ενδομήτριο εξάρτημα απελευθέρωσης λεβονοργεστρέλης, 52 mg/εξάρτημα	not available	18612	BAYER HELLAS SA	CY
Mirena®, ενδομήτριο εξάρτημα απελευθέρωσης λεβονοργεστρέλης, 52 mg/εξάρτημα	not available	20731/1-4-2008	BAYER HELLAS SA	GR
Norgeston®	not available	00010/0550	BAYER PLC	UK
NORLEVO 1,5 mg compresse	FR/H/0146/002	034884078	LABORATOIRE HRA PHARMA	IT
NORLEVO 1,5 mg compresse	FR/H/0146/002	034884080	LABORATOIRE HRA PHARMA	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
NORLEVO 1,5 mg compresse	FR/H/0146/002	034884092	LABORATOIRE HRA PHARMA	IT
NORLEVO 1,5 mg compresse	FR/H/0146/002	034884104	LABORATOIRE HRA PHARMA	IT
NORLEVO 1,5 mg compresse	FR/H/0146/002	034884066	LABORATOIRE HRA PHARMA	IT
Norlevo 1,5 mg comprimés	FR/H/0146/002	BE277356	LABORATOIRE HRA PHARMA	BE
Norlevo 1,5 mg comprimido	not available	5607288	LABORATOIRE HRA PHARMA	PT
NORLEVO 1,5 mg tableta	FR/H/0146/002	H/10/01127/001-005	LABORATOIRE HRA PHARMA	SI
NorLevo 1,5 mg tablett	FR/H/0146/002	22234	LABORATOIRE HRA PHARMA	SE
NORLEVO 1,5 mg tabletten	FR/H/0146/002	BE277356	LABORATOIRE HRA PHARMA	BE
NORLEVO 1,5 mg tabletten	FR/H/0146/002	BE277356	LABORATOIRE HRA PHARMA	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
Norlevo 1,5 mg Tabletten	FR/H/0146/002	2005100005	LABORATOIRE HRA PHARMA	LU
Norlevo 1,5 mg tabletter	not available	05-3426	LABORATOIRE HRA PHARMA	NO
NORLEVO 1,5 mg, comprimé	FR/H/0146/002	34009 565 510 2 6	LABORATOIRE HRA PHARMA	FR
NORLEVO 1,5 mg, comprimé	FR/H/0146/002	34009 565 511 9 4	LABORATOIRE HRA PHARMA	FR
NORLEVO 1,5 mg, comprimé	FR/H/0146/002	34009 565 512 5 5	LABORATOIRE HRA PHARMA	FR
NORLEVO 1,5 mg, comprimé	FR/H/0146/002	34009 565 513 1 6	LABORATOIRE HRA PHARMA	FR
NORLEVO 1,5 mg, comprimé	FR/H/0146/002	34009 364 137 2 6	LABORATOIRE HRA PHARMA	FR
NorLevo 1,5 mg, tabletten	FR/H/0146/002	RVG 32303	LABORATOIRE HRA PHARMA	NL
NorLevo 1,5mg comprimido	not available	67.770	LABORATOIRE HRA PHARMA	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
NORLEVO 1.5 mg tablet	FR/H/0146/002	PA 1166/2/1	LABORATOIRE HRA PHARMA	IE
Norlevo 750 mikrogram tabletter	not available	MTNR 99-7321	LABORATOIRE HRA PHARMA	NO
Norlevo δισκίο των 1,5 mg	not available	20222	LABORATOIRE HRA PHARMA	CY
Norlevo δισκίο των 1,5 mg	FR/H/0146/002	74875/20-12-2012	LABORATOIRE HRA PHARMA	GR
NorLevo, tabletter	FR/H/0146/002	37788	LABORATOIRE HRA PHARMA	DK
NORLEVO® 1,5 mg comprimés	FR/H/0146/002	2005100005	LABORATOIRE HRA PHARMA	LU
NORLEVO® 1,5 mg - tabletti	FR/H/0146/002	20803	LABORATOIRE HRA PHARMA	FI
NORLEVO® 1,5 mg - tabletti	FR/H/0146/002	20803	LABORATOIRE HRA PHARMA	FI
PiDaNa 1,5 mg Tablette	not available	75074.00.000	LABORATOIRE HRA PHARMA	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
POSTINOR - DUO 750 mikrogramų tabletės	not available	LT/1/03/3098/001	GEDEON RICHTER PLC.	LT
Postinor 1,5 mg comprimido	CZ/H/0939/001	67.515	GEDEON RICHTER PLC.	ES
Postinor 1,5 mg tablett	CZ/H/0939/001	22194	GEDEON RICHTER PLC.	SE
