28 January 2021
EMA/270645/2015
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: tramadol

Procedure no.: PSUSA/00003002/202005

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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
ADAMON 100 mg retard kemény kapszula	not available	OGYI-T-6970/04	MYLAN EPD KFT.	HU
ADAMON 100 mg retard kemény kapszula	not available	OGYI-T-6970/05	MYLAN EPD KFT.	HU
ADAMON 100 mg retard kemény kapszula	not available	OGYI-T-6970/06	MYLAN EPD KFT.	HU
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/08	MYLAN EPD KFT.	HU
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/07	MYLAN EPD KFT.	HU
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/09	MYLAN EPD KFT.	HU
ADAMON 200 mg retard kemény kapszula	not available	OGYI-T-6970/10	MYLAN EPD KFT.	HU
ADAMON 200 mg retard kemény kapszula	not available	OGYI-T-6970/11	MYLAN EPD KFT.	HU
ADAMON 200 mg retard kemény kapszula	not available	OGYI-T-6970/12	MYLAN EPD KFT.	HU
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/01	MYLAN EPD KFT.	HU
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/02	MYLAN EPD KFT.	HU
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/03	MYLAN EPD KFT.	HU
Adamon inject, 50 mg / ml Injektionslösung	DE/H/0306/002	28667.00.00	MEDA PHARMA GMBH & CO. KG	DE
Adamon Kaps 50 mg, Hartkapseln	DE/H/0306/001	28661.00.00	MEDA PHARMA GMBH & CO. KG	DE
Adamon long retard 150 mg-Filmtabletten	UK/H/0306/001	1-23430	MEDA PHARMA GMBH	AT
Adamon long retard 300 mg-Filmtabletten	UK/H/0306/003	1-23432	MEDA PHARMA GMBH	AT
Adamon SR 100 mg, tvrdé kapsuly s	UK/H/0301/002	65/0182/06-S	MYLAN IRE HEALTHCARE LIMITED	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
predĺženým uvoľňovaním				
Adamon SR 100, 100 mg, kapsułki o przedłużonym uwalnianiu	not available	9361	MYLAN HEALTHCARE SP. Z.O.O.	PL
Adamon SR 150 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/003	65/0183/06-S	MYLAN IRE HEALTHCARE LIMITED	SK
Adamon SR 150, 150 mg, kapsułki o przedłużonym uwalnianiu	not available	9362	MYLAN HEALTHCARE SP. Z.O.O.	PL
Adamon SR 200 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/004	65/0185/06-S	MYLAN IRE HEALTHCARE LIMITED	SK
Adamon SR 200, 200 mg, kapsułki o przedłużonym uwalnianiu	not available	9363	MYLAN HEALTHCARE SP. Z.O.O.	PL
Adamon SR 50 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/001	65/0184/06-S	MYLAN IRE HEALTHCARE LIMITED	SK
Adamon SR 50, 50 mg, kapsułki o przedłużonym uwalnianiu	not available	9360	MYLAN HEALTHCARE SP. Z.O.O.	PL
Adamon Tropfen 100 mg/ml Tropfen zum Einnehmen, Lösung	DE/H/0306/003	28664.00.00	MEDA PHARMA GMBH & CO. KG	DE
Adolonta 100 mg/ 2 ml solución inyectable	not available	59.086	GRÜNENTHAL PHARMA S.A.	ES
Adolonta 100 mg/ml gotas orales en solución	not available	61617	GRÜNENTHAL PHARMA S.A.	ES
Adolonta 50 mg cápsulas duras	not available	59.088	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA 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
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 200 mg comprimidos de liberación prolongada	DE/H/0108/003	61786	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 50 mg comprimidos de liberación prolongada	DE/H/0108/004	68570	GRÜNENTHAL PHARMA S.A.	ES
Amadol Retard 100 mg Hartkapseln, retardiert	UK/H/0301/002	46342.01.00	MEDA PHARMA GMBH & CO. KG	DE
Amadol Retard 150 mg Hartkapseln, retardiert	UK/H/0301/003	46342.02.00	MEDA PHARMA GMBH & CO. KG	DE
Amadol Retard 200 mg Hartkapseln, retardiert	UK/H/0301/004	46342.03.00	MEDA PHARMA GMBH & CO. KG	DE
Amadol Retard 50 mg Hartkapseln, retardiert	UK/H/0301/001	46342.00.00	MEDA PHARMA GMBH & CO. KG	DE
Amadol® Kapseln	not available	32283.00.00	TAD PHARMA GMBH	DE
Amadol® Kapseln	not available	32283.00.00	TAD PHARMA GMBH	DE
Awardix 100 mg otopina za injekciju/infuziju	HR/H/0104/ 002	HR-H-840360429	KRKA, D.D., NOVO MESTO	HR
Awardix 100 mg/ml oralne kapi, otopina	HR/H/0126/001	HR-H-910956418	KRKA, D.D., NOVO MESTO	HR
Awardix 50 mg otopina za injekciju/infuziju	HR/H/0104/ 002	HR-H-944643023	KRKA, D.D., NOVO MESTO	HR
Awardix 50 mg tvrde kapsule	HR/H/0104/ 003	HR-H-177611449	KRKA, D.D., NOVO MESTO	HR
Awardix, 100 mg toimeainet prolongeeritult vabastavad tabletid	EE/H/0269/001	975218	KRKA, D.D., NOVO MESTO	EE
Awardix, 150 mg toimeainet prolongeeritult	EE/H/0269/002	975318	KRKA, D.D., NOVO MESTO	EE

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vabastavad tabletid				
Awardix, 200 mg toimeainet prolongeeritult vabastavad tabletid	EE/H/0269/003	975418	KRKA, D.D., NOVO MESTO	EE
BIODALGIC 50 mg, COMPRIME EFFERVESCENT	not available	561 772-2	BIOCODEX	FR
BIODALGIC 50 mg, COMPRIME EFFERVESCENT	not available	350 673-4	BIOCODEX	FR
Brimisol PR 100 mg Prolonged-Release Tablet	not available	PL 17907/0134	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Brimisol PR 200 mg Prolonged-Release Tablet	not available	PL 17907/0136	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
By-Madol SR 100 mg prolonged-release capsules, hard	DE/H/0639/002	PA 0549/016/002	ETHYPHARM	IE
By-Madol SR 150 mg prolonged-release capsules, hard	DE/H/0639/003	PA 0549/016/003	ETHYPHARM	IE
By-Madol SR 200 mg prolonged-release capsules, hard	DE/H/0639/004	PA 0549/016/004	ETHYPHARM	IE
By-Madol SR 50 mg prolonged-release capsules, hard	DE/H/0639/001	PA 0549/016/001	ETHYPHARM	IE
CONTRAMAL 100 mg compresse a rilascio prolungato	not available	028853036	GRÜNENTHAL ITALIA S.R.L.	IT
Contramal 100 mg retard filmtabletta	not available	OGYI-T-4975/07	STADA ARZNEIMITTEL AG	HU
Contramal 100 mg retard filmtabletta	not available	OGYI-T-4975/08	STADA ARZNEIMITTEL AG	HU

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Contramal 100 mg végbélkúp	not available	OGYI-T-4975/06	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 100 mg/2 ml oplossing voor injectie of infusie	not available	BE 163037	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/2 ml solution injectable ou pour perfusion	not available	BE 163037	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/2 ml soluzione iniettabile	not available	028853063	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL 100 mg/2 ml, solution injectable	not available	34009 561 113-9 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Contramal 100 mg/ml belsőleges oldatos cseppek	not available	OGYI-T-4975/04	STADA ARZNEIMITTEL AG	HU
Contramal 100 mg/ml belsőleges oldatos cseppek	not available	OGYI-T-4975/05	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE 190836	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE 163046	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/ml gocce orali soluzione con contagocce	not available	028853024	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL 100 mg/ml solution buvable en gouttes	not available	BE 190836	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/ml solution buvable en gouttes	not available	BE 163046	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 100 mg/ml	not available	028853101	GRÜNENTHAL 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
soluzione orale con erogatore				
CONTRAMAL 100 mg/ml, solution buvable	not available	34009 362 068 3 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL 150 mg compresse a rilascio prolungato	DE/H/0108/002	028853 075	GRÜNENTHAL ITALIA S.R.L.	IT
Contramal 150 mg retard filmtabletta	not available	OGYI-T-4975/10	STADA ARZNEIMITTEL AG	HU
Contramal 150 mg retard filmtabletta	not available	OGYI-T-4975/09	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 200 mg compresse a rilascio prolungato	DE/H/0108/003	028853 087	GRÜNENTHAL ITALIA S.R.L.	IT
Contramal 200 mg retard filmtabletta	not available	OGYI-T-4975/12	STADA ARZNEIMITTEL AG	HU
Contramal 200 mg retard filmtabletta	not available	OGYI-T-4975/11	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 50 mg capsule rigide	not available	028853012	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL 50 mg capsules, hard	not available	BE 163055	SA GRÜNENTHAL N.V.	BE
CONTRAMAL 50 mg gélules	not available	BE163055	SA GRÜNENTHAL N.V.	BE
Contramal 50 mg kemény kapszula	not available	OGYI-T-4975/02	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 50 mg, gélule	not available	34009 300 022 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL 50 mg, gélule	not available	34009 550 009 4 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL 50 mg, gélule	not available	34009 561 112 2 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL 50 mg, gélule	not available	34009 348 081 6 6	LABORATOIRES GRÜNENTHAL 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
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/13	STADA ARZNEIMITTEL AG	HU
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/03	STADA ARZNEIMITTEL AG	HU
CONTRAMAL 50 mg/ml soluzione iniettabile	not available	028853051	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 348 276-1 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 351 629-9 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 561 140-6 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 561 141-2 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 351 628-2 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 100 mg, comprimé à libération prolongée	DE/H/0136/001	34009 348 275-5 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 150 mg, comprimé à libération prolongée	DE/H/0136/002	34009 561 142-9 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 150 mg, comprimé à libération prolongée	DE/H/0136/002	34009 348 278-4 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 150 mg, comprimé à libération prolongée	DE/H/0136/002	34009 351 630-7 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 150 mg,	DE/H/0136/002	34009 351 631-3 4	LABORATOIRES	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
CONTRAMAL LP 150 mg, comprimé à libération prolongée	DE/H/0136/002	34009 348 277-8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 150 mg, comprimé à libération prolongée	DE/H/0136/002	34009 561 143-5 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 351 677-3 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 351 676-7 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 348 279-0 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 348 280-9 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 561 144-1 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
CONTRAMAL LP 200 mg, comprimé à libération prolongée	DE/H/0136/003	34009 561 145-8 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à	DE/H/0108/001	BE364743	SA GRÜNENTHAL 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
libération prolongée				
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	BE364743	SA GRÜNENTHAL 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
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	2003070150	SA GRÜNENTHAL N.V.	LU
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE

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mg tabletten met verlengde afgifte				
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE364743	SA GRÜNENTHAL N.V.	BE
Contramal Retard 100 mg tabletten met verlengde afgifte	DE/H/0108/001	BE188483	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE

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libération prolongée				
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002/MR	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	2003070152	SA GRÜNENTHAL N.V.	LU
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE

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Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg tabletten met verlengde afgifte	DE/H/0108/002	BE188474	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200	DE/H/0108/003	BE364761	SA GRÜNENTHAL 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
mg comprimés à libération prolongée				
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à	DE/H/0108/003	BE364761	SA GRÜNENTHAL 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
libération prolongée				
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	2003070151	SA GRÜNENTHAL N.V.	LU
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL 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
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE364761	SA GRÜNENTHAL N.V.	BE
Contramal Retard 200 mg tabletten met verlengde afgifte	DE/H/0108/003	BE188465	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg	DE/H/0108/004	BE290376	SA GRÜNENTHAL 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
comprimés à libération prolongée				
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg comprimés à libération prolongée	DE/H/0108/004	2007040035	SA GRÜNENTHAL N.V.	LU
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde	DE/H/0108/004	BE290376	SA GRÜNENTHAL 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
afgifte				
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290385	SA GRÜNENTHAL N.V.	BE
Contramal Retard 50 mg tabletten met verlengde afgifte	DE/H/0108/004	BE290376	SA GRÜNENTHAL 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
Dolatramyl	DK/H/2942/001	48179	MYLAN AB	DK
Dolatramyl	DK/H/2942/001	48179	MYLAN AB	DK
Dolatramyl	DK/H/2942/001	48179	MYLAN AB	DK
Dolatramyl	DK/H/2942/001	48179	MYLAN AB	DK
Dolatramyl 100 mg depottabletter	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletter	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletter	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletter	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletter	DK/H/2942/001	10-8115	MYLAN AB	NO
Dolatramyl 100 mg depottabletter	DK/H/2942/001	10-8115	MYLAN AB	NO
Dolatramyl 100 mg depottabletter	DK/H/2942/001	10-8115	MYLAN AB	NO
Dolatramyl 100 mg depottabletter	DK/H/2942/001	10-8115	MYLAN AB	NO
Dolatramyl 100 mg depottabletter.	DK/H/2942/001	45573	MYLAN AB	SE
Dolatramyl 100 mg depottabletter.	DK/H/2942/001	45573	MYLAN AB	SE
Dolatramyl 100 mg depottabletter.	DK/H/2942/001	45573	MYLAN AB	SE
Dolatramyl 100 mg depottabletter.	DK/H/2942/001	45573	MYLAN AB	SE
Dolatramyl 100 mg depottabletti	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletti	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 100 mg depottabletti	DK/H/2942/001	29450	MYLAN AB	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
Dolatramyl 100 mg depottabletti	DK/H/2942/001	29450	MYLAN AB	FI
Dolatramyl 150 mg depottabletter	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletter	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletter	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletter	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletter	DK/H/2942/002	10-8116	MYLAN AB	NO
Dolatramyl 150 mg depottabletter	DK/H/2942/002	10-8116	MYLAN AB	NO
Dolatramyl 150 mg depottabletter	DK/H/2942/002	10-8116	MYLAN AB	NO
Dolatramyl 150 mg depottabletter	DK/H/2942/002	10-8116	MYLAN AB	NO
Dolatramyl 150 mg depottabletter.	DK/H/2942/002	45574	MYLAN AB	SE
Dolatramyl 150 mg depottabletter.	DK/H/2942/002	45574	MYLAN AB	SE
Dolatramyl 150 mg depottabletter.	DK/H/2942/002	45574	MYLAN AB	SE
Dolatramyl 150 mg depottabletter.	DK/H/2942/002	45574	MYLAN AB	SE
Dolatramyl 150 mg depottabletti	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletti	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletti	DK/H/2942/002	29451	MYLAN AB	FI
Dolatramyl 150 mg depottabletti	DK/H/2942/002	29451	MYLAN AB	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
Dolatramyl 200 mg depottabletter	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletter	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletter	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletter	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletter	DK/H/2942/003	10-8117	MYLAN AB	NO
Dolatramyl 200 mg depottabletter	DK/H/2942/003	10-8117	MYLAN AB	NO
Dolatramyl 200 mg depottabletter	DK/H/2942/003	10-8117	MYLAN AB	NO
Dolatramyl 200 mg depottabletter	DK/H/2942/003	10-8117	MYLAN AB	NO
Dolatramyl 200 mg depottabletter.	DK/H/2942/003	45575	MYLAN AB	SE
Dolatramyl 200 mg depottabletter.	DK/H/2942/003	45575	MYLAN AB	SE
Dolatramyl 200 mg depottabletter.	DK/H/2942/003	45575	MYLAN AB	SE
Dolatramyl 200 mg depottabletter.	DK/H/2942/003	45575	MYLAN AB	SE
Dolatramyl 200 mg depottabletti	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletti	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletti	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl 200 mg depottabletti	DK/H/2942/003	29452	MYLAN AB	FI
Dolatramyl, depottabletter	DK/H/2942/002	48180	MYLAN 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
Dolatramyl, depottabletter	DK/H/2942/003	48181	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/002	48180	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/003	48181	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/002	48180	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/003	48181	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/002	48180	MYLAN AB	DK
Dolatramyl, depottabletter	DK/H/2942/003	48181	MYLAN AB	DK
Dolol Retard UNO, depotkapsler, hårde	FI/H/0164/001	33911	TAKEDA PHARMA A/S	DK
Dolol Retard UNO, depotkapsler, hårde	FI/H/0164/002	33912	TAKEDA PHARMA A/S	DK
Dolol Retard UNO, depotkapsler, hårde	FI/H/0164/003	33913	TAKEDA PHARMA A/S	DK
Dolol, brusetabletter	not available	19173	TAKEDA PHARMA A/S	DK
Dolol, hårde kapsler	DK/H/0141/002	18099	TAKEDA PHARMA A/S	DK
Doloris 50 mg trde kapsule	not available	H/19/02529/001	LENIS FARMACEVTIKA D.O.O.	SI
Dolzam 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE168585	ZAMBON NV	BE
Dolzam 100 mg/ml solution buvable en gouttes	not available	BE168585	ZAMBON NV	BE
Dolzam 50 mg capsules, hard	not available	BE168594	ZAMBON NV	BE
Dolzam 50 mg gélules	not available	BE168594	ZAMBON NV	BE
Dolzam 50 mg/ml	not available	BE168603	ZAMBON 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
ampoules, solution injectable				
Dolzam 50 mg/ml ampullen, oplossing voor injectie	not available	BE168603	ZAMBON NV	BE
Dolzam Retard 100 mg comprimés à libération prolongée	UK/H/0330/002	BE232434	ZAMBON NV	BE
Dolzam Retard 100 mg comprimés à libération prolongée	UK/H/0330/002	BE370194	ZAMBON NV	BE
Dolzam Retard 100 mg comprimés à libération prolongée	UK/H/0330/002	2011071217	ZAMBON NV	LU
Dolzam Retard 100 mg tabletten met verlengde afgifte	UK/H/0330/002	BE232434	ZAMBON NV	BE
Dolzam Retard 100 mg tabletten met verlengde afgifte	UK/H/0330/002	BE370194	ZAMBON NV	BE
Dolzam Retard 150 mg comprimés à libération prolongée	UK/H/0330/003	BE232443	ZAMBON NV	BE
Dolzam Retard 150 mg comprimés à libération prolongée	UK/H/0330/003	BE370203	ZAMBON NV	BE
Dolzam Retard 150 mg comprimés à libération prolongée	UK/H/0330/003	2010120898	ZAMBON NV	LU
Dolzam Retard 150 mg tabletten met verlengde afgifte	UK/H/0330/003	BE232443	ZAMBON NV	BE
Dolzam Retard 150 mg tabletten met verlengde	UK/H/0330/003	BE370203	ZAMBON 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
afgifte				
Dolzam Retard 200 mg comprimés à libération prolongée	UK/H/0330/004	BE232452	ZAMBON NV	BE
Dolzam Retard 200 mg comprimés à libération prolongée	UK/H/0330/004	BE370212	ZAMBON NV	BE
Dolzam Retard 200 mg comprimés à libération prolongée	UK/H/0330/004	2010120899	ZAMBON NV	LU
Dolzam Retard 200 mg tabletten met verlengde afgifte	UK/H/0330/004	BE232452	ZAMBON NV	BE
Dolzam Retard 200 mg tabletten met verlengde afgifte	UK/H/0330/004	BE370212	ZAMBON NV	BE
Dolzam Retard 75 mg comprimés à libération prolongée	UK/H/0330/001	BE232425	ZAMBON NV	BE
Dolzam Retard 75 mg comprimés à libération prolongée	UK/H/0330/001	BE370185	ZAMBON NV	BE
Dolzam Retard 75 mg comprimés à libération prolongée	UK/H/0330/001	2010120896	ZAMBON NV	LU
Dolzam Retard 75 mg tabletten met verlengde afgifte	UK/H/0330/001	BE232425	ZAMBON NV	BE
Dolzam Retard 75 mg tabletten met verlengde afgifte	UK/H/0330/001	BE370185	ZAMBON NV	BE
FORTRADOL 100 mg compresse a rilascio prolungato	not available	028878092	ALFASIGMA 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
FORTRADOL 100 mg/2 ml soluzione iniettabile	not available	028878128	ALFASIGMA S.P.A.	IT
FORTRADOL 100 mg/ml gocce orali soluzione con contagocce	not available	028878080	ALFASIGMA S.P.A.	IT
FORTRADOL 150 mg compresse a rilascio prolungato	not available	028878142	ALFASIGMA S.P.A.	IT
FORTRADOL 200 mg compresse a rilascio prolungato	not available	028878155	ALFASIGMA S.P.A.	IT
FORTRADOL 50 mg capsule rigide	not available	028878078	ALFASIGMA S.P.A.	IT
FORTRADOL 50 mg/ml soluzione iniettabile	not available	028878116	ALFASIGMA S.P.A.	IT
GELOTRADOL 100 mg cápsulas duras de liberación prolongada	DE/H/0639/002	68.501	FERRER INTERNACIONAL, S.A.	ES
GELOTRADOL 150 mg cápsulas duras de liberación prolongada	DE/H/0639/003	68.502	FERRER INTERNACIONAL, S.A.	ES
GELOTRADOL 200 mg cápsulas duras de liberación prolongada	DE/H/0639/004	68503	FERRER INTERNACIONAL, S.A.	ES
GELOTRADOL 50 mg cápsulas duras de liberación prolongada	DE/H/0639/001	68.500	FERRER INTERNACIONAL, S.A.	ES
GELOTRALIB 100 mg cápsulas de libertação prolongada	DE/H/0639/002	5010731	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 100 mg cápsulas de libertação prolongada	DE/H/0639/002	5010749	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 100 mg	DE/H/0639/002	5010756	FERRER INTERNACIONAL,	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
cápsulas de libertação prolongada			S.A.	
GELOTRALIB 150 mg cápsulas de libertação prolongada	DE/H/0639/003	5010764	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 150 mg cápsulas de libertação prolongada	DE/H/0639/003	5010772	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 150 mg cápsulas de libertação prolongada	DE/H/0639/003	5010806	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 200 mg cápsulas de libertação prolongada	DE/H/0639/004	5010814	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 200 mg cápsulas de libertação prolongada	DE/H/0639/004	5010822	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 200 mg cápsulas de libertação prolongada	DE/H/0639/004	5010830	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 50 mg cápsulas de libertação prolongada	DE/H/0639/001	5010707	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 50 mg cápsulas de libertação prolongada	DE/H/0639/001	5010715	FERRER INTERNACIONAL, S.A.	PT
GELOTRALIB 50 mg cápsulas de libertação prolongada	DE/H/0639/001	5010723	FERRER INTERNACIONAL, S.A.	PT
Gemadol 100 mg depotkapsel, hård	UK/H/0225/002	14064	MEDA AB	SE
Gemadol 150 mg depotkapsel, hård	UK/H/0225/003	14065	MEDA AB	SE
Gemadol 200 mg	UK/H/0225/004	14066	MEDA 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
depotkapsel, hård				
Gemadol 50 mg depotkapsel, hård	UK/H/0225/001	14063	MEDA AB	SE
Gemadol Retard	DE/H/5587/002	19326	MYLAN DENMARK APS	DK
Gemadol Retard	DE/H/5587/003	19327	MYLAN DENMARK APS	DK
Gemadol Retard	DE/H/5587/001	19325	MYLAN DENMARK APS	DK
Gemadol Retard	DE/H/5587/004	19328	MYLAN DENMARK APS	DK
Lanalget retard 100 mg-Filmtabletten	AT/H/0117/001	1–23123	G.L. PHARMA GMBH	AT
Lanalget retard 150 mg-Filmtabletten	AT/H/0117/002	1–24030	G.L. PHARMA GMBH	AT
Lanalget retard 200 mg-Filmtabletten	AT/H/0117/003	1–24031	G.L. PHARMA GMBH	AT
Lanalget retard 200 mg-Filmtabletten	AT/H/0117/003	1–24031	G.L. PHARMA GMBH	AT
Lanalget retard 200 mg-Filmtabletten	AT/H/0117/003	1–24031	G.L. PHARMA GMBH	AT
Lanalget retard 200 mg-Filmtabletten	AT/H/0117/003	1–24031	G.L. PHARMA GMBH	AT
Larapam 100mg SR Tablets	NL/H/0483/001	PL 04416/0596	SANDOZ LTD	UK
Larapam 150mg SR Tablets	NL/H/0483/002	PL 04416/0597	SANDOZ LTD	UK
Larapam 200mg SR Tablets	NL/H/0483/003	PL 04416/0598	SANDOZ LTD	UK
LUMIDOL 100 mg otopina za injekciju/infuziju	not available	HR-H-521675660	BELUPO D.D.	HR
LUMIDOL 100 mg/ml oralne kapi, otopina	not available	HR-H-513243658	BELUPO D.D.	HR
LUMIDOL 50 mg otopina za injekciju/infuziju	not available	HR-H-433645088	BELUPO D.D.	HR
LUMIDOL 50 mg tvrde kapsule	not available	HR-H-231277570	BELUPO D.D.	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
Lumidol retard 100 mg tablete s produljenim oslobađanjem	not available	HR-H-864156880	BELUPO D.D.	HR
Lumidol retard 100 mg tablete s produljenim oslobađanjem	not available	HR-H-864156880	BELUPO D.D.	HR
Lumidol retard 200 mg tablete s produljenim oslobađanjem	not available	HR-H-947867682	BELUPO D.D.	HR
MABRON 100 mg/2 ml soluţie injectabilă	not available	12343/2019/01	MEDOCHEMIE LTD.	RO
MABRON 100 mg/2 ml soluţie injectabilă	not available	12343/2019/02	MEDOCHEMIE LTD.	RO
MABRON 100 mg/2 ml soluţie injectabilă	not available	12343/2019/03	MEDOCHEMIE LTD.	RO
MABRON 100 mg/2 ml, injekcní roztok	not available	65/788/94-C	MEDOCHEMIE LTD.	CZ
MABRON 100 mg/2 ml, injekcní roztok	not available	65/788/94-C	MEDOCHEMIE LTD.	CZ
MABRON 100 mg/2 ml, injekcný roztok	not available	65/0459/94-S	MEDOCHEMIE LTD.	SK
Mabron 100mg Prolonged Release Tablets	NL/H/0538/001	PL 20117/0003	MORNINGSIDE HEALTHCARE LTD	UK
Mabron 100mg/2ml solution for injection/infusion	not available	13185	MEDOCHEMIE LTD.	CY
Mabron 150mg Prolonged Release Tablets	NL/H/0538/002	PL 20117/0004	MORNINGSIDE HEALTHCARE LTD	UK
Mabron 200mg Prolonged Release Tablets	NL/H/0538/003	PL 20117/0005	MORNINGSIDE HEALTHCARE LTD	UK
MABRON 50 mg capsule	not available	8728/2016/01	MEDOCHEMIE LTD.	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
MABRON 50 mg capsule	not available	8728/2016/02	MEDOCHEMIE LTD.	RO
MABRON 50 mg capsule	not available	8728/2016/03	MEDOCHEMIE LTD.	RO
Mabron 50 mg capsules	not available	12119	MEDOCHEMIE LTD.	CY
Mabron 50 mg cietās kapsulas	not available	02-0247	MEDOCHEMIE LTD.	LV
MABRON 50 mg tvrdé kapsuly	not available	65/0076/93-S	MEDOCHEMIE LTD.	SK
MABRON 50 mg tvrdé tobolky	not available	65/1000/93-C	MEDOCHEMIE LTD.	CZ
MABRON 50 mg tvrdé tobolky	not available	65/1000/93-C	MEDOCHEMIE LTD.	CZ
MABRON 50 mg/ml injekcinis ar infuzinis tirpalas	not available	LT/1/98/3123/001	MEDOCHEMIE LTD.	LT
Mabron 50 mg/ml šķīdums injekcijām vai infūzijām	not available	02-0248	MEDOCHEMIE LTD.	LV
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/01	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/02	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/03	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/04	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/05	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare	not available	12340/2019/06	MEDOCHEMIE LTD.	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
prelungită				
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/07	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/08	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/09	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/10	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/11	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/12	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/13	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/14	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/15	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/16	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/17	MEDOCHEMIE LTD.	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
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/18	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/19	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/20	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/21	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/22	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/23	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/24	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/25	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/26	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/27	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/28	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg	not available	12340/2019/29	MEDOCHEMIE LTD.	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
comprimate cu eliberare prelungită				
Mabron retard 100 mg comprimate cu eliberare prelungită	not available	12340/2019/30	MEDOCHEMIE LTD.	RO
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/001	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/002	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/003	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/004	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/005	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/006	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/007	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/008	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/009	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto	NL/H/0889/001	LT/1/07/0754/010	MEDOCHEMIE LTD.	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
atpalaidavimo tabletes				
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/011	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/012	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/013	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/014	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/015	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/016	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/017	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/018	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/019	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/020	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/021	MEDOCHEMIE LTD.	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
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/022	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/023	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/024	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/025	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/026	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/027	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/028	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/029	MEDOCHEMIE LTD.	LT
MABRON RETARD 100 mg pailginto atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/030	MEDOCHEMIE LTD.	LT
Mabron Retard 100 mg prolonged-release tablets	NL/H/0889/001	MA032/06701	MEDOCHEMIE LTD.	MT
MABRON RETARD 100 mg tablety s predĺženým uvoľňovaním	NL/H/0538/001	65/0275/05-S	MEDOCHEMIE LTD.	SK
MABRON RETARD 100 mg tablety s	NL/H/0538/001	65/369/05-C	MEDOCHEMIE LTD.	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
prodlouženým uvolňováním				
Mabron retard 100 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/001	20240	MEDOCHEMIE LTD.	CY
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/01	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/02	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/03	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/04	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/05	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/06	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/07	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/08	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/09	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare	not available	12341/2019/10	MEDOCHEMIE LTD.	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
prelungită				
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/11	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/12	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/13	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/14	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/15	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/16	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/17	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/18	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/19	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/20	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/21	MEDOCHEMIE LTD.	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
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/22	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/23	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/24	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/25	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/26	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/27	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/28	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/29	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungită	not available	12341/2019/30	MEDOCHEMIE LTD.	RO
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/031	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/032	MEDOCHEMIE LTD.	LT
MABRON RETARD 150	NL/H/0889/002	LT/1/07/0754/033	MEDOCHEMIE LTD.	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
mg pailginto atpalaidavimo tabletes				
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/034	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/035	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/036	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/037	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/038	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/039	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/040	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/041	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/042	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/043	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto	NL/H/0889/002	LT/1/07/0754/044	MEDOCHEMIE LTD.	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
atpalaidavimo tabletes				
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/045	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/046	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/047	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/048	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/049	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/050	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/051	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/052	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/053	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/055	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/056	MEDOCHEMIE LTD.	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
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/057	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/058	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/059	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/060	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletės	NL/H/0889/002	LT/1/07/0754/054	MEDOCHEMIE LTD.	LT
Mabron Retard 150 mg prolonged-release tablets	NL/H/0889/002	MA032/06702	MEDOCHEMIE LTD.	MT
MABRON RETARD 150 mg tablety s predĺženým uvoľňovaním	NL/H/0538/002	65/0285/05-S	MEDOCHEMIE LTD.	SK
MABRON RETARD 150 mg tablety s prodlouženým uvolňováním	NL/H/0538/002	65/370/05-C	MEDOCHEMIE LTD.	CZ
Mabron retard 150 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/002	20241	MEDOCHEMIE LTD.	CY
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/01	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/02	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg	not available	12342/2019/03	MEDOCHEMIE LTD.	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
comprimate cu eliberare prelungită				
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/04	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/05	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/06	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/07	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/08	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/09	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/10	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/11	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/12	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/13	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare	not available	12342/2019/15	MEDOCHEMIE LTD.	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
prelungită				
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/16	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/17	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/18	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/19	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/14	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/20	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/21	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/22	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/23	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/24	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/25	MEDOCHEMIE LTD.	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
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/26	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/27	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/28	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/29	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungită	not available	12342/2019/30	MEDOCHEMIE LTD.	RO
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/062	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/064	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/065	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/066	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/067	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/068	MEDOCHEMIE LTD.	LT
MABRON RETARD 200	NL/H/0889/003	LT/1/07/0754/069	MEDOCHEMIE LTD.	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
mg pailginto atpalaidavimo tabletes				
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/070	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/071	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/072	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/073	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/074	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/075	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/076	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/077	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/078	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/079	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto	NL/H/0889/003	LT/1/07/0754/080	MEDOCHEMIE LTD.	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
atpalaidavimo tabletes				
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/081	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/082	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/083	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/084	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/085	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/086	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/087	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/088	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/089	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/090	MEDOCHEMIE LTD.	LT
MABRON RETARD 200 mg pailginto atpalaidavimo tabletės	NL/H/0889/003	LT/1/07/0754/061	MEDOCHEMIE LTD.	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
MABRON RETARD 200 mg pailginto atpalaidavimo tabletės	NL/H/0889/003	LT/1/07/0754/063	MEDOCHEMIE LTD.	LT
Mabron Retard 200 mg prolonged-release tablets	NL/H/0889/003	MA032/06703	MEDOCHEMIE LTD.	MT
MABRON RETARD 200 mg tablety s predĺženým uvoľňovaním	NL/H/0538/003	65/0286/05-S	MEDOCHEMIE LTD.	SK
MABRON RETARD 200 mg tablety s prodlouženým uvolňováním	NL/H/0538/003	65/371/05-C	MEDOCHEMIE LTD.	CZ
Mabron retard 200 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/003	20239	MEDOCHEMIE LTD.	CY
Mabron retard, 100 mg toimeainet prolongeeritult vabastavad tabletid	NL/H/0538/001	493405	MEDOCHEMIE LTD.	EE
Mabron retard, 150 mg toimeainet prolongeeritult vabastavad tabletid	NL/H/0538/002	493605	MEDOCHEMIE LTD.	EE
Mabron retard, 200 mg toimeainet prolongeeritult vabastavad tabletid	NL/H/0538/003	493505	MEDOCHEMIE LTD.	EE
MABRON SR 100 mg ilgstošās darbības tabletes	NL/H/0538/001	05-0500	MEDOCHEMIE LTD.	LV
MABRON SR 150 mg ilgstošās darbības tabletes	NL/H/0538/002	05-0501	MEDOCHEMIE LTD.	LV
MABRON SR 200 mg	NL/H/0538/003	05-0502	MEDOCHEMIE LTD.	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
ilgstošās darbības tabletes				
Mabron, 50 mg/ml süste- või infusioonilahus	not available	340701	MEDOCHEMIE LTD.	EE
Mandolgin	not available	19352	SANDOZ A/S	DK
Mandolgin	not available	18666	SANDOZ A/S	DK
Mandolgin	not available	18665	SANDOZ A/S	DK
Mandolgin Retard	not available	19424	SANDOZ A/S	DK
Mandolgin Retard	not available	19426	SANDOZ A/S	DK
Mandolgin Retard	not available	19425	SANDOZ A/S	DK
Maneo 100 mg Prolonged-release tablets.	DK/H/2942/001	PL 04569/1406	GENERICS [UK] LIMITED	UK
Maneo 100 mg Prolonged-release tablets.	DK/H/2942/001	PL 04569/1406	GENERICS [UK] LIMITED	UK
Maneo 100 mg Prolonged-release tablets.	DK/H/2942/001	PL 04569/1406	GENERICS [UK] LIMITED	UK
Maneo 100 mg Prolonged-release tablets.	DK/H/2942/001	PL 04569/1406	GENERICS [UK] LIMITED	UK
Maneo 150 mg Prolonged-release tablets.	DK/H/2942/002	PL 04569/1407	GENERICS [UK] LIMITED	UK
Maneo 150 mg Prolonged-release tablets.	DK/H/2942/002	PL 04569/1407	GENERICS [UK] LIMITED	UK
Maneo 150 mg Prolonged-release tablets.	DK/H/2942/002	PL 04569/1407	GENERICS [UK] LIMITED	UK
Maneo 150 mg Prolonged-release tablets.	DK/H/2942/002	PL 04569/1407	GENERICS [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
Maneo 200 mg Prolonged-release tablets.	DK/H/2942/003	PL 04569/1408	GENERICS [UK] LIMITED	UK
Maneo 200 mg Prolonged-release tablets.	DK/H/2942/003	PL 04569/1408	GENERICS [UK] LIMITED	UK
Maneo 200 mg Prolonged-release tablets.	DK/H/2942/003	PL 04569/1408	GENERICS [UK] LIMITED	UK
Maneo 200 mg Prolonged-release tablets.	DK/H/2942/003	PL 04569/1408	GENERICS [UK] LIMITED	UK
Marol 100mg Prolonged-release tablets	not available	PL 20117/0045	MORNINGSIDE HEALTHCARE LTD	UK
Marol 150mg Prolonged-release tablets	not available	PL 20117/0046	MORNINGSIDE HEALTHCARE LTD	UK
Marol 200mg Prolonged-release tablets	not available	PL 20117/0047	MORNINGSIDE HEALTHCARE LTD	UK
MAXITRAM SR 100 mg prolonged-release capsule, hard	DE/H/0639/002	PL 06934/0235	ETHYPHARM	UK
MAXITRAM SR 150 mg prolonged-release capsule, hard	DE/H/0639/003	PL 06934/0236	ETHYPHARM	UK
MAXITRAM SR 200 mg prolonged-release capsule, hard	DE/H/0639/004	PL 06934/0237	ETHYPHARM	UK
MAXITRAM SR 50 mg prolonged-release capsule, hard	DE/H/0639/001	PL 06934/0234	ETHYPHARM	UK
Metamizole Kalceks 500 mg/ml solution for injection	CZ/H/0701/001	20200069	KALCEKS	BG
MONOCRIXO L.P. 100	FI/H/0164/001	34009 362 470 6 2	THERABEL LUCIEN PHARMA	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
mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)			S.A.	