Postinor 1,5 mg tablett	CZ/H/0939/001	05-3275	GEDEON RICHTER PLC.	NO
Postinor 1,5 mg tabletti	FI/H/1041/001	32491	GEDEON RICHTER PLC.	FI
Postinor 1,5 mg, tafla	CZ/H/0939/001	IS/I/05/016/01	GEDEON RICHTER PLC.	IS
Postinor 1500 microgram tablet.	CZ/H/0939/001	BE285196	GEDEON RICHTER PLC.	BE
Postinor 1500 microgram, tablet	CZ/H/0939/001	RVG 32253	GEDEON RICHTER PLC.	NL
Postinor 1500 microgramas comprimido	UK/H/0803/001	5693189	GEDEON RICHTER PLC.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Postinor 1500 microgrammes comprimé.	CZ/H/0939/001	BE285196	GEDEON RICHTER PLC.	BE
POSTINOR 1500 microgrammes comprimé.	CZ/H/0939/001	2006010037	GEDEON RICHTER PLC.	LU
Postinor 1500 Mikrogramm - Tablette	CZ/H/0939/001	1-26693	GEDEON RICHTER PLC.	AT
Postinor 1500 Mikrogramm Tablette	CZ/H/0939/001	62588.00.00	GEDEON RICHTER PLC.	DE
Postinor 1500 Mikrogramm Tabletten	CZ/H/0939/001	BE285196	GEDEON RICHTER PLC.	BE
Postinor 1500 μικρογραμμάρια δισκίο	UK/H/0803/001	39664/9-6-2015	GEDEON RICHTER PLC.	GR
POSTINOR, 1500 mikrogrammi tablett	UK/H/4569/001	781812	GEDEON RICHTER PLC.	EE
Postinor-1 1 500 mikrogramov tableta	UK/H/4569/001	56/0201/12-S	GEDEON RICHTER PLC.	SK
Postinor-2 750 micrograme comprimate	not available	8348/2015/01	GEDEON RICHTER PLC.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
POSTINOR-2 750 mikrogramů tablety	not available	17/834/99-C	GEDEON RICHTER PLC.	CZ
POSTINOR-DUO 750 mikrogrami tabletes	not available	96-0366	GEDEON RICHTER PLC.	LV
Prevenelle 1500 microgram tablet	UK/H/0803/001	PA1330/021/001	GEDEON RICHTER PLC.	IE
Ramonna 1,5 mg tableta	CZ/H/0940/001	H/12/01318/001	GEDEON RICHTER PLC.	SI
Ramonna 1500 micrograme comprimate	UK/H/4569/001	4737/2012/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Ramonna 1500 mikrogrami tablete	UK/H/4569/001	12-0131	GEDEON RICHTER PLC.	LV
Ramonna 1500 mikrogramů tableta	CZ/H/0940/001	17/126/12-C	GEDEON RICHTER PLC.	CZ
RAMONNA 1500 mikrogramų tabletė	CZ/H/0940/001	LT/1/12/2856/001	GEDEON RICHTER PLC.	LT
Ramonna, 1500 mikrogramów, tabletka	UK/H/4569/001	20088	GEDEON RICHTER POLSKA 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
Rigesoft 750 mikrogramm tabletta	not available	OGYI-T-6372/01	GEDEON RICHTER PLC.	HU
Upostelle 1500 microgram tablet	not available	PL04854/0106	GEDEON RICHTER PLC.	UK
Vikela 1,5 mg - Tablette	FR/H/0146/002	1-26057	LABORATOIRE HRA PHARMA	AT
Джейдас 13,5 mg вътрематочна лекарстводоставяща система	SE/H/1186/001	20130075	BAYER AG	BG
Ескапеле 1,5 mg таблетка	not available	20060032	GEDEON RICHTER PLC.	BG
ЛЕВОСЕРТ 20 МИКРОГРАМА/24 ЧАСА ВЪТРЕМАТОЧНА ЛЕКАРСТВОДОСТАВЯЩА СИСТЕМА	UK/H/3030/001	20130077	GEDEON RICHTER PLC.	BG
Мирена 20 микрограма/24 часа вътрематочна лекарстводоставяща система	not available	9800344	BAYER AG	BG
Постинор-Дуо 750 микрограма таблетки	not available	20000358	GEDEON RICHTER PLC.	BG
Рамона 1500 микрограма таблетка	CZ/H/0940/001	20120195	GEDEON RICHTER PLC.	BG