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 471 2 3	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 472 9 1	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 473 5 2	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 474 1 3	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 475 8 1	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 476 4 2	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 477 0 3	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE	FI/H/0164/002	34009 362 478 7 1	THERABEL LUCIEN PHARMA S.A.	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
QUOTIDIENNE)				
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 479 3 2	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 200 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/003	34009 362 480 1 4	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 200 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/003	34009 362 481 8 2	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 200 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/003	34009 362 482 4 3	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 200 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/003	34009 362 483 0 4	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 200 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/003	34009 362 484 7 2	THERABEL LUCIEN PHARMA S.A.	FR
Nobligan 100 mg/ml orala droppar, lösning	not available	14305	GRÜNENTHAL GMBH	SE
Nobligan 50 mg kapslar, hårda	not available	12845	GRÜNENTHAL GMBH	SE
Nobligan 50 mg kapsler, harde	not available	00-8268	GRÜNENTHAL GMBH	NO
Nobligan Retard 100 mg depottabletter	not available	98-2970	GRÜNENTHAL GMBH	NO
Nobligan retard 100 mg	DE/H/0108/001	13287	GRÜNENTHAL GMBH	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
depottabletter				
Nobligan retard 100 mg Retardtabletten	DE/H/0136/001	37291.00.00	GRÜNENTHAL GMBH	DE
Nobligan Retard 150 mg depottabletter	not available	98-2971	GRÜNENTHAL GMBH	NO
Nobligan retard 150 mg depottabletter	DE/H/0108/002	13288	GRÜNENTHAL GMBH	SE
Nobligan retard 150 mg Retardtabletten	DE/H/0136/002	37291.01.00	GRÜNENTHAL GMBH	DE
Nobligan Retard 200 mg depottabletter	not available	98-2972	GRÜNENTHAL GMBH	NO
Nobligan retard 200 mg depottabletter	DE/H/0108/003	13289	GRÜNENTHAL GMBH	SE
Nobligan retard 200 mg, Retardtabletten	DE/H/0136/003	37291.02.00	GRÜNENTHAL GMBH	DE
Nobligan retard 50 mg Retardtabletten	DE/H/0136/004	60445.00.00	GRÜNENTHAL GMBH	DE
Nobligan Retard, depottabletter 100 mg	DE/H/0108/001	18540	GRÜNENTHAL GMBH	DK
Nobligan Retard, depottabletter 150 mg	DE/H/0108/002	18541	GRÜNENTHAL GMBH	DK
Nobligan Retard, depottabletter 200 mg	DE/H/0108/003	18542	GRÜNENTHAL GMBH	DK
Nobligan, kapsler, hårde	not available	14623	GRÜNENTHAL GMBH	DK
Nobligan, orale dråber, opløsning	not available	14624	GRÜNENTHAL GMBH	DK
Olteron, 100 mg toimeainet prolongeeritult vabastavad tabletid	EE/H/0262/001	976318	KRKA, D.D., NOVO MESTO	EE
Olteron, 150 mg toimeainet prolongeeritult vabastavad tabletid	EE/H/0262/002	976418	KRKA, D.D., NOVO MESTO	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
Olteron, 200 mg toimeainet prolongeeritult vabastavad tabletid	EE/H/0262/003	976518	KRKA, D.D., NOVO MESTO	EE
ORATRAM 100 100 mg, tabletki o przedluzonym uwalnianiu	NL/H/0888/001	14313	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
ORATRAM 150 150 mg, tabletki o przedłużonym uwalnianiu	NL/H/0888/002	14315	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
ORATRAM 200 200 mg, tabletki o przedłużonym uwalnianiu	NL/H/0888/003	14314	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
PAXILFAR 100 mg/2 ml solução injetável	not available	9515411	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
PAXILFAR 100 mg/2 ml solução injetável	not available	9515445	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
PAXILFAR 100 mg/2 ml solução injetável	not available	9515457	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
Paxilfar 100mg tablets	not available	6/13/84	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
Poltram 100, 50 mg/ml, roztwór do wstrzykiwań	not available	9689	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Poltram 50, 50 mg/ml, roztwór do wstrzykiwań	not available	9688	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Poltram Retard 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	9676	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	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
Poltram Retard 150, 150 mg, tabletki o przedłużonym uwalnianiu	not available	9677	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Poltram Retard 200, 200 mg, tabletki o przedłużonym uwalnianiu	not available	9678	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Poltram, 100 mg/ml, krople doustne, roztwór	not available	9690	MEDANA PHARMA SPOLKA AKCYJNA	PL
Poltram, 50 mg, kapsułki	not available	9687	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074042	NEOPHARMED GENTILI SPA	IT
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074067	NEOPHARMED GENTILI SPA	IT
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074055	NEOPHARMED GENTILI SPA	IT
PRONTALGIN 50 mg/ml soluzione iniettabile	NL/H/0113/001	033074028	NEOPHARMED GENTILI SPA	IT
PRONTALGIN Capsule, 50 mg capsule rigide.	NL/H/0113/002	033074030	NEOPHARMED GENTILI SPA	IT
PRONTALGIN Gocce, 100 mg/ml gocce orali, soluzione.	NL/H/0113/003	033074016	NEOPHARMED GENTILI SPA	IT
Ralgen 50 mg kemény kapszula	not available	OGYI-T-20310/15	ZENTIVA, K.S.	HU
Ralgen 50 mg kemény kapszula	not available	OGYI-T-20310/14	ZENTIVA, K.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/05	ZENTIVA, K.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/04	ZENTIVA, K.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/03	ZENTIVA, K.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
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/02	ZENTIVA, K.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/09	ZENTIVA, K.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/08	ZENTIVA, K.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/06	ZENTIVA, K.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/07	ZENTIVA, K.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/13	ZENTIVA, K.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/11	ZENTIVA, K.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/12	ZENTIVA, K.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/10	ZENTIVA, K.S.	HU
Rofy 100mg/2ml, solution for injection or infusion	not available	20016	CODAL SYNTO LTD	CY
Rofy 50 mg capsules	not available	20017	CODAL SYNTO LTD	CY
Rofy retard 100 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0888/001	20236	CODAL SYNTO LTD	CY
Rofy retard 150 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0888/002	20237	CODAL SYNTO LTD	CY
Rofy retard 200 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0888/003	20238	CODAL SYNTO LTD	CY
Tadol 100 mg svečke	not available	H/93/01481/005	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg tablete s podaljšanim sproščanjem	not available	H/93/01481/001	KRKA, D.D., NOVO MESTO	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
Tadol 100 mg/ml peroralne kapljice, raztopina	not available	H/93/01481/006	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg/ml peroralne kapljice, raztopina	not available	H/93/01481/007	KRKA, D.D., NOVO MESTO	SI
Tadol 150 mg tablete s podaljšanim sproščanjem	not available	H/93/01481/002	KRKA, D.D., NOVO MESTO	SI
Tadol 200 mg tablete s podaljšanim sproščanjem	not available	H/93/01481/003	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg trde kapsule	not available	H/93/01481/004	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	not available	H/93/01481/010	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	not available	H/93/01481/011	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	not available	H/93/01481/008	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	not available	H/93/01481/009	KRKA, D.D., NOVO MESTO	SI
Tadol, kapsler, hårde	DE/H/0282/001	32972	STADA ARZNEIMITTEL AG	DK
Tadol, suppositorier	DE/H/0282/003	32974	STADA ARZNEIMITTEL AG	DK
TAKADOL 100 mg, comprimé effervescent sécable	not available	34009 355 316 5 0	LABORATOIRES EXPANSCIENCE	FR
THERADOL 50 mg Bruistabletten	NL/H/0113/005	RVG24573	THERABEL PHARMA N.V.	NL
THERADOL Capsules, capsules 50 mg, hard	NL/H/0113/002	RVG 17779	THERABEL PHARMA N.V.	NL
THERADOL Druppels,	NL/H/0113/003	RVG 17780	THERABEL PHARMA N.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
druppels 100 mg/ml				
THERADOL Injectie, oplossing voor injectie 50 mglml	NL/H/0113/001	RVG 17778	THERABEL PHARMA N.V.	NL
Tilodol SR 100 mg Prolonged-Release Tablets	DE/H/0448/001	PL 04416/1237	SANDOZ LTD	UK
Tilodol SR 150 mg Prolonged-Release Tablets	DE/H/0448/002	PL 04416/1238	SANDOZ LTD	UK
Tilodol SR 200 mg Prolonged-Release Tablets	DE/H/0448/003	PL 04416/1239	SANDOZ LTD	UK
Tioner 100 mg/ml solución oral	not available	62015	ARISTO PHARMA IBERIA, S.L.	ES
Tioner 50 mg cápsulas duras.	not available	62.016	ARISTO PHARMA IBERIA, S.L.	ES
Tioner retard 100 mg comprimidos de liberación prolongada	not available	62919	ARISTO PHARMA IBERIA, S.L.	ES
Tioner retard 150 mg comprimidos de liberación prolongada	not available	62920	ARISTO PHARMA IBERIA, S.L.	ES
Tioner retard 200 mg comprimidos de liberación prolongada	not available	62921	ARISTO PHARMA IBERIA, S.L.	ES
Tiparol 50 mg brustablett	not available	14106	BLUEFISH PHARMACEUTICALS AB	SE
Tiparol 50 mg brustablett	not available	14106	BLUEFISH PHARMACEUTICALS AB	SE
T-long ® 150 mg Retardkapseln Hartkapsel, retardiert Wirkstoff:	FI/H/0164/002	54607.01.00	LABORATOIRES SMB S.A.	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
Tramadolhydrochlorid				
T-long ® 200 mg Retardkapseln Hartkapsel, retardiert Wirkstoff: Tramadolhydrochlorid	FI/H/0164/003	54607.02.00	LABORATOIRES SMB S.A.	DE
T-long® 100 mg Retardkapseln Hartkapsel, retardiert Wirkstoff: Tramadolhydrochlorid	FI/H/0164/001	54607.00.00	LABORATOIRES SMB S.A.	DE
Toram 50 mg tvrde kapsule	not available	HR-H-151686235	MIBE PHARMACEUTICALS D.O.O.	HR
TRADOGUT 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875020	MEDA PHARMA S.P.A.	IT
TRADOGUT 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875018	MEDA PHARMA S.P.A.	IT
TRADOGUT 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875032	MEDA PHARMA S.P.A.	IT
TRADOGUT 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875044	MEDA PHARMA S.P.A.	IT
TRADOGUT 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875057	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg capsule rigide	DE/H/0306/001	035875095	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg capsule rigide	DE/H/0306/001	035875107	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg capsule rigide	DE/H/0306/001	035875121	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg capsule rigide	DE/H/0306/001	035875119	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg/ml soluzione iniettabile	DE/H/0306/002	035875071	MEDA PHARMA S.P.A.	IT
TRADOGUT 50 mg/ml	DE/H/0306/002	035875069	MEDA PHARMA 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
soluzione iniettabile				
TRADOGUT 50 mg/ml soluzione iniettabile	DE/H/0306/002	035875083	MEDA PHARMA S.P.A.	IT
Tradol 50 mg Effervescent Tablets	not available	PA0711/029/002	ROWEX LTD	IE
Tradol 50mg Hard Capsules	not available	PA0711/029/001	ROWEX LTD	IE
Tradol 50mg/ml Solution for Injection or Infusion	not available	PA0711/029/003	ROWEX LTD	IE
Tradol SR 100 mg Prolonged Release Tablets	not available	PA 0711/029/005	ROWEX LTD	IE
Tradol SR 150 mg Prolonged Release Tablets	not available	PA 0711/029/006	ROWEX LTD	IE
Tradol SR 200 mg Prolonged Release Tablets	not available	PA 0711/029/007	ROWEX LTD	IE
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan	not available	18233	G.L. PHARMA GMBH	DK
Tradolan 100 mg-Ampullen	not available	1-21804	G.L. PHARMA GMBH	AT
Tradolan 100 mg-Zäpfchen	not available	1-21810	G.L. PHARMA GMBH	AT
Tradolan 50 mg filmdragerade tabletter	not available	13068	G.L. PHARMA GMBH	SE
Tradolan 50 mg filmuhúðaðar töflur	not available	960128	G.L. PHARMA GMBH	IS
Tradolan 50 mg filmuhúðaðar töflur	not available	960128	G.L. PHARMA GMBH	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
Tradolan 50 mg filmuhúðaðar töflur	not available	960128	G.L. PHARMA GMBH	IS
Tradolan 50 mg filmuhúðaðar töflur	not available	960128	G.L. PHARMA GMBH	IS
Tradolan 50 mg filmuhúðaðar töflur	not available	960128	G.L. PHARMA GMBH	IS
Tradolan 50 mg/ml injektionsvätska, lösning	not available	13067	G.L. PHARMA GMBH	SE
Tradolan 50 mg/ml injektionsvätska, lösning	not available	13067	G.L. PHARMA GMBH	SE
Tradolan 50 mg/ml injektionsvätska, lösning	not available	13067	G.L. PHARMA GMBH	SE
Tradolan 50 mg-Ampullen	not available	1-21802	G.L. PHARMA GMBH	AT
Tradolan 50 mg-Filmtabletten	not available	1-21806	G.L. PHARMA GMBH	AT
Tradolan Retard	AT/H/0117/003	32518	G.L. PHARMA GMBH	DK
Tradolan Retard	AT/H/0117/001	32516	G.L. PHARMA GMBH	DK
Tradolan Retard	AT/H/0117/002	32517	G.L. PHARMA GMBH	DK
Tradolan Retard	AT/H/0117/003	32518	G.L. PHARMA GMBH	DK
Tradolan Retard	AT/H/0117/003	32518	G.L. PHARMA GMBH	DK
Tradolan Retard	AT/H/0117/003	32518	G.L. PHARMA GMBH	DK
Tradolan Retard 100 mg depottabletter	AT/H/0117/001	17256	G.L. PHARMA GMBH	SE
Tradolan Retard 100 mg depottabletti	AT/H/0117/001	16503	G.L. PHARMA GMBH	FI
Tradolan retard 100 mg-Filmtabletten	not available	1-21255	G.L. PHARMA GMBH	AT
Tradolan Retard 150 mg comprimate cu eliberare prelungită	not available	12191/2019/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Tradolan Retard 150 mg comprimate cu eliberare prelungită	not available	12191/2019/02	LANNACHER HEILMITTEL GES.M.B.H.,	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
Tradolan Retard 150 mg depottabletter	AT/H/0117/002	17257	G.L. PHARMA GMBH	SE
Tradolan Retard 150 mg depottabletti	AT/H/0117/002	16504	G.L. PHARMA GMBH	FI
Tradolan retard 150 mg-Filmtabletten	not available	1-21254	G.L. PHARMA GMBH	AT
Tradolan Retard 200 mg comprimate cu eliberare prelungită	not available	12192/2019/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Tradolan Retard 200 mg comprimate cu eliberare prelungită	not available	12192/2019/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Tradolan Retard 200 mg depottabletter	AT/H/0117/003	17258	G.L. PHARMA GMBH	SE
Tradolan Retard 200 mg depottabletter	AT/H/0117/003	17258	G.L. PHARMA GMBH	SE
Tradolan Retard 200 mg depottabletter	AT/H/0117/003	17258	G.L. PHARMA GMBH	SE
Tradolan Retard 200 mg depottabletter	AT/H/0117/003	17258	G.L. PHARMA GMBH	SE
Tradolan Retard 200 mg depottabletti	AT/H/0117/003	16505	G.L. PHARMA GMBH	FI
Tradolan Retard 200 mg depottabletti	AT/H/0117/003	16505	G.L. PHARMA GMBH	FI
Tradolan Retard 200 mg depottabletti	AT/H/0117/003	16505	G.L. PHARMA GMBH	FI
Tradolan Retard 200 mg depottabletti	AT/H/0117/003	16505	G.L. PHARMA GMBH	FI
Tradolan retard 200 mg-Filmtabletten	not available	1-21239	G.L. PHARMA GMBH	AT
Tradolan-Tropfen	not available	1-21808	G.L. PHARMA GMBH	AT
Tradonal retard 100 mg cápsulas duras de liberación prolongada	UK/H/0225/002	62.111	MYLAN IRE HEALTHCARE LIMITED	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
Tradonal retard 150 mg cápsulas duras de liberación prolongada	UK/H/0225/003	62.112	MYLAN IRE HEALTHCARE LIMITED	ES
Tradonal retard 200 mg cápsulas duras de liberación prolongada	UK/H/0225/004	62.113	MYLAN IRE HEALTHCARE LIMITED	ES
TRADONAL Retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	BE195447	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	2011010940	MYLAN EPD BVBA/SPRL	LU
TRADONAL Retard 100 mg, capsules met verlengde afgifte, hard	UK/H/0225/002	BE195447	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 100 mg, gélules à libération prolongée	UK/H/0225/002	BE195447	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 100 mg, gélules à libération prolongée	UK/H/0225/002	2011010940	MYLAN EPD BVBA/SPRL	LU
TRADONAL Retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	BE195456	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	2011010941	MYLAN EPD BVBA/SPRL	LU
TRADONAL Retard 150 mg, capsules met verlengde afgifte, hard	UK/H/0225/003	BE195456	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 150 mg, gélules à libération prolongée	UK/H/0225/003	BE195456	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 150	UK/H/0225/003	2011010941	MYLAN EPD BVBA/SPRL	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
mg, gélules à libération prolongée				
TRADONAL Retard 200 mg Hartkapseln, retardiert	UK/H/0225/004	BE195465	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 200 mg Hartkapseln, retardiert	UK/H/0225/004	2011010942	MYLAN EPD BVBA/SPRL	LU
TRADONAL Retard 200 mg, capsules met verlengde afgifte, hard	UK/H/0225/004	BE195465	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 200 mg, gélules à libération prolongée	UK/H/0225/004	BE195465	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 200 mg, gélules à libération prolongée	UK/H/0225/004	2011010942	MYLAN EPD BVBA/SPRL	LU
Tradonal retard 50 mg cápsulas duras de liberación prolongada.	UK/H/0225/001	62110	MYLAN IRE HEALTHCARE LIMITED	ES
TRADONAL Retard 50 mg Hartkapseln, retardiert	UK/H/0225/001	BE195377	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 50 mg Hartkapseln, retardiert	UK/H/0225/001	2011010939	MYLAN EPD BVBA/SPRL	LU
TRADONAL Retard 50 mg, capsules met verlengde afgifte, hard	UK/H/0225/001	BE195377	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 50 mg, gélules à libération prolongée	UK/H/0225/001	BE195377	MYLAN EPD BVBA/SPRL	BE
TRADONAL Retard 50 mg, gélules à libération prolongée	UK/H/0225/001	2011010939	MYLAN EPD BVBA/SPRL	LU
TRADONAL S.R. 100 mg	UK/H/0225/002	034233027	MEDA PHARMA 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
capsule rigide a rilascio prolungato				
TRADONAL S.R. 100 mg capsule rigide a rilascio prolungato	UK/H/0225/002	034233066	MEDA PHARMA S.P.A.	IT
TRADONAL S.R. 50 mg capsule rigide a rilascio prolungato	UK/H/0225/001	034233054	MEDA PHARMA S.P.A.	IT
TRADONAL S.R. 50 mg capsule rigide a rilascio prolungato	UK/H/0225/001	034233015	MEDA PHARMA S.P.A.	IT
TRADONAL, 100 mg, druppels voor oraal gebruik	not available	BE 177633	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, Injektionslösung	not available	BE 177615	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, Injektionslösung	not available	2003087661	MYLAN EPD BVBA/SPRL	LU
TRADONAL, 100 mg, oplossing voor injectie	not available	BE 177615	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, solution buvable en gouttes	not available	BE 177633	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, solution buvable en gouttes	not available	2003/087560	MYLAN EPD BVBA/SPRL	LU
TRADONAL, 100 mg, solution injectable	not available	BE 177615	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, solution injectable	not available	2003087661	MYLAN EPD BVBA/SPRL	LU
TRADONAL, 100 mg, Tropfen zum Einnehmen	not available	BE 177633	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 100 mg, Tropfen zum Einnehmen	not available	2003/087560	MYLAN EPD BVBA/SPRL	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
TRADONAL, 50 mg, capsules	not available	BE 177642	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 50 mg, gélules	not available	BE 177642	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 50 mg, gélules	not available	2003/087559	MYLAN EPD BVBA/SPRL	LU
TRADONAL, 50 mg, Kapseln	not available	BE 177642	MYLAN EPD BVBA/SPRL	BE
TRADONAL, 50 mg, Kapseln	not available	2003/087559	MYLAN EPD BVBA/SPRL	LU
TRADONAL™ S.R. 150 mg Capsule	UK/H/0225/003	034233039	MEDA PHARMA S.P.A.	IT
TRADONAL™ S.R. 150 mg Capsule	UK/H/0225/003	034233078	MEDA PHARMA S.P.A.	IT
TRADONAL™ S.R. 200 mg Capsule	UK/H/0225/004	034233041	MEDA PHARMA S.P.A.	IT
TRADONAL™ S.R. 200 mg Capsule	UK/H/0225/004	034233080	MEDA PHARMA S.P.A.	IT
TRALGIT 100 inj. injekčný roztok	not available	65/0172/03-S	ZENTIVA, A.S.	SK
Tralgit 100 mg/2 ml injekční roztok	not available	65/028/04-C	ZENTIVA, A.S.	CZ
Tralgit 100 mg/ml perorální roztok	not available	65/195/03-C	ZENTIVA, A.S.	CZ
TRALGIT 50 mg kapsuly	not available	65/0169/03-S	ZENTIVA, A.S.	SK
Tralgit 50 mg tvrdé tobolky	not available	65/105/03-C	ZENTIVA, K.S.	CZ
Tralgit 50 mg/ml injekční roztok	not available	65/027/04-C	ZENTIVA, A.S.	CZ
TRALGIT gtt 100 mg/ml perorálne roztokové kvapky	not available	65/0170/03-S	ZENTIVA, A.S.	SK
Tralgit Orotab 50 mg tablety dispergovatelné v	not available	65/575/10-C	ZENTIVA, K.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
ústech				
TRALGIT Sprint 50 mg orodispergovateľné tablety	not available	65/0417/10-S	ZENTIVA, A.S.	SK
TRALGIT Sprint 50 mg orodispergovateľné tablety	not available	65/0417/10-S	ZENTIVA, A.S.	SK
TRALGIT SR 100 100 mg tablety s predĺženým uvoľňovaním	not available	65/0258/02-S	ZENTIVA, A.S.	SK
Tralgit SR 100 mg tablety s prodlouženým uvolňováním	not available	65/357/01-C	ZENTIVA, K.S.	CZ
TRALGIT SR 150 150 mg tablety s predĺženým uvoľňovaním	not available	65/0259/02-S	ZENTIVA, A.S.	SK
Tralgit SR 150 mg tablety s prodlouženým uvolňováním	not available	65/048/03-C	ZENTIVA, K.S.	CZ
TRALGIT SR 200 200 mg tablety s predĺženým uvoľňovaním	not available	65/0260/02-S	ZENTIVA, A.S.	SK
Tralgit SR 200 mg tablety s prodlouženým uvolňováním	not available	65/049/03-C	ZENTIVA, K.S.	CZ
Tralodie® 100 mg Capsule rigide a rilascio prolungato.	FI/H/0164/001	035986037	NEOPHARMED GENTILI SPA	IT
Tralodie® 100 mg Capsule rigide a rilascio prolungato.	FI/H/0164/001	035986025	NEOPHARMED GENTILI SPA	IT
Tralodie® 100 mg Capsule rigide a rilascio prolungato.	FI/H/0164/001	035986049	NEOPHARMED GENTILI 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
Tralodie® 100 mg Capsule rigide a rilascio prolungato.	FI/H/0164/001	035986013	NEOPHARMED GENTILI SPA	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato.	FI/H/0164/002	035986088	NEOPHARMED GENTILI SPA	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato.	FI/H/0164/002	035986052	NEOPHARMED GENTILI SPA	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato.	FI/H/0164/002	035986064	NEOPHARMED GENTILI SPA	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato.	FI/H/0164/002	035986076	NEOPHARMED GENTILI SPA	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato.	FI/H/0164/003	035986090	NEOPHARMED GENTILI SPA	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato.	FI/H/0164/003	035986114	NEOPHARMED GENTILI SPA	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato.	FI/H/0164/003	035986102	NEOPHARMED GENTILI SPA	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato.	FI/H/0164/003	035986126	NEOPHARMED GENTILI SPA	IT
Tramabene - Tropfen	not available	1-21799	TEVA B.V	AT
Tramabene 100 mg - Ampullen	not available	1-21797	TEVA B.V	AT
Tramabene 100 mg – Retardtabletten	not available	1-23913	TEVA B.V	AT
Tramabene 150 mg Retardtabletten	NL/H/0892/002	1-27076	TEVA B.V	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
Tramabene 200 mg Retardtabletten	NL/H/0892/003	1-27077	TEVA B.V	AT
Tramabene 50 mg - Ampullen	not available	1-21796	TEVA B.V	AT
Tramabene 50 mg - Kapseln	not available	1-21798	TEVA B.V	AT
TRAMABENE kapky	not available	65/155/99-C	RATIOPHARM GMBH	CZ
Tramabene Perorálne kvapky	not available	65/0120/99-S	TEVA B.V	SK
Tramabeta 100 mg/ml; Tropfen zum Einnehmen, Lösung	not available	25528.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 100 mg Retardtabletten	not available	44604.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 150 mg Retardtabletten	not available	44604.01.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 200 mg Retardtabletten	not available	44604.02.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramadex 50 mg Capsules	not available	019207	DELORBIS PHARMACEUTICALS LTD	CY
Tramadin 50 mg hård kapsel	not available	12346	RATIOPHARM GMBH	FI
Tramadin 50 mg kapseli, kova	not available	12346	RATIOPHARM GMBH	FI
Tramadol "Aurobindo", hårde kapsler	NL/H/2480/001	49659	AUROBINDO PHARMA (MALTA) LIMITED	DK
Tramadol "Krka", depottabletter	EE/H/0262/001	MTNR. 60644	KRKA, D.D., NOVO MESTO	DK
Tramadol "Krka", depottabletter	EE/H/0262/002	60645	KRKA, D.D., NOVO MESTO	DK
Tramadol "Krka", depottabletter	EE/H/0262/003	60646	KRKA, D.D., NOVO MESTO	DK
Tramadol "Krka", hårde kapsler	HR/H/0104/003	59094	KRKA, D.D., NOVO MESTO	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
Tramadol "Medical Valley", hårde kapsler	DK/H/2693/001	58385	MEDICAL VALLEY INVEST AB	DK
Tramadol ”Actavis”, kapsler, hårde	SE/H/1774/001	31829	ACTAVIS GROUP PTC EHF.	DK
Tramadol ”Vitabalans”, tabletter	FI/H/0779/001	49370	VITABALANS OY	DK
Tramadol 100 injekt - 1A-Pharma, 100mg/2 ml Injektionslösung	not available	32751.00.00	1 A PHARMA GMBH	DE
Tramadol 100 mg supozitoare	not available	10410/2017/01	KRKA, D.D., NOVO MESTO	RO
Tramadol 100 mg/2 ml ssoluţie injectabilă	not available	10412/2017/01	KRKA, D.D., NOVO MESTO	RO
Tramadol 100 mg/ml oral drops, solution	UK/H/5154/001	PL 12762/0453	MERCURY PHARMACEUTICALS LTD.	UK
Tramadol 100 ret - 1 A Pharma, 100 mg Retardtabletten	not available	49470.00.00	1 A PHARMA GMBH	DE
Tramadol 150 ret - 1 A Pharma, 150 mg Retardtabletten	not available	49470.01.00	1 A PHARMA GMBH	DE
Tramadol 1A Pharma 100 mg/ml - Tropfen	not available	1-24618	1A PHARMA GMBH	AT
Tramadol 1A Pharma 50 mg - Kapseln	not available	1-24617	1A PHARMA GMBH	AT
Tramadol 200 ret - 1 A Pharma, 200 mg Retardtabletten	not available	49470.02.00	1 A PHARMA GMBH	DE
Tramadol 2care4 100 mg kapslar, hårda	not available	55031	2CARE4 GENERICS APS	SE
Tramadol 2care4 50 mg kapslar, hårda	not available	55030	2CARE4 GENERICS APS	SE
Tramadol 50 Kapseln - 1A-Pharma 50 mg	not available	32751.00.02	1 A 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
Hartkapseln				
TRAMADOL 50 mg capsule	not available	10413/2017/01	KRKA, D.D., NOVO MESTO	RO
Tramadol 50 mg Capsules	not available	PL 17907/0110	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Tramadol 50 mg Capsules, Hard	UK/H/6747/001	PL 36722/0129	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Tramadol 50 mg capsules, hard	NL/H/2480/001	PL 16363/0335	MILPHARM LIMITED	UK
Tramadol 50 mg/1 ml soluţie injectabilă	not available	10411/2017/01	KRKA, D.D., NOVO MESTO	RO
Tramadol 50 mg/ml solution for injection/infusion	LV/H/0152/001	PL 47015/0002	KALCEKS	UK
Tramadol 50 tabs - 1 A Pharma, 50 mg Tabletten	not available	30798.00.00	1 A PHARMA GMBH	DE
Tramadol 50mg Capsules	UK/H/0953/001	PA1128/003/001	RELON CHEM LIMITED	IE
Tramadol 50mg Capsules	UK/H/0953/001	PA1128/003/001	RELON CHEM LIMITED	IE
Tramadol 50mg Capsules	not available	PL 00289/1603	TEVA UK LIMITED	UK
Tramadol 50mg Capsules	UK/H/0953/001	PL 20395/0065	RELON CHEM LIMITED	UK
Tramadol 50mg Capsules	UK/H/0953/001	PL 20395/0065	RELON CHEM LIMITED	UK
Tramadol 50mg/ml Solution for Injection or Infusion.	not available	PL 18157/0014	BEACON PHARMACEUTICALS LIMITED	UK
Tramadol AbZ 100 mg Retardkapseln	not available	54390.01.00	ABZ-PHARMA GMBH	DE
Tramadol AbZ 100 mg/ml Tropfen Tropfen zum Einnehmen, Lösung	not available	31518.00.01	ABZ-PHARMA GMBH	DE
Tramadol AbZ 150 mg Retardkapseln	not available	62758.01.00	ABZ-PHARMA GMBH	DE
Tramadol AbZ 200 mg Retardkapseln	not available	62758.02.00	ABZ-PHARMA GMBH	DE
Tramadol actas 100 mg	not available	49469.00.00	ARISTO 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
Retardtabletten			(ART 57)	
Tramadol actas 150 mg Retardtabletten	not available	49469.01.00	ARISTO PHARMA GMBH (ART 57)	DE
Tramadol actas 200 mg Retardtabletten	not available	49469.02.00	ARISTO PHARMA GMBH (ART 57)	DE
Tramadol Actavis 50 mg gélule	FR/H/0403/001	NL 37429	ACTAVIS GROUP PTC EHF.	FR
Tramadol Actavis 50 mg harde kapsler.	FR/H/0403/001	09-6559	ACTAVIS GROUP PTC EHF.	NO
Tramadol Actavis 50 mg hylki, hörð	FR/H/0403/001	IS/1/09/064/01	ACTAVIS GROUP PTC EHF.	IS
Tramadol Actavis 50 mg hylki, hörð	FR/H/0403/001	IS/1/09/064/01	ACTAVIS GROUP PTC EHF.	IS
Tramadol Actavis 50 mg kapslar, hårda	SE/H/1774/001	16600	ACTAVIS GROUP PTC EHF.	SE
Tramadol Actavis 50mg capsules, hard	FR/H/0403/001	09-6559	ACTAVIS GROUP PTC EHF.	NO
Tramadol AL 100 Ampullen	not available	30903.00.00	ALIUD PHARMA GMBH	DE
Tramadol AL 100 mg Retardtabletten	AT/H/0118/001	53804.00.00	ALIUD PHARMA GMBH	DE
Tramadol AL 150 mg Retardtabletten	AT/H/0118/002	53804.01.00	ALIUD PHARMA GMBH	DE
Tramadol AL 200 mg Retardtabletten	AT/H/0118/003	53804.02.00	ALIUD PHARMA GMBH	DE
Tramadol AL 50 Kapseln	not available	30903.00.02	ALIUD PHARMA GMBH	DE
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/03	STADA ARZNEIMITTEL AG	HU
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/02	STADA ARZNEIMITTEL AG	HU
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/01	STADA ARZNEIMITTEL AG	HU
Tramadol AL Tropfen Tramadolhydrochlorid	not available	30903.00.01	ALIUD 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
100 mg/ml Lösung zum Einnehmen				
Tramadol Almus 50 mg cápsulas duras EFG	ES/H/0700/001	83921	ALMUS FARMACEUTICA S.A	ES
Tramadol Alvogen 100 mg/ml belsőleges oldatos cseppek	not available	OGYI-T-7724/01	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Alvogen 50 mg kemény kapszula	not available	OGYI-T-7724/06	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Alvogen 50 mg kemény kapszula	not available	OGYI-T-7724/07	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Alvogen 50 mg kemény kapszula	not available	OGYI-T-7724/05	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Alvogen 50 mg/ml oldatos injekció	not available	OGYI-T-7724/03	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Alvogen 50 mg/ml oldatos injekció	not available	OGYI-T-7724/04	ALVOGEN PHARMA TRADING EUROPE EOOD	HU
Tramadol Andrómaco 100 mg/2 ml solución inyectable y para perfusión EFG	not available	83760	ANDRÓMACO PHARMA S.L.	ES
Tramadol Andrómaco 100 mg/ml solución oral EFG	not available	83759	ANDRÓMACO PHARMA S.L.	ES
Tramadol Andrómaco 50 mg cápsulas duras EFG	not available	83758	ANDRÓMACO PHARMA S.L.	ES
Tramadol Apotex 50 mg cápsulas duras EFG	not available	65724	APOTEX EUROPE B.V.	ES
Tramadol Aristo 100 mg comprimidos de liberación prolongada EFG	not available	65.833	ARISTO PHARMA IBERIA, S.L.	ES
Tramadol Aristo 150 mg comprimidos de	not available	65.834	ARISTO PHARMA IBERIA, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
liberación prolongada EFG				
Tramadol Aristo 200 mg comprimidos de liberación prolongada EFG	not available	65.835	ARISTO PHARMA IBERIA, S.L.	ES
TRAMADOL ARISTO 50 mg cápsulas E.F.G.	not available	63.451	ARISTO PHARMA IBERIA, S.L.	ES
Tramadol Aristo® 100 mg Retardtabletten	not available	50307.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Tramadol Aristo® 150 mg Retardtabletten	not available	50307.01.00	ARISTO PHARMA GMBH (ART 57)	DE
Tramadol Aristo® 200 mg Retardtabletten	not available	50307.02.00	ARISTO PHARMA GMBH (ART 57)	DE
TRAMADOL ARROW 100 mg/2 mL, solution injectable/pour perfusion	not available	NL 46364	ARROW GENERIQUES	FR
TRAMADOL ARROW 50 mg, comprimé	not available	NL 26860	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 100 mg, comprimé à libération prolongée	not available	NL 36433	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 100 mg, gélule à libération prolongée	not available	NL32723	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 150 mg, comprimé à libération prolongée	not available	NL 36434	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 150 mg, gélule à libération prolongée	not available	NL 32724	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 200 mg, comprimé à libération prolongée	not available	NL 36435	ARROW GENERIQUES	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
TRAMADOL ARROW L.P. 200 mg, gélule à libération prolongée	not available	NL 32725	ARROW GENERIQUES	FR
TRAMADOL ARROW LAB 50 mg, gélule	NL/H/2480/001	NL 41670	ARROW GENERIQUES	FR
TRAMADOL ASTA Medica 100 mg Solución Inyectable EFG	not available	63465	MEDA PHARMA S.L.	ES
Tramadol ASTA Medica 100 mg/ml gotas orales en solución EFG.	not available	62925	MEDA PHARMA S.L.	ES
Tramadol ASTA Medica 50 mg cápsulas duras EFG	not available	61849	MEDA PHARMA S.L.	ES
Tramadol Aurobindo 50 mg capsules, hard	NL/H/2480/001	MA807/05401	AUROBINDO PHARMA (MALTA) LIMITED	MT
Tramadol Aurobindo 50 mg kapslar, hårda	NL/H/2480/001	46985	AUROBINDO PHARMA (MALTA) LIMITED	SE
Tramadol Aurobindo, 50 mg, kapsułki, twarde	NL/H/2480/001	20928	AUROBINDO PHARMA (MALTA) LIMITED	PL
Tramadol Aurovitas 100 mg comprimidos de libertação prolongada	NL/H/0890/001	5032016	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 100 mg comprimidos de libertação prolongada	NL/H/0890/001	5580527	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 100 mg comprimidos de libertação prolongada	NL/H/0890/001	5032024	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 100 mg comprimidos de libertação prolongada	NL/H/0890/001	5032032	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 150 mg comprimidos de	NL/H/0890/002	5032040	GENERIS FARMACÊUTICA, 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
libertação prolongada				
Tramadol Aurovitas 150 mg comprimidos de libertação prolongada	NL/H/0890/002	5580535	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 150 mg comprimidos de libertação prolongada	NL/H/0890/002	5032057	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 150 mg comprimidos de libertação prolongada	NL/H/0890/002	5032065	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 200 mg comprimidos de libertação prolongada	NL/H/0890/003	5032073	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 200 mg comprimidos de libertação prolongada	NL/H/0890/003	5580543	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 200 mg comprimidos de libertação prolongada	NL/H/0890/003	5032107	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 200 mg comprimidos de libertação prolongada	NL/H/0890/003	5032115	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 50 mg cápsulas	PT/H/1576/001	5700257	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 50 mg cápsulas	PT/H/1576/001	PT/H/1576/001	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 50 mg cápsulas	PT/H/1576/001	PT/H/1576/001	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 50 mg cápsulas duras EFG	PT/H/1576/001	81.624	AUROVITAS SPAIN,S.A.U.	ES
Tramadol Aurovitas 50 mg tvrdé tobolky	PT/H/1576/001	65/806/15-C	AUROVITAS, SPOL. S R.O.	CZ
Tramadol Aurovitas 50 mg/ml solução injetável	PT/H/1548/001	5703368	GENERIS FARMACÊUTICA, 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
ou para perfusão				
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	5703376	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Aurovitas 50 mg/ml solución inyectable y para perfusión EFG	PT/H/1548/001	81.910	AUROVITAS SPAIN,S.A.U.	ES
Tramadol Aurovitas, 50 mg, kapsułki, twarde	PT/H/1576/001	24556	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
Tramadol axcount 100 mg retard, Retardtabletten	not available	50310.00.00	AXCOUNT GENERIKA GMBH	DE
Tramadol axcount 150 mg retard, Retardtabletten	not available	50310.01.00	AXCOUNT GENERIKA GMBH	DE
Tramadol axcount 200 mg retard, Retardtabletten	not available	50310.02.00	AXCOUNT GENERIKA GMBH	DE
Tramadol axcount 50 mg Brausetabletten	not available	37018.00.00	AXCOUNT GENERIKA GMBH	DE
Tramadol axcount Tropfen, Lösung zum Einnehmen	not available	35164.00.00	AXCOUNT GENERIKA GMBH	DE
Tramadol Azevedos 50 mg cápsulas	not available	5012273	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50 mg cápsulas	not available	5012307	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50 mg cápsulas	not available	5012315	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50	not available	5012331	LABORATÓRIOS AZEVEDOS	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
mg cápsulas			- INDÚSTRIA FARMACÊUTICA, S.A.	
Tramadol Azevedos 50 mg cápsulas	not available	5012349	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50 mg cápsulas	not available	5012356	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50 mg cápsulas	not available	5012323	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Azevedos 50 mg cápsulas	not available	5012265	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Basi, 100 mg/2 ml Solução Injectável	not available	497 87 97	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Basi, 100 mg/2 ml Solução Injectável	not available	324 25 91	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Basi, 50 mg/1 ml Solução Injectável.	not available	497 86 98	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
Tramadol Basi, 50 mg/1 ml Solução Injectável.	not available	324 27 99	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
TRAMADOL BGR L.P. 100 mg, comprimé à libération prolongée	not available	3400938675382	BIOGARAN	FR
TRAMADOL BGR L.P. 100 mg, comprimé à libération prolongée	not available	3400957311391	BIOGARAN	FR
TRAMADOL BGR L.P. 100 mg, comprimé à	not available	3400938675214	BIOGARAN	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
libération prolongée				
TRAMADOL BGR L.P. 100 mg, comprimé à libération prolongée	not available	3400930176764	BIOGARAN	FR
TRAMADOL BGR L.P. 150 mg, comprimé à libération prolongée	not available	3400938747393	BIOGARAN	FR
TRAMADOL BGR L.P. 150 mg, comprimé à libération prolongée	not available	3400938747454	BIOGARAN	FR
TRAMADOL BGR L.P. 150 mg, comprimé à libération prolongée	not available	3400957326531	BIOGARAN	FR
TRAMADOL BGR L.P. 200 mg, comprimé à libération prolongée	not available	3400938675443	BIOGARAN	FR
TRAMADOL BGR L.P. 200 mg, comprimé à libération prolongée	not available	3400957311452	BIOGARAN	FR
TRAMADOL BGR L.P. 200 mg, comprimé à libération prolongée	not available	3400938675504	BIOGARAN	FR
TRAMADOL BIOGARAN 50 mg, gélule	not available	3400935248442	BIOGARAN	FR
TRAMADOL BIOGARAN 50 mg, gélule	not available	3400955047490	BIOGARAN	FR
TRAMADOL BIOGARAN 50 mg, gélule	not available	3400930127209	BIOGARAN	FR
Tramadol Ciclum 100 mg/ml solução oral	not available	2668986	CICLUM FARMA UNIPESSOAL LDA.	PT
Tramadol Ciclum 100 mg/ml solução oral	not available	2709988	CICLUM FARMA UNIPESSOAL LDA.	PT
Tramadol Ciclum 100 mg/ml solução oral	not available	3081387	CICLUM FARMA UNIPESSOAL LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramadol Ciclum 50 mg cápsulas	not available	2679686	CICLUM FARMA UNIPESSOAL LDA.	PT
Tramadol Ciclum 50 mg cápsulas	not available	2679587	CICLUM FARMA UNIPESSOAL LDA.	PT
tramadol cinfa 50 mg cápsulas duras EFG	not available	63.440	LABORATORIOS CINFA, S.A.	ES
TRAMADOL CRISTERS LP 100 mg, comprimé pelliculé à libération prolongée	not available	34009 493 635 9 9	CRISTERS	FR
TRAMADOL CRISTERS LP 100 mg, comprimé pelliculé à libération prolongée	not available	34009 493 636 5 0	CRISTERS	FR
TRAMADOL CRISTERS LP 100 mg, comprimé pelliculé à libération prolongée	not available	34009 493 637 1 1	CRISTERS	FR
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 493 638 8 9	CRISTERS	FR
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 493 639 4 0	CRISTERS	FR
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 493 640 2 2	CRISTERS	FR
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 493 641 9 0	CRISTERS	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
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 578 149 1 5	CRISTERS	FR
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 578 151 6 5	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 493 785 0 0	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 493 786 7 8	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 493 787 3 9	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 493 789 6 8	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 578 169 2 6	CRISTERS	FR
TRAMADOL CRISTERS LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 578 170 0 8	CRISTERS	FR
Tramadol Denk 100 mg Injektionslösung	not available	38542.00.00	DENK PHARMA 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
Tramadol Denk 50 mg Brause	not available	34728.00.00	DENK PHARMA GMBH & CO. KG	DE
Tramadol EG 100 mg/2 ml Injektionslösung	DE/H/0282/004	BE245786	EUROGENERICS N.V./S.A.	BE
Tramadol EG 100 mg/2 ml oplossing voor injectie	DE/H/0282/004	BE245786	EUROGENERICS N.V./S.A.	BE
Tramadol EG 100 mg/2 ml solution injectable	DE/H/0282/004	0019/08039713	EUROGENERICS N.V./S.A.	LU
Tramadol EG 100 mg/2ml solution injectable	DE/H/0282/004	BE245786	EUROGENERICS N.V./S.A.	BE
Tramadol EG 100 mg/ml druppels voor oraal gebruik, oplossing	DE/H/0282/002	BE245707	EUROGENERICS N.V./S.A.	BE
Tramadol EG 100 mg/ml solution buvable en gouttes	DE/H/0282/002	BE245707	EUROGENERICS N.V./S.A.	BE
Tramadol EG 100 mg/ml solution buvable en gouttes	DE/H/0282/002	0019/08039712	EUROGENERICS N.V./S.A.	LU
Tramadol EG 100 mg/ml Tropfen zum Einnehmen, Lösung	DE/H/0282/002	BE245707	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg comprimés	BE/H/0298/001	BE216474	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg comprimés	BE/H/0298/001	0019/04060001	EUROGENERICS N.V./S.A.	LU
Tramadol EG 50 mg tabletten	BE/H/0298/001	BE216474	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg Tabletten	BE/H/0298/001	BE216474	EUROGENERICS N.V./S.A.	BE
TRAMADOL EG 50 mg, comprimé	not available	NL24562	EG LABO LABORATOIRES EUROGENERICS	FR
TRAMADOL EG L.P. 100	not available	NL37921	EG LABO LABORATOIRES	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
mg, comprimé à libération prolongée			EUROGENERICS	
TRAMADOL EG L.P. 150 mg, comprimé à libération prolongée	not available	NL37922	EG LABO LABORATOIRES EUROGENERICS	FR
TRAMADOL EG L.P. 200 mg, comprimé à libération prolongée	not available	NL37923	EG LABO LABORATOIRES EUROGENERICS	FR
TRAMADOL ETHYPHARM 100 mg, Hartkapsel, retardiert	DE/H/0639/002	54160.01.00	ETHYPHARM	DE
TRAMADOL ETHYPHARM 150 mg, Hartkapsel, retardiert	DE/H/0639/003	54160.02.00	ETHYPHARM	DE
Tramadol Ethypharm 200mg, Hartkapsel, retardiert	DE/H/0639/004	54160.03.00	ETHYPHARM	DE
TRAMADOL ETHYPHARM 50 mg, Hartkapsel, retardiert	DE/H/0639/001	54160.00.00	ETHYPHARM	DE
TRAMADOL EVOLUGEN 50 mg, gélule	not available	34009 266 909 0 8	EVOLUPHARM	FR
TRAMADOL EVOLUGEN L.P. 100 mg, comprimé à libération prolongée	not available	34009 301 803 9 6	EVOLUPHARM	FR
TRAMADOL EVOLUGEN L.P. 150 mg, comprimé à libération prolongée	not available	34009 301 804 4 0	EVOLUPHARM	FR
TRAMADOL EVOLUGEN L.P. 200 mg, comprimé à libération prolongée	not available	34009 301 804 8 8	EVOLUPHARM	FR
TRAMADOL FARMALIDER 100 mg/ml, solución oral .E.F.G	not available	69404	FARMALIDER, 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
Tramadol Generis 100 mg/ml solução oral	not available	3759396	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis 100 mg/ml solução oral	not available	5598990	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis 50 mg cápsulas	not available	5723606	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis Phar, 50 mg, Cápsula	not available	3759495	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis Phar, 50 mg, Cápsula	not available	3759594	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis, 100 mg/2 ml, solução injectável	not available	3759198	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis, 100 mg/2 ml, solução injectável	not available	3759297	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol HCl Apotex 50 mg, capsules	not available	RVG 21626	APOTEX EUROPE B.V.	NL
Tramadol HCl Aurobindo 50 mg, capsules, hard	NL/H/2480/001	RVG 110742	AUROBINDO PHARMA B.V.	NL
Tramadol HCl Aurobindo Retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0890/001	RVG 33553	AUROBINDO PHARMA B.V.	NL
Tramadol HCl Aurobindo Retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0890/002	RVG 33554	AUROBINDO PHARMA B.V.	NL
Tramadol HCl Aurobindo Retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0890/003	RVG 33555	AUROBINDO PHARMA B.V.	NL
Tramadol HCl capsule CF 50 mg, capsules	DE/H/0282/001	RVG 27163	CENTRAFARM B.V.	NL
Tramadol HCl druppels CF 100 mg/ml, druppels voor orale toediening.	DE/H/0282/002	RVG 27165	CENTRAFARM 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
Tramadol HCl Duiven retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0539/001	RVG 30577	ICC B.V.	NL
Tramadol HCl Duiven retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0538/001	RVG 30562	ICC B.V.	NL
Tramadol HCl Duiven retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0539/002	RVG 30578	ICC B.V.	NL
Tramadol HCl Duiven retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0538/002	RVG 30563	ICC B.V.	NL
Tramadol HCl Duiven retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0539/003	RVG 30579	ICC B.V.	NL
Tramadol HCl Duiven retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0538/003	RVG 30564	ICC B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl Nevik 50 mg, capsules.	not available	RVG 21682	NEVIK LIMITED	NL
Tramadol HCl Nevik 50 mg, capsules.	not available	RVG 21682	NEVIK LIMITED	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
Tramadol HCl Nevik 50 mg, capsules.	not available	RVG 21682	NEVIK LIMITED	NL
Tramadol HCl Retard 100 mg Teva, tabletten met gereguleerde afgifte	not available	RVG 33751	TEVA NEDERLAND B.V.	NL
Tramadol HCl Retard 100 mg, tabletten met gereguleerde afgifte	not available	RVG 35254	ICC B.V.	NL
Tramadol HCl retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0888/001	RVG 32921	ICC B.V.	NL
Tramadol HCl retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0892/001	RVG 33754	ICC B.V.	NL
Tramadol HCl retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0889/001	RVG 32924	ICC B.V.	NL
Tramadol HCl Retard 150 mg Teva, tabletten met gereguleerde afgifte	not available	RVG 33752	TEVA NEDERLAND B.V.	NL
Tramadol HCl Retard 150 mg, tabletten met gereguleerde afgifte	not available	RVG 35255	ICC B.V.	NL
Tramadol HCl retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0888/002	RVG 32922	ICC B.V.	NL
Tramadol HCl retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0892/002	RVG 33755	ICC B.V.	NL
Tramadol HCl retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0889/002	RVG 32925	ICC B.V.	NL
Tramadol HCl Retard 200 mg Teva, tabletten met	not available	RVG 33753	TEVA NEDERLAND 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
gereguleerde afgifte				
Tramadol HCl Retard 200 mg, tabletten met gereguleerde afgifte	not available	RVG 35256	ICC B.V.	NL
Tramadol HCl retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0888/003	RVG 32923	ICC B.V.	NL
Tramadol HCl retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0892/003	RVG 33756	ICC B.V.	NL
Tramadol HCl retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0889/003	RVG 32926	ICC B.V.	NL
Tramadol HCl Retard Mylan 100 mg, harde capsules met verlengde afgifte	UK/H/0225/002	RVG 22326	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met gereguleerde afgifte	not available	RVG 25699	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met gereguleerde afgifte	not available	RVG 25699	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met gereguleerde afgifte	not available	RVG 25699	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met gereguleerde afgifte	not available	RVG 25699	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met gereguleerde afgifte	not available	RVG 25699	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten	not available	RVG 25699	MYLAN 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
met gereguleerde afgifte				
Tramadol HCl Retard Mylan 100 mg, tabletten met verlengde afgifte	DK/H/2942/001	RVG 109064	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met verlengde afgifte	DK/H/2942/001	RVG 109064	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met verlengde afgifte	DK/H/2942/001	RVG 109064	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 100 mg, tabletten met verlengde afgifte	DK/H/2942/001	RVG 109064	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, harde capsules met verlengde afgifte	UK/H/0225/003	RVG 22327	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met gereguleerde afgifte	not available	RVG 25700	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met gereguleerde afgifte	not available	RVG 25700	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met gereguleerde afgifte	not available	RVG 25700	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met gereguleerde afgifte	not available	RVG 25700	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met gereguleerde afgifte	not available	RVG 25700	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten	not available	RVG 25700	MYLAN 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
met gereguleerde afgifte				
Tramadol HCl Retard Mylan 150 mg, tabletten met verlengde afgifte	DK/H/2942/002	RVG 109065	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met verlengde afgifte	DK/H/2942/002	RVG 109065	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met verlengde afgifte	DK/H/2942/002	RVG 109065	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 150 mg, tabletten met verlengde afgifte	DK/H/2942/002	RVG 109065	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, harde capsules met verlengde afgifte	UK/H/0225/004	RVG 22328	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met gereguleerde afgifte	not available	RVG 25701	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met gereguleerde afgifte	not available	RVG 25701	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met gereguleerde afgifte	not available	RVG 25701	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met gereguleerde afgifte	not available	RVG 25701	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met gereguleerde afgifte	not available	RVG 25701	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten	not available	RVG 25701	MYLAN 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
met gereguleerde afgifte				
Tramadol HCl Retard Mylan 200 mg, tabletten met verlengde afgifte	DK/H/2942/003	RVG 109066	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met verlengde afgifte	DK/H/2942/003	RVG 109066	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met verlengde afgifte	DK/H/2942/003	RVG 109066	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 200 mg, tabletten met verlengde afgifte	DK/H/2942/003	RVG 109066	MYLAN B.V.	NL
Tramadol HCl Retard Mylan 50 mg, harde capsules met verlengde afgifte	UK/H/0225/001	RVG 22325	MYLAN B.V.	NL
Tramadol HCl Sandoz bruis 50 mg, bruistabletten	DE/H/0144/001	RVG 23527	SANDOZ B.V.	NL
Tramadol HCl Sandoz capsule 50 mg, capsules	not available	RVG 21690	SANDOZ B.V.	NL
Tramadol HCl Sandoz retard 100, tabletten met verlengde afgifte 100 mg	NL/H/0483/001	RVG 25693	SANDOZ B.V.	NL
Tramadol HCl Sandoz retard 150, tabletten met verlengde afgifte 150 mg	NL/H/0483/002	RVG 25694	SANDOZ B.V.	NL
Tramadol HCl Sandoz retard 200, tabletten met verlengde afgifte 200 mg	NL/H/0483/003	RVG 25695	SANDOZ B.V.	NL
Tramadol HCl Teva 100 mg/ml, druppels voor oraal gebruik, oplossing	not available	RVG 27177	TEVA 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
Tramadol HCl Teva 50 mg, capsules	not available	RVG 22031	TEVA NEDERLAND B.V.	NL
Tramadol HCl zetpil CF 100 mg, zetpillen	DE/H/0282/003	RVG 27166	CENTRAFARM B.V.	NL
TRAMADOL HCS 100 mg/2 mL, solution injectable/pour perfusion	MT/H/0260/001	34009 550 583 2 8	HCS BVBA	FR
TRAMADOL HCS 100 mg/2 mL, solution injectable/pour perfusion	MT/H/0260/001	34009 550 583 3 5	HCS BVBA	FR
Tramadol HCS 50 mg hard capsules	MT/H/0259/001	MA1082/00101	HCS BVBA	MT
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 301 519 0 7	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 301 519 1 4	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 301 519 2 1	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 301 519 3 8	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 301 519 4 5	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 550 566 2 1	HCS BVBA	FR
TRAMADOL HCS 50 mg, gélule	MT/H/0259/001	34009 550 566 3 8	HCS BVBA	FR
TRAMADOL HCS 50 mg/1 mL, solution injectable/pour perfusion	MT/H/0260/001	34009 550 583 0 4	HCS BVBA	FR
TRAMADOL HCS 50 mg/1 mL, solution injectable/pour perfusion	MT/H/0260/001	34009 550 583 1 1	HCS BVBA	FR
Tramadol Heumann Tropfen Tropfen zum	not available	25530.00.00	HEUMANN PHARMA GMBH & CO. GENERICA 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
Einnehmen, Lösung mit 100 mg Tramadolhydrochlorid/ml				
Tramadol Hexal 100 mg kapslar, hårda	not available	13466	HEXAL A/S	SE
Tramadol Hexal 100 mg kovat kapselit	not available	12731	HEXAL A/S	FI
Tramadol HEXAL 50 mg kapsel, hard	not available	99-7755	HEXAL A/S	NO
Tramadol Hexal 50 mg kapslar, hårda	not available	13465	HEXAL A/S	SE
Tramadol Hexal 50 mg kovat kapselit	not available	12729	HEXAL A/S	FI
Tramadol Hydrochloride 50 mg Capsules	not available	PL 44041/0037	NOUMED LIFE SCIENCES	UK
Tramadol Hydrochloride 50 mg Capsules	not available	PL 21880/0167	MEDREICH PLC	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride	not available	PL 25298/0240	BROWN & BURK 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
50 mg capsules, hard				
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg capsules, hard	not available	PL 25298/0240	BROWN & BURK UK LIMITED	UK
Tramadol hydrochloride 50 mg Hard Capsules.	IE/H/0591	PA 2315/214/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Tramadol hydrochloride 50 mg/ml solution for injection or infusion	PT/H/1548/001	PL 16363/0467	MILPHARM LIMITED	UK
Tramadol hydrochloride 50 mg/ml solution for injection or infusion	UK/H/5235/001	PL 01502/0085	HAMELN PHARMA LTD	UK
Tramadol Hydrochloride 50mg Capsules	not available	PL 11311/0084	TILLOMED LABORATORIES LTD	UK
Tramadol Hydrochloride 50mg Capsules	not available	PL 20117/0086	MORNINGSIDE HEALTHCARE LTD	UK
Tramadol Hydrochloride 50mg Capsules	not available	PL 30464/0037	ATHLONE PHARMACEUTICALS LIMITED	UK
TRAMADOL HYDROCHLORIDE 50mg CAPSULES	UK/H/0380/001	PL 0142/0484	ACTAVIS UK LIMITED	UK
Tramadol Hydrochloride 50mg Effervescent Tablets	not available	PL 36722/0119	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Tramadol Hydrochloride Capsules 50mg	not available	PL 20075/0290	ACCORD HEALTHCARE LIMITED	UK
Tramadol hydrochloride Krka 100 mg prolonged-release tablets	EE/H/0262/001	PL 01656/0264	KRKA, D.D., NOVO MESTO	UK
Tramadol hydrochloride Krka 150 mg prolonged-	EE/H/0262/002	PL 01656/0265	KRKA, D.D., NOVO MESTO	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
release tablets				
Tramadol hydrochloride Krka 200 mg prolonged-release tablets	EE/H/0262/003	PL 01656/0266	KRKA, D.D., NOVO MESTO	UK
Tramadol Kalceks 100 mg/2 ml injekční/infuzní roztok	LV/H/0152/001	65/169/17-C	KALCEKS	CZ
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/001	KALCEKS	LT
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/002	KALCEKS	LT
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/003	KALCEKS	LT
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/004	KALCEKS	LT
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/005	KALCEKS	LT
Tramadol Kalceks 50 mg/ml injekcinis ar infuzinis tirpalas	LV/H/0152/001	LT/1/17/4142/006	KALCEKS	LT
Tramadol Kalceks 50 mg/ml Injektions-/Infusionslösung	LV/H/0152/001	137857	KALCEKS	AT
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/01	KALCEKS	HU
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/02	KALCEKS	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
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/03	KALCEKS	HU
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/04	KALCEKS	HU
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/05	KALCEKS	HU
Tramadol Kalceks 50 mg/ml oldatos injekció vagy infúzió	LV/H/0152/001	OGYI-T-23256/06	KALCEKS	HU
Tramadol Kalceks 50 mg/ml šķīdums injekcijām/infūzijām	LV/H/0152/001	16-0115	KALCEKS	LV
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/01	KALCEKS	RO
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/02	KALCEKS	RO
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/03	KALCEKS	RO
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/04	KALCEKS	RO
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/05	KALCEKS	RO
Tramadol Kalceks 50mg/ml soluție injectabilă/perfuzabilă	LV/H/0152/001	10308/2017/06	KALCEKS	RO
Tramadol Kalceks, 50	LV/H/0152/001	24390	KALCEKS	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
mg/ml, roztwór do wstrzykiwań/infuzji				
TRAMADOL KERN PHARMA 50 mg cápsulas EFG	not available	63.920	KERN PHARMA, S.L.	ES
Tramadol Krka 100 mg čepići	not available	HR-H-767374749	KRKA-FARMA D.O.O.	HR
Tramadol Krka 100 mg comprimidos de liberación prolongada EFG	EE/H/0262/001	83784	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 100 mg depottabletti	EE/H/0262/001	35806	KRKA, D.D., NOVO MESTO	FI
Tramadol Krka 100 mg forðatöflur.	EE/H/0262/001	IS/1/18/119/01	KRKA, D.D., NOVO MESTO	IS
Tramadol Krka 100 mg ilgstošās darbības tabletes	not available	00-0666	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 100 mg otopina za injekciju/infuziju	not available	HR-H-380362751	KRKA-FARMA D.O.O.	HR
Tramadol Krka 100 mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1055/007	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/25	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/26	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/27	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/28	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/29	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/30	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/31	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/32	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/33	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/34	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/35	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg retard tabletta	EE/H/0262/001	OGYI-T-23380/36	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg tablete s produljenim oslobađanjem	EE/H/0262/001	HR-H-141300728	KRKA-FARMA D.O.O.	HR
Tramadol Krka 100 mg tabletten met verlengde afgifte	EE/H/0262/001	BE535786	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 100 mg tablety s predĺženým uvoľňovaním	EE/H/0262/001	65/0366/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 100 mg tablety s prodlouženým uvolňováním	EE/H/0262/001	65/609/17-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 100 mg/ 2 ml šķīdums injekcijām	not available	01-0148	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 100 mg/2 ml injekcinis tirpalas	not available	LT/1/94/1055/006	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 100 mg/2 ml injekční/infuzní roztok	HR/H/0104/ 002	65/1050/16-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 100 mg/2 ml injekčný roztok/	HR/H/0104/002	65/0108/18-S	KRKA, D.D., NOVO MESTO	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
infúzny roztok				
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/07	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/08	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/09	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/10	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/11	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/12	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/2 ml oplossing voor injectie/infusie	HR/H/0104/ 002	BE530222	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 100 mg/2 ml solución inyectable y para perfusión EFG	HR/H/0104/002	83132	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 100 mg/ml belsőleges oldatos cseppek	HR/H/0126/001	OGYI-T-23380/20	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/ml belsőleges oldatos cseppek	HR/H/0126/001	OGYI-T-23380/21	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/ml belsőleges oldatos cseppek	HR/H/0126/001	OGYI-T-23380/22	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 100 mg/ml belsőleges oldatos cseppek	HR/H/0126/001	OGYI-T-23380/23	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/ml belsőleges oldatos cseppek	HR/H/0126/001	OGYI-T-23380/24	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 100 mg/ml druppels voor oraal gebruik, oplossing	HR/H/0126/001	BE533982	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 100 mg/ml druppels voor oraal gebruik, oplossing	HR/H/0126/001	BE533991	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 100 mg/ml geriamieji lašai (tirpalas)	not available	LT/1/94/1055/003	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 100 mg/ml geriamieji lašai (tirpalas)	not available	LT/1/94/1055/004	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 100 mg/ml gotas orais, solução	HR/H/0126/	5756309	KRKA, D.D., NOVO MESTO	PT
Tramadol Krka 100 mg/ml gotas orais, solução	HR/H/0126/	5756416	KRKA, D.D., NOVO MESTO	PT
Tramadol Krka 100 mg/ml gotas orales en solución EFG	MT/H/0272/001	83837	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 100 mg/ml oralne kapi, otopina	not available	HR-H-121650469	KRKA-FARMA D.O.O.	HR
Tramadol Krka 100 mg/ml perorálne roztokové kvapky	HR/H/0126/001	65/0259/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 100	HR/H/0126/001	65/328/17-C	KRKA, D.D., NOVO MESTO	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
mg/ml perorální roztok				
Tramadol Krka 100 mg/ml picături orale, soluție	HR/H/0126/001	11210/2018/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Krka 100 mg/ml picături orale, soluție	HR/H/0126/001	11210/2018/02	KRKA, D.D., NOVO MESTO	RO
Tramadol Krka 100 mg/ml picături orale, soluție	HR/H/0126/001	11210/2018/03	KRKA, D.D., NOVO MESTO	RO
Tramadol Krka 100 mg/ml picături orale, soluție	HR/H/0126/001	11210/2018/04	KRKA, D.D., NOVO MESTO	RO
Tramadol Krka 100 mg/ml picături orale, soluție	HR/H/0126/001	11210/2018/05	KRKA, D.D., NOVO MESTO	RO
Tramadol Krka 150 mg comprimidos de liberación prolongada EFG	EE/H/0262/002	83785	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 150 mg depottabletti	EE/H/0262/002	35807	KRKA, D.D., NOVO MESTO	FI
Tramadol Krka 150 mg forðatöflur.	EE/H/0262/002	IS/1/18/119/02	KRKA, D.D., NOVO MESTO	IS
Tramadol Krka 150 mg ilgstošās darbības tabletes	not available	03-0206	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 150 mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1055/008	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/37	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/38	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/39	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/40	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/41	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/42	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/43	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/44	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/45	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/46	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/47	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg retard tabletta	EE/H/0262/002	OGYI-T-23380/48	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 150 mg tablete s produljenim oslobađanjem	EE/H/0262/002	HR-H-213943343	KRKA-FARMA D.O.O.	HR
Tramadol Krka 150 mg tabletten met verlengde afgifte	EE/H/0262/002	BE535795	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 150 mg tablety s predĺženým uvoľňovaním	EE/H/0262/002	65/0367/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 150 mg tablety s prodlouženým uvolňováním	EE/H/0262/002	65/610/17-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 200 mg comprimidos de	EE/H/0262/003	83786	KRKA, D.D., NOVO MESTO	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
liberación prolongada EFG				
Tramadol Krka 200 mg depottabletti	EE/H/0262/003	35808	KRKA, D.D., NOVO MESTO	FI
Tramadol Krka 200 mg forðatöflur.	EE/H/0262/003	IS/1/18/119/03	KRKA, D.D., NOVO MESTO	IS
Tramadol Krka 200 mg ilgstošās darbības tabletes	not available	03-0207	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 200 mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1055/009	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/49	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/50	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/51	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/52	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/53	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/54	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/55	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/56	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/57	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/58	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/59	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 200 mg retard tabletta	EE/H/0262/003	OGYI-T-23380/60	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 200 mg tablete s produljenim oslobađanjem	EE/H/0262/003	HR-H-038452159	KRKA-FARMA D.O.O.	HR
Tramadol Krka 200 mg tabletten met verlengde afgifte	EE/H/0262/003	BE535804	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 200 mg tablety s predĺženým uvoľňovaním	EE/H/0262/003	65/0368/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 200 mg tablety s prodlouženým uvolňováním	EE/H/0262/003	65/611/17-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 50 mg cápsulas	HR/H/0104/003	5736327	KRKA, D.D., NOVO MESTO	PT
Tramadol Krka 50 mg cápsulas duras EFG	HR/H/0104/003	83133	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 50 mg cietās kapsulas	not available	96-0110	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 50 mg hard capsules	HR/H/0104/003	PA1347/079/001	KRKA, D.D., NOVO MESTO	IE
Tramadol Krka 50 mg hard capsules	HR/H/0104/003	PL 01656/0248	KRKA, D.D., NOVO MESTO	UK
Tramadol Krka 50 mg harde capsules	HR/H/0104/003	BE526604	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 50 mg kapslar, hårda	HR/H/0104/003	56007	KRKA, D.D., NOVO MESTO	SE
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/003	OGYI-T-23380/13	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/003	OGYI-T-23380/14	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/ 003	OGYI-T-23380/15	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/ 003	OGYI-T-23380/16	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/ 003	OGYI-T-23380/17	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/ 003	OGYI-T-23380/18	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kemény kapszula	HR/H/0104/ 003	OGYI-T-23380/19	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg kietosios kapsulės	not available	LT/1/94/1055/001	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 50 mg otopina za injekciju/infuziju	not available	HR-H-883684879	KRKA-FARMA D.O.O.	HR
Tramadol Krka 50 mg tvrde kapsule	not available	HR-H-083262314	KRKA-FARMA D.O.O.	HR
Tramadol Krka 50 mg tvrdé kapsuly	HR/H/0104/003	65/0142/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 50 mg tvrdé tobolky	HR/H/0104/ 003	65/036/17-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 50 mg/1 ml injekční/infuzní roztok	HR/H/0104/ 002	65/826/16-C	KRKA, D.D., NOVO MESTO	CZ
Tramadol Krka 50 mg/1 ml oplossing voor injectie/infusie	HR/H/0104/ 002	BE530213	KRKA, D.D., NOVO MESTO	BE
Tramadol Krka 50 mg/1 ml solución inyectable y para perfusión EFG	HR/H/0104/001	83131	KRKA, D.D., NOVO MESTO	ES
Tramadol Krka 50 mg/ml injekcinis tirpalas	not available	LT/1/94/1055/005	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 50 mg/ml injekčný roztok/ infúzny roztok	HR/H/0104/002	65/0107/18-S	KRKA, D.D., NOVO MESTO	SK
Tramadol Krka 50 mg/ml oldatos injekció vagy	HR/H/0104/ 002	OGYI-T-23380/01	KRKA, D.D., NOVO MESTO	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
infúzió				
Tramadol Krka 50 mg/ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/02	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg/ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/03	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg/ml oldatos injekció vagy infúzió	HR/H/0104/002	OGYI-T-23380/04	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg/ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/05	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg/ml oldatos injekció vagy infúzió	HR/H/0104/ 002	OGYI-T-23380/06	KRKA, D.D., NOVO MESTO	HU
Tramadol Krka 50 mg/ml šķīdums injekcijām	not available	96-0107	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 50 mg/ml süstelahus	not available	135196	KRKA, D.D., NOVO MESTO	EE
TRAMADOL KRKA LP 100 mg, comprimé à libération prolongée	EE/H/0262/001	34009 301 650 1 0	KRKA, D.D., NOVO MESTO	FR
TRAMADOL KRKA LP 100 mg, comprimé à libération prolongée	EE/H/0262/001	34009 301 650 2 7	KRKA, D.D., NOVO MESTO	FR
TRAMADOL KRKA LP 150 mg, comprimé à libération prolongée	EE/H/0262/002	34009 301 649 9 0	KRKA, D.D., NOVO MESTO	FR
TRAMADOL KRKA LP 150 mg, comprimé à libération prolongée	EE/H/0262/002	34009 301 650 0 3	KRKA, D.D., NOVO MESTO	FR
TRAMADOL KRKA LP 200 mg, comprimé à	EE/H/0262/003	34009 301 649 7 6	KRKA, D.D., NOVO MESTO	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
libération prolongée				
TRAMADOL KRKA LP 200 mg, comprimé à libération prolongée	EE/H/0262/003	34009 301 649 8 3	KRKA, D.D., NOVO MESTO	FR
Tramadol Krka, 100 mg toimeainet prolongeeritult vabastavad tabletid	not available	331200	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 100 mg, tabletki o przedłużonym uwalnianiu	EE/H/0262/001	25264	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 100 mg/2 mL, roztwór do wstrzykiwań/infuzji	HR/H/0104/002	24585	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 100 mg/ml suukaudsed tilgad, lahus	not available	148696	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 100 mg/ml, krople doustne, roztwór	HR/H/0126/	25023	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 150 mg, tabletki o przedłużonym uwalnianiu	EE/H/0262/002	25265	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 200 mg, tabletki o przedłużonym uwalnianiu	EE/H/0262/003	25266	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 50 mg kõvakapslid	not available	139596	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 50 mg, kapsułki, twarde	HR/H/0104/ 003	24847	KRKA, D.D., NOVO MESTO	PL
Tramadol Krka, 50 mg/mL, roztwór do wstrzykiwań/infuzji	HR/H/0104/002	24584	KRKA, D.D., NOVO MESTO	PL
Tramadol Labesfal, 100	not available	4348587	GENERIS FARMACÊUTICA,	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
mg, comprimidos de libertaテァテ」o prolongada			S.A.	
Tramadol Labesfal, 100 mg, comprimidos de libertaテァテ」o prolongada	not available	5639638	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Labesfal, 100 mg/2 ml, solução injetável	not available	3210994	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 100 mg/2 ml, solução injetável	not available	4842399	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 50 mg cテ。psulas	not available	3827599	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Labesfal, 50 mg cテ。psulas	not available	3827698	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Labesfal, 50 mg/1 ml, solução injetável	not available	3210796	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 50 mg/1 ml, solução injetável	not available	4842290	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Lannacher 100 mg ilgstosas darbibas tabletes	not available	00-1168	G.L. PHARMA GMBH	LV
Tramadol Lannacher 100 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/002	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg pailginto atpalaidavimo tabletės	not available	LT/1/97/1431/001	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg, toimeainet prolongeeritult vabastavad tabletid	not available	345101	G.L. PHARMA GMBH	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
Tramadol Lannacher 100 mg/2 ml injekcinis tirpalas	not available	LT/1/97/1431/010	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg/2 ml injekcinis tirpalas	not available	LT/1/97/1431/011	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg/2 ml šķīdums injekcijām	not available	99-0235	G.L. PHARMA GMBH	LV
Tramadol Lannacher 100 mg/2 ml süstelahus	not available	321600	G.L. PHARMA GMBH	EE
Tramadol Lannacher 100 mg/ml geriamieji lašai, tirpalas	not available	LT/1/97/1431/012	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg/ml pilieni iekšķīgai lietošanai, šķīdums	not available	99-0514	G.L. PHARMA GMBH	LV
Tramadol Lannacher 150 mg ilgstosas darbibas tabletes	not available	03-0120	G.L. PHARMA GMBH	LV
Tramadol Lannacher 150 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/004	G.L. PHARMA GMBH	LT
Tramadol Lannacher 150 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/003	G.L. PHARMA GMBH	LT
Tramadol Lannacher 150 mg, toimeainet prolongeeritult vabastavad tabletid	not available	392202	G.L. PHARMA GMBH	EE
Tramadol Lannacher 200 mg ilgstosas darbibas tabletes	not available	03-0121	G.L. PHARMA GMBH	LV
Tramadol Lannacher 200	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	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
mg pailginto atpalaidavimo tabletes				
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	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
atpalaidavimo tabletes				
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/006	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg pailginto atpalaidavimo tabletes	not available	LT/1/97/1431/005	G.L. PHARMA GMBH	LT
Tramadol Lannacher 200 mg, toimeainet prolongeeritult vabastavad tabletid	not available	392302	G.L. PHARMA GMBH	EE
Tramadol Lannacher 50 mg apvalkotās tabletes	not available	99-0233	G.L. PHARMA GMBH	LV
Tramadol Lannacher 50 mg plėvele dengtos tabletės	not available	LT/1/97/1431/007	G.L. PHARMA GMBH	LT
Tramadol Lannacher 50 mg plėvele dengtos tabletės	not available	LT/1/97/1431/008	G.L. PHARMA GMBH	LT
Tramadol Lannacher, 100 mg/ml suukaudsed tilgad	not available	321700	G.L. PHARMA GMBH	EE
Tramadol Lannacher, 50 mg õhukese polümeerikilega kaetud tabletid	not available	274199	G.L. PHARMA GMBH	EE
TRAMADOL LAVOISIER 50 mg/ ml, solution injectable	not available	3400957486020	LABORATOIRES CHAIX ET DU MARAIS	FR
Tramadol LIBRAPHARM 100 mg, Injektionslösung	not available	13144.01.03	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM 100 mg/ml Lösung zum	not available	13144.00.02	LIBRA-PHARM 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
Einnehmen				
Tramadol LIBRAPHARM 50 mg, Injektionslösung	not available	13144.00.03	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM Kapseln, 50 mg, Hartkapseln	not available	13144.00.01	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM retard 100 mg Retardtabletten	DE/H/1093/002	69192.00.00	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM retard 150 mg Retardtabletten	DE/H/1093/003	69193.00.00	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM retard 200 mg Retardtabletten	DE/H/1093/004	69194.00.00	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM retard 50 mg Retardtabletten	DE/H/1093/001	69191.00.00	LIBRA-PHARM GMBH	DE
Tramadol LIBRAPHARM Zäpfchen, 100 mg	not available	13144.00.00	LIBRA-PHARM GMBH	DE
Tramadol Lösung - 1A-Pharma 100 mg/ml Tropfen zum Einnehmen, Lösung	not available	32751.00.01	1 A PHARMA GMBH	DE
Tramadol Medical Valley 50 mg hårda kapslar	DK/H/2693/001	55298	MEDICAL VALLEY INVEST AB	SE
Tramadol Medical Valley 50 mg harde kapsler	DK/H/2693/001	16-11326	MEDICAL VALLEY INVEST AB	NO
Tramadol Medical Valley 50 mg hörð hylki.	DK/H/2693/001	IS/1/18/019/01	MEDICAL VALLEY INVEST AB	IS
Tramadol MEIJI 50 mg cápsulas duras EFG	not available	83.616	MEIJI PHARMA SPAIN, S.A.	ES
Tramadol Mibe 100 mg tablete s produljenim	not available	HR-H-671676641	MIBE PHARMACEUTICALS 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
oslobađanjem				
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463179	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463211	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463318	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463302	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463203	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463310	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463179	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463211	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463318	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463302	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463203	MYLAN, 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
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463310	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463179	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463211	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463318	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463302	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463203	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463310	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463179	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463211	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463318	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463302	MYLAN, LDA	PT
Tramadol Mylan 100 mg	DK/H/2942/001	5463203	MYLAN, 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
comprimidos de libertação prolongada				
Tramadol Mylan 100 mg comprimidos de libertação prolongada	DK/H/2942/001	5463310	MYLAN, LDA	PT
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN GERMANY GMBH	DE
Tramadol Mylan 100 mg tablety s predĺženým uvoľňovaním	DK/H/2942/001	65/0556/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 100 mg tablety s predĺženým uvoľňovaním	DK/H/2942/001	65/0556/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 100 mg tablety s predĺženým uvoľňovaním	DK/H/2942/001	65/0556/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 100 mg tablety s predĺženým uvoľňovaním	DK/H/2942/001	65/0556/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 100 mg, tablety s prodlouženým uvolňováním	DK/H/2942/001	65/120/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 100 mg, tablety s prodlouženým	DK/H/2942/001	65/120/13-C	MYLAN IRELAND LIMITED	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
uvolňováním				
Tramadol Mylan 100 mg, tablety s prodlouženým uvolňováním	DK/H/2942/001	65/120/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 100 mg, tablety s prodlouženým uvolňováním	DK/H/2942/001	65/120/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
Tramadol Mylan 100 mg/ml Lösung zum Einnehmen	not available	6006921.00.00	KREWEL MEUSELBACH GMBH	DE
TRAMADOL Mylan 100mg/2ml SOLUÇÃO INJECTÁVEL	not available	2831188	MYLAN, LDA	PT
TRAMADOL Mylan 100mg/ml GOTAS ORAIS SOLUÇÃO	not available	3081080	MYLAN, 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
TRAMADOL Mylan 100mg/ml GOTAS ORAIS SOLUÇÃO	not available	2831089	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463245	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463229	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463336	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463237	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463344	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463351	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463245	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463229	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463336	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463237	MYLAN, LDA	PT
Tramadol Mylan 150 mg	DK/H/2942/002	5463344	MYLAN, 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
comprimidos de libertação prolongada				
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463351	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463245	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463229	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463336	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463237	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463344	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463351	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463245	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463229	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463336	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de	DK/H/2942/002	5463237	MYLAN, 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
libertação prolongada				
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463344	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	DK/H/2942/002	5463351	MYLAN, LDA	PT
Tramadol Mylan 150 mg tablety s predĺženým uvoľňovaním	DK/H/2942/002	65/0557/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 150 mg tablety s predĺženým uvoľňovaním	DK/H/2942/002	65/0557/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 150 mg tablety s predĺženým uvoľňovaním	DK/H/2942/002	65/0557/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 150 mg tablety s predĺženým uvoľňovaním	DK/H/2942/002	65/0557/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 150 mg tablety s prodlouženým uvolňováním	DK/H/2942/002	65/121/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 150 mg tablety s prodlouženým uvolňováním	DK/H/2942/002	65/121/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 150 mg tablety s prodlouženým uvolňováním	DK/H/2942/002	65/121/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 150 mg tablety s prodlouženým uvolňováním	DK/H/2942/002	65/121/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463260	MYLAN, 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
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463252	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463369	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463377	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463401	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463278	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463260	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463252	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463369	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463377	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463401	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463278	MYLAN, LDA	PT
Tramadol Mylan 200 mg	DK/H/2942/003	5463260	MYLAN, 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
comprimidos de libertação prolongada				
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463252	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463369	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463377	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463401	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463278	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463260	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463252	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463369	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463377	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	DK/H/2942/003	5463401	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de	DK/H/2942/003	5463278	MYLAN, 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
libertação prolongada				
Tramadol Mylan 200 mg tablety s predĺženým uvoľňovaním	DK/H/2942/003	65/0558/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 200 mg tablety s predĺženým uvoľňovaním	DK/H/2942/003	65/0558/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 200 mg tablety s predĺženým uvoľňovaním	DK/H/2942/003	65/0558/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 200 mg tablety s predĺženým uvoľňovaním	DK/H/2942/003	65/0558/12-S	MYLAN IRELAND LIMITED	SK
Tramadol Mylan 200 mg tablety s prodlouženým uvolňováním	DK/H/2942/003	65/122/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 200 mg tablety s prodlouženým uvolňováním	DK/H/2942/003	65/122/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 200 mg tablety s prodlouženým uvolňováním	DK/H/2942/003	65/122/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 200 mg tablety s prodlouženým uvolňováním	DK/H/2942/003	65/122/13-C	MYLAN IRELAND LIMITED	CZ
Tramadol Mylan 50 mg, comprimé effervescent	not available	NL 23973	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, comprimé effervescent	not available	NL 23973	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, comprimé effervescent	not available	NL 23973	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, comprimé effervescent	not available	NL 23973	MYLAN S.A.S	FR
Tramadol Mylan 50 mg,	not available	NL 23973	MYLAN 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
comprimé effervescent				
Tramadol Mylan 50 mg, comprimé effervescent	not available	NL 23973	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
Tramadol Mylan 50 mg, gélule	not available	NL 24060	MYLAN S.A.S	FR
TRAMADOL Mylan 50mg CÁPSULAS	not available	2830982	MYLAN, LDA	PT
TRAMADOL MYLAN GENERIQUES LP 100 mg, comprimé à libération prolongée	NL/H/0888/001	NL 33388	MYLAN S.A.S	FR
TRAMADOL MYLAN GENERIQUES LP 150 mg, comprimé à libération prolongée	NL/H/0888/002	NL 33389	MYLAN S.A.S	FR
TRAMADOL MYLAN GENERIQUES LP 200 mg, comprimé à libération prolongée	NL/H/0888/003	NL 33390	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 100 mg, comprimé à libération prolongée	DK/H/2942/001	NL 40858	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 100 mg, comprimé à	DK/H/2942/001	NL 40858	MYLAN 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
libération prolongée				
TRAMADOL MYLAN LP 100 mg, comprimé à libération prolongée	DK/H/2942/001	NL 40858	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 100 mg, comprimé à libération prolongée	DK/H/2942/001	NL 40858	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 150 mg, comprimé à libération prolongée	DK/H/2942/002	NL 40859	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 150 mg, comprimé à libération prolongée	DK/H/2942/002	NL 40859	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 150 mg, comprimé à libération prolongée	DK/H/2942/002	NL 40859	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 150 mg, comprimé à libération prolongée	DK/H/2942/002	NL 40859	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 200 mg, comprimé à libération prolongée	DK/H/2942/003	NL 40860	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 200 mg, comprimé à libération prolongée	DK/H/2942/003	NL 40860	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 200 mg, comprimé à libération prolongée	DK/H/2942/003	NL 40860	MYLAN S.A.S	FR
TRAMADOL MYLAN LP 200 mg, comprimé à libération prolongée	DK/H/2942/003	NL 40860	MYLAN S.A.S	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération	DE/H/0798/002	34009 300 141 0 3	LABORATOIRES GRÜNENTHAL 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
prolongée				
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 550 049 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 128 1 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 125 2 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 129 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 121 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 130 6 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 571 534 7 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 127 5 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN	DE/H/0798/002	34009 382 131 2 6	LABORATOIRES	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
PHARMA LP 100 mg, comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 122 3 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 120 0 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 124 6 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 132 9 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 571 533 0 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 126 9 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 300 141 1 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération	DE/H/0798/003	34009 550 049 6 7	LABORATOIRES GRÜNENTHAL 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
prolongée				
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 223 4 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 225 7 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 228 6 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 220 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 230 0 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 229 2 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 222 8 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 571 539 9 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN	DE/H/0798/003	34009 382 231 7 0	LABORATOIRES	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
PHARMA LP 150 mg, comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 224 0 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 571 538 2 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 226 3 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 221 1 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 232 3 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 300 141 2 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 550 049 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération	DE/H/0798/004	34009 382 259 9 0	LABORATOIRES GRÜNENTHAL 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
prolongée				
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 258 2 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 243 5 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 241 2 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 247 0 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 257 6 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 249 3 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 253 0 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 248 7 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN	DE/H/0798/004	34009 571 540 7 3	LABORATOIRES	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
PHARMA LP 200 mg, comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 571 541 3 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 245 8 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 254 7 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 255 3 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Tramadol Neogen 100 mg cápsulas	not available	5785043	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785035	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785068	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785050	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785076	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5784954	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785019	NEOGEN N.V.	PT
Tramadol Neogen 100	not available	5784962	NEOGEN N.V.	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
mg cápsulas				
Tramadol Neogen 100 mg cápsulas	not available	5785027	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5784970	NEOGEN N.V.	PT
Tramadol Neogen 100 mg cápsulas	not available	5785001	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784921	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784913	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784905	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784939	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784947	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784848	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784871	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784863	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784830	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784855	NEOGEN N.V.	PT
Tramadol Neogen 50 mg cápsulas	not available	5784822	NEOGEN N.V.	PT
Tramadol Normon 100 mg/2 ml solución inyectable y para perfusión EFG	not available	63.734	LABORATORIOS NORMON, S.A.	ES
Tramadol Normon 50 mg	not available	63910	LABORATORIOS NORMON,	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
cápsulas duras EFG			S.A.	
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 100 mg/2 ml Injektionslösung	not available	38543.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 50 mg/1 ml Injektionslösung	not available	38449.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 50 mg/1 ml Injektionslösung	not available	38449.00.00	PANPHARMA GMBH	DE
TRAMADOL PANPHARMA 50 mg/1 ml Injektionslösung	not available	38449.00.00	PANPHARMA 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
TRAMADOL PANPHARMA 50 mg/1 ml Injektionslösung	not available	38449.00.00	PANPHARMA GMBH	DE
Tramadol ratiopharm 50 mg cápsulas duras EFG	not available	63.472	RATIOPHARM ESPAÑA S.A.,	ES
Tramadol Retard "Hexal"	DE/H/0448/001	36837	HEXAL A/S	DK
Tramadol Retard "Hexal"	DE/H/0448/003	36839	HEXAL A/S	DK
Tramadol Retard "Hexal"	DE/H/0448/002	36838	HEXAL A/S	DK
Tramadol Retard "Actavis", depottabletter	SE/H/1717/001	40152	ACTAVIS GROUP PTC EHF.	DK
Tramadol Retard "Actavis", depottabletter	SE/H/1717/002	40153	ACTAVIS GROUP PTC EHF.	DK
Tramadol Retard "Actavis", depottabletter	SE/H/1717/003	40154	ACTAVIS GROUP PTC EHF.	DK
Tramadol retard 100 mg comprimate cu eliberare prelungită	not available	4146/2011/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Retard 150 mg comprimate cu eliberare prelungită	not available	6137/2014/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Retard 200 mg comprimate cu eliberare prelungită	not available	6138/2014/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Retard Actavis 100 mg depottabletter	SE/H/1717/001	24560	ACTAVIS GROUP PTC EHF.	SE
Tramadol Retard Actavis 100 mg tablety s predĺženým uvoľňovaním	SE/H/1717/001	65/0216/07-S	ACTAVIS GROUP PTC EHF.	SK
Tramadol Retard Actavis 150 mg depottabletter	SE/H/1717/002	24561	ACTAVIS GROUP PTC EHF.	SE
Tramadol Retard Actavis 150 mg tablety s predĺženým uvoľňovaním	SE/H/1717/002	65/0217/07-S	ACTAVIS GROUP PTC EHF.	SK
Tramadol Retard Actavis	SE/H/1717/003	24562	ACTAVIS GROUP PTC EHF.	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
200 mg depottabletter				
Tramadol Retard Actavis 200 mg tablety s predĺženým uvoľňovaním	SE/H/1717/003	65/0218/07-S	ACTAVIS GROUP PTC EHF.	SK
Tramadol retard Andrómaco 100 mg comprimidos de liberación prolongada EFG	not available	83727	ANDRÓMACO PHARMA S.L.	ES
Tramadol retard Andrómaco 150 mg comprimidos de liberación prolongada EFG	not available	83725	ANDRÓMACO PHARMA S.L.	ES
Tramadol retard Andrómaco 200 mg comprimidos de liberación prolongada EFG	not available	83728	ANDRÓMACO PHARMA S.L.	ES
Tramadol retard Andrómaco 50 mg comprimidos de liberación prolongada EFG	not available	83726	ANDRÓMACO PHARMA S.L.	ES
Tramadol Retard Aurovitas Spain 100 mg comprimidos de liberación prolongada EFG	NL/H/0539/001	72.427	AUROVITAS SPAIN,S.A.U.	ES
Tramadol Retard Aurovitas Spain 150 mg comprimidos de liberación prolongada EFG	NL/H/0539/002	71.710	AUROVITAS SPAIN,S.A.U.	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
Tramadol Retard Aurovitas Spain 200 mg comprimidos de liberación prolongada EFG	NL/H/0539/003	72.430	AUROVITAS SPAIN,S.A.U.	ES
Tramadol retard Combix 100 mg comprimidos de liberación prolongada EFG	not available	74.252	LABORATORIOS COMBIX, S.L.U.	ES
Tramadol retard Combix 150 mg comprimidos de liberación prolongada EFG	not available	74.253	LABORATORIOS COMBIX, S.L.U.	ES
Tramadol retard Combix 200 mg comprimidos de liberación prolongada EFG	not available	74.254	LABORATORIOS COMBIX, S.L.U.	ES
Tramadol Retard EG 100 mg comprimés à libération prolongée	NL/H/0888/001	BE300203	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg comprimés à libération prolongée	NL/H/0888/001	BE300212	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg comprimés à libération prolongée	NL/H/0888/001	BE300221	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg comprimés à libération prolongée	NL/H/0888/001	0019/08060054	EUROGENERICS N.V./S.A.	LU
Tramadol Retard EG 100 mg Retardtabletten	NL/H/0888/001	BE300212	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg Retardtabletten	NL/H/0888/001	BE300221	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100	NL/H/0888/001	BE300203	EUROGENERICS 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
mg Retardtabletten				
Tramadol Retard EG 100 mg tabletten met verlengde afgifte	NL/H/0888/001	BE300203	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg tabletten met verlengde afgifte	NL/H/0888/001	BE300212	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg tabletten met verlengde afgifte	NL/H/0888/001	BE300221	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg comprimés à libération prolongée	NL/H/0888/002	BE300237	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg comprimés à libération prolongée	NL/H/0888/002	BE300246	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg comprimés à libération prolongée	NL/H/0888/002	BE300255	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg comprimés à libération prolongée	NL/H/0888/002	0019/08060056	EUROGENERICS N.V./S.A.	LU
Tramadol Retard EG 150 mg Retardtabletten	NL/H/0888/002	BE300246	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg Retardtabletten	NL/H/0888/002	BE300237	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg Retardtabletten	NL/H/0888/002	BE300255	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg tabletten met verlengde afgifte	NL/H/0888/002	BE300237	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 150 mg tabletten met verlengde afgifte	NL/H/0888/002	BE300246	EUROGENERICS 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
Tramadol Retard EG 150 mg tabletten met verlengde afgifte	NL/H/0888/002	BE300255	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg comprimés à libération prolongée	NL/H/0888/003	BE300264	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg comprimés à libération prolongée	NL/H/0888/003	BE300273	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg comprimés à libération prolongée	NL/H/0888/003	BE300282	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg comprimés à libération prolongée	NL/H/0888/003	0019/08060055	EUROGENERICS N.V./S.A.	LU
Tramadol Retard EG 200 mg Retardtabletten	NL/H/0888/003	BE300273	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg Retardtabletten	NL/H/0888/003	BE300264	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg Retardtabletten	NL/H/0888/003	BE300282	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg tabletten met verlengde afgifte	NL/H/0888/003	BE300282	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg tabletten met verlengde afgifte	NL/H/0888/003	BE300273	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 200 mg tabletten met verlengde afgifte	NL/H/0888/003	BE300264	EUROGENERICS N.V./S.A.	BE
Tramadol Retard Hexal 100 mg depottablett	DE/H/0448/001	21235	HEXAL A/S	SE
Tramadol Retard HEXAL 100 mg depottabletti	DE/H/0448/001	19650	HEXAL 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
Tramadol Retard Hexal 150 mg depottablett	DE/H/0448/002	21236	HEXAL A/S	SE
Tramadol Retard Hexal 200 mg depottablett	DE/H/0448/003	21237	HEXAL A/S	SE
Tramadol Retard HEXAL 200 mg depottabletti	DE/H/0448/003	19651	HEXAL A/S	FI
Tramadol Retard Krka 100 mg depottabletter	EE/H/0262/001	57398	KRKA, D.D., NOVO MESTO	SE
Tramadol Retard Krka 150 mg depottabletter	EE/H/0262/002	57399	KRKA, D.D., NOVO MESTO	SE
Tramadol Retard Krka 200 mg depottabletter	EE/H/0262/003	57400	KRKA, D.D., NOVO MESTO	SE
Tramadol Retard Medartuum 100 mg depottabletter	NL/H/0889/001	24429	MEDARTUUM MEDICAL AB	SE
Tramadol Retard Medartuum 150 mg depottabletter	NL/H/0889/002	24430	MEDARTUUM MEDICAL AB	SE
Tramadol Retard Medartuum 200 mg depottabletter	NL/H/0889/003	24431	MEDARTUUM MEDICAL AB	SE
Tramadol Retard Mylan 100 mg comprimidos de liberación prolongada EFG.	DK/H/2942/001	76572	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan 100 mg comprimidos de liberación prolongada EFG.	DK/H/2942/001	76572	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan 100 mg comprimidos de liberación prolongada EFG.	DK/H/2942/001	76572	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan	DK/H/2942/001	76572	MYLAN PHARMACEUTICALS	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
100 mg comprimidos de liberación prolongada EFG.			S.L.	
Tramadol Retard Mylan 150 mg comprimidos de liberación prolongada EFG.	DK/H/2942/002	76573	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan 150 mg comprimidos de liberación prolongada EFG.	DK/H/2942/002	76573	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan 150 mg comprimidos de liberación prolongada EFG.	DK/H/2942/002	76573	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard Mylan 150 mg comprimidos de liberación prolongada EFG.	DK/H/2942/002	76573	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 200 mg comprimidos de liberación prolongada EFG	DK/H/2942/003	76574	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 200 mg comprimidos de liberación prolongada EFG	DK/H/2942/003	76574	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 200 mg comprimidos de liberación prolongada EFG	DK/H/2942/003	76574	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 200 mg comprimidos de liberación prolongada	DK/H/2942/003	76574	MYLAN PHARMACEUTICALS S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFG				
Tramadol retard NORMON 100 mg comprimidos de liberación prolongada EFG	not available	76122	LABORATORIOS NORMON, S.A.	ES
Tramadol retard NORMON 150 mg comprimidos de liberación prolongada EFG	not available	76124	LABORATORIOS NORMON, S.A.	ES
Tramadol retard NORMON 200 mg comprimidos de liberación prolongada EFG	not available	76125	LABORATORIOS NORMON, S.A.	ES
Tramadol retard ratiopharm 100 mg comprimidos de liberación prolongada EFG	not available	77217	RATIOPHARM ESPAÑA S.A.	ES
Tramadol retard ratiopharm 150 mg comprimidos de liberación prolongada EFG	not available	77218	RATIOPHARM ESPAÑA S.A.	ES
Tramadol retard ratiopharm 200 mg comprimidos de liberación prolongada EFG	not available	77219	RATIOPHARM ESPAÑA S.A.	ES
Tramadol Retard Sandoz 150 mg depottabletti	NL/H/0483/002	19304	SANDOZ A/S	FI
Tramadol retard STADA	not available	77.200	LABORATORIO STADA, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
100 mg comprimidos de liberación prolongada EFG				
Tramadol retard STADA 150 mg comprimidos de liberación prolongada EFG	not available	77.198	LABORATORIO STADA, S.L.	ES
Tramadol retard STADA 200 mg comprimidos de liberación prolongada EFG	not available	77.202	LABORATORIO STADA, S.L.	ES
Tramadol retard Teva 100 mg comprimidos de liberación prolongada EFG	not available	77206	TEVA PHARMA S.L.U.,	ES
Tramadol retard Teva 150 mg comprimidos de liberación prolongada EFG	not available	77232	TEVA PHARMA S.L.U.	ES
Tramadol retard Teva 200 mg comprimidos de liberación prolongada EFG	not available	77231	TEVA PHARMA S.L.U.	ES
Tramadol Sandoz 100 mg Brausetabletten	not available	39935.00.00	HEXAL AG	DE
Tramadol Sandoz 100 mg, tabletten met verlengde afgifte	DE/H/0288/001	BE235557	SANDOZ N.V.	BE
Tramadol Sandoz 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE279325	SANDOZ N.V.	BE
Tramadol Sandoz 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE279343	SANDOZ 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
Tramadol Sandoz 150 mg, tabletten met verlengde afgifte	DE/H/0288/002	BE235566	SANDOZ N.V.	BE
Tramadol Sandoz 200 mg, tabletten met verlengde afgifte	DE/H/0288/003	BE235575	SANDOZ N.V.	BE
Tramadol Sandoz 50 mg harde capsules	not available	BE254344	SANDOZ N.V.	BE
TRAMADOL SANDOZ 50 mg, comprimé	not available	34009 351 065 8 2	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	34009 351 066 4 3	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	34009 351 068 7 2	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	34009 368 193 4 4	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 383 986 1 8	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 383 987 8 6	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 383 988 4 7	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 383 989 0 8	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé	not available	34009 385 524 5 4	SANDOZ	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
pelliculé à libération prolongée				
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 385 525 1 5	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 572 785 3 3	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 572 787 6 2	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 383 990 9 7	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 383 991 5 8	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 572 175 0 1	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 572 176 7 9	SANDOZ	FR
Tramadol Sandoz Retard 200 mg tablety s prodlouženým uvolňováním	DE/H/0576/001	65/392/06-C	SANDOZ GMBH	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
Tramadol Sandoz UNO 200 mg Retardtabletten	DE/H/0576/001	44610.02.00	HEXAL AG	DE
Tramadol SOPHARMA 50 mg/ml oldatos injekcio	not available	OGYI-T-22657/02	SOPHARMA AD	HU
Tramadol SOPHARMA 50 mg/ml oldatos injekció	not available	OGYI-T-22657/01	SOPHARMA AD	HU
TRAMADOL SR ZENTIVA 100 mg comprimate cu eliberare prelungită	not available	8763/2016/04	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 100 mg comprimate cu eliberare prelungită	not available	8763/2016/03	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 100 mg comprimate cu eliberare prelungită	not available	8763/2016/02	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 100 mg comprimate cu eliberare prelungită	not available	8763/2016/01	ZENTIVA, A.S.	RO
Tramadol Stada 50 mg cápsulas duras EFG	not available	64.215	LABORATORIO STADA, S.L.	ES
Tramadol STADA 50 mg kapslar, hårda	DE/H/0282/001	17807	STADA ARZNEIMITTEL AG	SE
Tramadol STADA® 100 mg Injektionslösung	DE/H/0282/004	25036.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 100 mg Retardtabletten	AT/H/0117/001	51845.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 100 mg Zäpfchen	DE/H/0282/003	25037.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 100 mg/ml Tropfen zum Einnehmen, Lösung	DE/H/0282/002	25038.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 150 mg Retardtabletten	AT/H/0117/002	51845.01.00	STADAPHARM GMBH	DE
Tramadol STADA® 200	AT/H/0117/003	51845.02.00	STADAPHARM 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
mg Retardtabletten				
Tramadol STADA® 200 mg Retardtabletten	AT/H/0117/003	51845.02.00	STADAPHARM GMBH	DE
Tramadol STADA® 200 mg Retardtabletten	AT/H/0117/003	51845.02.00	STADAPHARM GMBH	DE
Tramadol STADA® 200 mg Retardtabletten	AT/H/0117/003	51845.02.00	STADAPHARM GMBH	DE
Tramadol STADA® 50 mg Hartkapseln	DE/H/0282/001	25035.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 50 mg Tabs Tabletten zur Herstellung einer Lösung zum Einnehmen	not available	37814.00.00	STADAPHARM GMBH	DE
TRAMADOL SYNTEZA 100 mg/ml krople doustne	not available	R/2947	SYNTEZA SP. Z O.O.	PL
TRAMADOL SYNTEZA 50 mg kapsułki twarde	not available	R/2946	SYNTEZA SP. Z O.O.	PL
Tramadol TAD 100 mg comprimidos de liberación prolongada EFG	EE/H/0269/001	83816	TAD PHARMA GMBH	ES
Tramadol TAD 100 mg comprimidos de libertação prolongada	EE/H/0262/001	5760103	TAD PHARMA GMBH	PT
Tramadol TAD 100 mg comprimidos de libertação prolongada	EE/H/0262/001	5760111	TAD PHARMA GMBH	PT
Tramadol TAD 100 mg Injektions-/Infusionslösung	HR/H/0104/002	99124.00.00	TAD PHARMA GMBH	DE
Tramadol TAD 100 mg/2 ml solution for injection/infusion	MT/H/0260/001	MA982/01502	TAD PHARMA GMBH	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
Tramadol TAD 100 mg/ml oral drops, solution	MT/H/0272/001	MA982/01601	TAD PHARMA GMBH	MT
Tramadol TAD 150 mg comprimidos de liberación prolongada EFG	EE/H/0269/002	83817	TAD PHARMA GMBH	ES
Tramadol TAD 150 mg comprimidos de libertação prolongada	EE/H/0262/002	5760129	TAD PHARMA GMBH	PT
Tramadol TAD 150 mg comprimidos de libertação prolongada	EE/H/0262/002	5760137	TAD PHARMA GMBH	PT
Tramadol TAD 200 mg comprimidos de liberación prolongada EFG	EE/H/0269/003	83818	TAD PHARMA GMBH	ES
Tramadol TAD 200 mg comprimidos de libertação prolongada	EE/H/0262/003	5760145	TAD PHARMA GMBH	PT
Tramadol TAD 200 mg comprimidos de libertação prolongada	EE/H/0262/003	5760152	TAD PHARMA GMBH	PT
Tramadol TAD 50 mg Injektions-/Infusionslösung	HR/H/0104/002	98646.00.00	TAD PHARMA GMBH	DE
Tramadol TAD 50 mg/ml solution for injection/infusion	MT/H/0260/001	MA982/01501	TAD PHARMA GMBH	MT
Tramadol Tarbis 100 mg comprimidos de liberación prolongada EFG	not available	65.800	TARBIS FARMA, S.L.	ES
Tramadol Tarbis 150 mg	not available	65.801	TARBIS FARMA, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de liberación prolongada EFG				
Tramadol Tarbis 200 mg comprimidos de liberación prolongada EFG	not available	65.408	TARBIS FARMA, S.L.	ES
TRAMADOL TEVA 50 mg, comprimé	not available	NL25151	TEVA SANTÉ	FR
TRAMADOL TEVA L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	NL38989	TEVA SANTÉ	FR
TRAMADOL TEVA L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	NL39006	TEVA SANTÉ	FR
TRAMADOL TEVA L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	NL39007	TEVA SANTÉ	FR
TRAMADOL TEVA SANTE 50 mg, gélule	not available	NL25606	TEVA SANTÉ	FR
Tramadol UNO Sandoz 200 mg tabletten met verlengde afgifte	DE/H/0576/001	BE296651	SANDOZ N.V.	BE
Tramadol VIR 100 mg/ml gotas orales en solución EFG	not available	72399	INDUSTRIA QUÍMICA Y FARMACEÚTICA VIR, S.A.	ES
Tramadol Vitabalans 50 mg tablete	FI/H/0779/001	H/13/01554/004	VITABALANS OY	SI
Tramadol Vitabalans 50 mg tablete	FI/H/0779/001	H/13/01554/003	VITABALANS OY	SI
Tramadol Vitabalans 50	FI/H/0779/001	H/13/01554/002	VITABALANS OY	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
mg tablete				
Tramadol Vitabalans 50 mg tablete	FI/H/0779/001	H/13/01554/001	VITABALANS OY	SI
Tramadol Vitabalans 50 mg tabletes	FI/H/0779/001	12-0309	VITABALANS OY	LV
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/001	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/002	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/003	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/004	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/005	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/006	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/007	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/008	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/009	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/010	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/011	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/012	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletės	FI/H/0779/001	LT/1/13/3221/013	VITABALANS OY	LT
Tramadol Vitabalans 50 mg tabletit	FI/H/0779/001	30017	VITABALANS OY	FI
Tramadol Vitabalans 50	FI/H/0779/001	OGYI-T-22315/01	VITABALANS OY	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
mg tabletta				
Tramadol Vitabalans 50 mg tabletta	FI/H/0779/001	OGYI-T-22315/02	VITABALANS OY	HU
Tramadol Vitabalans 50 mg tabletta	FI/H/0779/001	OGYI-T-22315/03	VITABALANS OY	HU
Tramadol Vitabalans 50 mg tabletta	FI/H/0779/001	OGYI-T-22315/04	VITABALANS OY	HU
Tramadol Vitabalans 50 mg tabletter	FI/H/0779/001	11-8528	VITABALANS OY	NO
Tramadol Vitabalans 50 mg tabletter	FI/H/0779/001	46613	VITABALANS OY	SE
Tramadol Vitabalans 50 mg tablety	FI/H/0779/001	65/045/13-C	VITABALANS OY	CZ
Tramadol Vitabalans 50 mg tablety	FI/H/0779/001	65/0395/13-S	VITABALANS OY	SK
Tramadol Vitabalans, 50 mg tabletid	FI/H/0779/001	796212	VITABALANS OY	EE
Tramadol Vitabalans, 50 mg, tabletki	FI/H/0779/001	20718	VITABALANS OY	PL
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 276 0 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 286 6 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 277 7 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 281 4 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 571 545 9 2	LABORATOIRES GRÜNENTHAL 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
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 283 7 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 278 3 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 284 3 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 571 546 5 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 275 4 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 280 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 282 0 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 287 2 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 100 mg, comprimé à libération prolongée	DE/H/1093/002	34009 382 274 8 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 294 9 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 299 0 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP	DE/H/1093/003	34009 382 300 9 3	LABORATOIRES	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
150 mg, comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 298 4 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 290 3 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 288 9 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 293 2 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 297 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 296 1 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 571 548 8 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 571 547 1 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 289 5 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 292 6 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 150 mg, comprimé à	DE/H/1093/003	34009 382 295 5 4	LABORATOIRES GRÜNENTHAL 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
libération prolongée				
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 301 5 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 302 1 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 306 7 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 311 0 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 571 549 4 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 309 6 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 303 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 571 550 2 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 304 4 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 313 3 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 307 3 4	LABORATOIRES GRÜNENTHAL 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
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 312 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 310 4 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 305 0 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 272 5 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 265 9 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 269 4 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 571 544 2 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 263 6 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 270 2 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 268 8 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 271 9 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP	DE/H/1093/001	34009 382 261 3 3	LABORATOIRES	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
50 mg, comprimé à libération prolongée			GRÜNENTHAL S.A.S.	
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 267 1 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 571 543 6 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 266 5 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 273 1 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 264 2 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 126 9 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 127 5 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 128 1 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 129 8 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 131 2 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 492 7 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 493 3 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 132 9 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50	UK/H/0380/001	34009 357 133 5 3	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, gélule				
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 134 1 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 135 8 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 136 4 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 137 0 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 495 6 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 496 2 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 576 208 0 6	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 130 6 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 126 9 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 127 5 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 128 1 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 129 8 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 131 2 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 492 7 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 493 3 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 132 9 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50	UK/H/0380/001	34009 357 133 5 3	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, gélule				
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 134 1 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 135 8 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 136 4 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 137 0 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 495 6 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 563 496 2 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 576 208 0 6	ZYDUS FRANCE	FR
TRAMADOL ZYDUS 50 mg, gélule	UK/H/0380/001	34009 357 130 6 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 374 901 7 7	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 374 902 3 8	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 374 899 2 8	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 374 901 7 7	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P.	not available	34009 374 902 3 8	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
100 mg, comprimé pelliculé à libération prolongée				
TRAMADOL ZYDUS L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	34009 374 899 2 8	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 144 7 8	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 174 3 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 176 6 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 575 107 6 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 575 108 2 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 173 7 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération	not available	34009 394 144 7 8	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 174 3 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 176 6 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 575 107 6 3	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 575 108 2 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	34009 394 173 7 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 177 2 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 179 5 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 180 3 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P.	not available	34009 575 109 9 2	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
200 mg, comprimé pelliculé à libération prolongée.				
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 575 110 7 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 178 9 9	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 177 2 1	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 179 5 0	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 394 180 3 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 575 109 9 2	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération prolongée.	not available	34009 575 110 7 4	ZYDUS FRANCE	FR
TRAMADOL ZYDUS L.P. 200 mg, comprimé pelliculé à libération	not available	34009 394 178 9 9	ZYDUS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée.				
Tramadol-hameln 100 mg	ENR2138540	35342.01.00	HAMELN PHARMA GMBH	DE
Tramadolhydrochlorid Actavis 150 mg Retardtabletten	SE/H/1717/002	1-26986	ACTAVIS GROUP PTC EHF.	AT
Tramadolhydrochlorid Actavis 200 mg Retardtabletten	SE/H/1717/003	1-26987	ACTAVIS GROUP PTC EHF.	AT
Tramadolhydrochlorid G.L. 100 mg-Ampullen	not available	1-21803	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid G.L. 100 mg-Zäpfchen	not available	1-21809	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid G.L. 50 mg-Ampullen	not available	1-21801	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid G.L. 50 mg-Filmtabletten	not available	1-21805	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid G.L.-Tropfen	not available	1-21807	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid Lannacher retard 100 mg-Filmtabletten	AT/H/0118/001	1-24188	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid Lannacher retard 100 mg-Filmtabletten	AT/H/0118/001	1070/03/02/0014	LANNACHER HEILMITTEL GES.M.B.H.,	LU
Tramadolhydrochlorid Lannacher retard 150 mg-Filmtabletten	AT/H/0118/002	1-24186	G.L. PHARMA GMBH	AT
Tramadolhydrochlorid Lannacher retard 150 mg-Filmtabletten	AT/H/0118/002	1070/03/02/0016	LANNACHER HEILMITTEL GES.M.B.H.,	LU
Tramadolhydrochlorid Lannacher retard 200 mg-Filmtabletten	AT/H/0118/003	1-24187	G.L. 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
Tramadolhydrochlorid Lannacher retard 200 mg-Filmtabletten	AT/H/0118/003	1070/03/02/0015	LANNACHER HEILMITTEL GES.M.B.H.,	LU
Tramadolhydrochloride capsules, capsules 50 mg	not available	RVG 18941	MYLAN B.V.	NL
Tramadolhydrochloride capsules, capsules 50 mg	not available	RVG 18941	MYLAN B.V.	NL
Tramadolis SANITAS 50 mg/ml injekcinis ar infuzinis tirpalas	not available	LT/1/2000/1577/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Tramadolis SANITAS 50 mg/ml injekcinis ar infuzinis tirpalas	not available	LT/1/2000/1577/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
TRAMADOLO ABC 100 mg/ml gocce orali, soluzione.	not available	037941010	ABC FARMACEUTICI S.P.A.	IT
TRAMADOLO ARISTO, 100 mg supposte.	not available	035918059	ARISTO PHARMA GMBH (ART 57)	IT
TRAMADOLO ARISTO, 100 mg/2 ml soluzione iniettabile.	not available	035918034	ARISTO PHARMA GMBH (ART 57)	IT
TRAMADOLO ARISTO, 100mg/ml gocce orali, soluzione.	not available	035918046	ARISTO PHARMA GMBH (ART 57)	IT
TRAMADOLO ARISTO, 50 mg capsule rigide	not available	035918010	ARISTO PHARMA GMBH (ART 57)	IT
TRAMADOLO ARISTO, 50 mg/ml soluzione iniettabile.	not available	035918022	ARISTO PHARMA GMBH (ART 57)	IT
Tramadolo EG 100 mg soluzione iniettabile	DE/H/0282/004	035847096	EG S.P.A.	IT
Tramadolo EG 100 mg	DE/H/0282/004	035847084	EG 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
soluzione iniettabile				
Tramadolo EG 100 mg soluzione iniettabile	DE/H/0282/004	035847108	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847060	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847021	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847058	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847072	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847019	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847033	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847045	EG S.P.A.	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847262	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847185	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847197	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847247	EG S.P.A.	IT
Tramadolo EG 50 mg	DE/H/0282/001	035847211	EG 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
capsule rigide				
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847223	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847235	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847250	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847110	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847146	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847122	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847209	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847159	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847173	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847134	EG S.P.A.	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847161	EG S.P.A.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133042	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133067	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133079	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio	NL/H/0538/001	037133081	GERMED PHARMA 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
prolungato				
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133093	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133105	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133117	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133129	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133319	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133321	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133333	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133345	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133358	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133360	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133372	GERMED PHARMA 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
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133384	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133396	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133408	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133055	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 100 mg compresse a rilascio prolungato	NL/H/0538/001	037133016	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133143	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133156	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133168	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133170	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133182	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133194	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150	NL/H/0538/002	037133206	GERMED PHARMA 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
mg compresse a rilascio prolungato				
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133218	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133410	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133422	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133434	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133446	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133459	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133461	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133473	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133485	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133497	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio	NL/H/0538/002	037133509	GERMED PHARMA 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
prolungato				
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133131	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 150 mg compresse a rilascio prolungato	NL/H/0538/002	037133028	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133232	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133244	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133257	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133269	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133271	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133283	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133295	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133307	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133511	GERMED PHARMA 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
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133523	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133535	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133547	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133550	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133562	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133574	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133586	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133598	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133600	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133220	GERMED PHARMA S.R.L.	IT
Tramadolo GERMED 200 mg compresse a rilascio prolungato	NL/H/0538/003	037133030	GERMED PHARMA S.R.L.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697011	SANDOZ 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
Tramadolo HCl Sandoz	NL/H/0483/001	036697074	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697023	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697047	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697035	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697050	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697062	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697086	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697098	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697100	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697112	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697124	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697136	SANDOZ 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
Tramadolo HCl Sandoz 100 mg, compresse a rilascio prolungato	NL/H/0483/001	036697148	SANDOZ S.P.A.	IT
Tramadolo Hexal 100 mg/2 ml soluzione iniettabile.	not available	033998030	SANDOZ S.P.A.	IT
TRAMADOLO HEXAL 100 mg/ml gocce orali, soluzione	not available	033998055	SANDOZ S.P.A.	IT
Tramadolo Hexal 50 mg capsule rigide.	not available	033998016	SANDOZ S.P.A.	IT
TRAMADOLO HEXAL AG 100 mg/2 ml soluzione iniettabile	not available	033531029	HEXAL AG	IT
TRAMADOLO HEXAL AG 100 mg/ml gocce orali, soluzione	not available	033531017	HEXAL AG	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569011	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569023	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569035	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569047	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569050	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569062	KRKA, D.D., NOVO MESTO	IT
Tramadolo Krka 50 mg capsule rigide	HR/H/0104/003	A.I.C. N. 045569074	KRKA, D.D., NOVO MESTO	IT
TRAMADOLO S.A.L.F. 100 mg/2 ml soluzione iniettabile	not available	044718029	S.A.L.F. SPA LABORATORIO FARMACOLOGICO	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
TRAMADOLO S.A.L.F. 50 mg/ml soluzione iniettabile	not available	044718017	S.A.L.F. SPA LABORATORIO FARMACOLOGICO	IT
Tramadolor 100 injekt, 100 mg/2 ml Injektionslösung	not available	6137845.01.00	HEXAL AG	DE
Tramadolor 100 mg - Ampullen	not available	1-23700	HEXAL PHARMA GMBH	AT
Tramadolor 100 mg / ml Lösung, Tropfen zum Einnehmen, Lösung	not available	6301608.00.00	HEXAL AG	DE
Tramadolor 100 mg Brause, Brausetabletten	not available	37201.01.00	HEXAL AG	DE
Tramadolor 100 mg ID, Retardtabletten	DE/H/0448/001	11558.00.00	HEXAL AG	DE
Tramadolor 100 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/01	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 100 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/02	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 100 mg/ml - Tropfen	not available	1-23701	HEXAL PHARMA GMBH	AT
Tramadolor 150 mg ID, Retardtabletten	DE/H/0448/002	45017.01.00	HEXAL AG	DE
Tramadolor 150 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/09	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 150 mg módosított hatóanyagleadású	not available	OGYI-T-8179/08	SANDOZ HUNGÁRIA KFT	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
tabletta				
Tramadolor 200 mg ID, Retardtabletten	DE/H/0448/003	45017.02.00	HEXAL AG	DE
Tramadolor 200 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/06	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 200 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/05	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 50 injekt, 50 mg/ml Injektionslösung	not available	35345.00.00	HEXAL AG	DE
Tramadolor 50 mg - Kapseln	not available	1-23708	HEXAL PHARMA GMBH	AT
Tramadolor 50 mg kemény kapszula	not available	OGYI-T-8179/03	SANDOZ HUNGÁRIA KFT	HU
Tramadolor 50 mg kemény kapszula	not available	OGYI-T-8179/04	SANDOZ HUNGÁRIA KFT	HU
Tramadolor ID 200 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/011	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 200 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/013	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 200 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/012	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 100 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/002	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 100 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/003	SANDOZ PHARMACEUTICALS D.D.	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
Tramadolor ID 150 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/008	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 150 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/009	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor ID 150 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/99/0247/010	SANDOZ PHARMACEUTICALS D.D.	LT
Tramadolor Kapseln, 50 mg Hartkapseln	not available	32747.00.02	HEXAL AG	DE
Tramadolor long 100, 100 mg Hartkapseln, retardiert	not available	42433.01.00	HEXAL AG	DE
Tramadolor long 150, 150 mg Hartkapseln, retardiert	not available	42433.02.00	HEXAL AG	DE
Tramadolor long 200, 200 mg Hartkapseln, retardiert	not available	42433.03.00	HEXAL AG	DE
Tramadolor long 50, 50 mg Hartkapseln, retardiert	not available	42433.00.00	HEXAL AG	DE
Tramadolor retard 100 mg - Tabletten	DE/H/0288/001	1-24377	HEXAL PHARMA GMBH	AT
Tramadolor retard 100 mg tablete s produljenim oslobađanjem	not available	HR-H-835899381	SANDOZ D.O.O.	HR
Tramadolor retard 150 mg - Tabletten	DE/H/0288/002	1-24382	HEXAL PHARMA GMBH	AT
Tramadolor retard 150 mg tablete s produljenim oslobađanjem	not available	HR-H-008744405	SANDOZ D.O.O.	HR
Tramadolor retard 200	DE/H/0288/003	1-24383	HEXAL 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
mg - Tabletten				
Tramadolor retard 200 mg tablete s produljenim oslobađanjem	not available	HR-H-026332681	SANDOZ D.O.O.	HR
Tramadolor tabs, 50 mg Tabletten	not available	6584337.00.00	HEXAL AG	DE
Tramadol-Q® 100 mg Retardtabletten	NL/H/0888/001	67444.00.00	JUTA PHARMA GMBH	DE
Tramadol-Q® 150 mg Retardtabletten	NL/H/0888/002	67445.00.00	JUTA PHARMA GMBH	DE
Tramadol-Q® 200 mg Retardtabletten	NL/H/0888/003/MR	67446.00.00	JUTA PHARMA GMBH	DE
Tramadol-ratiopharm® 100 mg Retardtabletten	not available	39925.00.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 100 mg/2 ml Injektionslösung	not available	6025891.00.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 100 mg/ml Tropfen Tropfen zum Einnehmen, Lösung	not available	31467.00.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 150 mg Retardkapseln	not available	54390.02.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 200 mg Retardkapseln	not available	54390.03.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 50 mg Hartkapseln	not available	31518.00.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 50 mg Retardkapseln	not available	54390.00.00	RATIOPHARM GMBH	DE
Tramadol-ratiopharm® 50 mg/ml Injektionslösung	not available	32305.00.00	RATIOPHARM GMBH	DE
Tramadol-saar 100 mg Ampulle	not available	32285.00.00	MIP 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
Tramadol-saar 50 mg Ampulle	not available	35344.00.00	MIP PHARMA GMBH	DE
Tramadol-Sandoz 100 mg / ml Tropfen zum Einnehmen, Lösung	not available	30508.00.00	HEXAL AG	DE
Tramag 50	not available	10415/2017/01	MAGISTRA C&C	RO
Tramagetic OD 150 mg depottabletter	not available	99-7688	MUNDIPHARMA AS.	NO
Tramagetic OD 200 mg depottabletter	not available	99-7689	MUNDIPHARMA AS.	NO
Tramagetic OD 300 mg depottabletter	not available	99-7690	MUNDIPHARMA AS.	NO
TRAMAGETIC Once-Daily 150 mg tabletten	not available	RVG 22232	MUNDIPHARMA PHARMACEUTICALS BV	NL
TRAMAGETIC Once-Daily 200 tabletten	not available	RVG 22233	MUNDIPHARMA PHARMACEUTICALS BV	NL
TRAMAGETIC Once-Daily 300 tabletten	not available	RVG 22234	MUNDIPHARMA PHARMACEUTICALS BV	NL
TRAMAGETIC Once-Daily 400 mg tabletten	not available	RVG 22235	MUNDIPHARMA PHARMACEUTICALS BV	NL
Tramagetic Retard 100 mg depottabletter	not available	99-7782	MUNDIPHARMA AS.	NO
Tramagetic Retard 150 mg depottabletter	not available	99-7783	MUNDIPHARMA AS.	NO
Tramagetic Retard 200 mg depottabletter	not available	99-7784	MUNDIPHARMA AS.	NO
Tramagetic Retard 75 mg depottabletter	not available	99-7781	MUNDIPHARMA AS.	NO
Tramagit® 100 mg Retardtabletten	not available	39924.00.00	KREWEL MEUSELBACH GMBH	DE
Tramagit® 100 mg/ml Lösung zum Einnehmen	not available	29737.00.01	KREWEL MEUSELBACH GMBH	DE
Tramagit® 150 mg Retardtabletten	not available	79350.00.00	KREWEL MEUSELBACH 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
Tramagit® 200 mg Retardtabletten	not available	79351.00.00	KREWEL MEUSELBACH GMBH	DE
Tramagit® 50 mg Tabletten	not available	29737.00.00	KREWEL MEUSELBACH GMBH	DE
Tramagit® Tabletten 50 mg	not available	6140209.00.00	KREWEL MEUSELBACH GMBH	DE
Tramake 100mg Tablets	not available	PA 22680/3/2	GALEN PHARMA IRELAND LIMITED	IE
Tramake 100mg Tablets	not available	PA 22680/3/2	GALEN PHARMA IRELAND LIMITED	IE
Tramake 100mg Tablets	not available	PA 22680/3/2	GALEN PHARMA IRELAND LIMITED	IE
Tramake 50mg Tablets	not available	PA 22680/3/1	GALEN PHARMA IRELAND LIMITED	IE
Tramake 50mg Tablets	not available	PA 22680/3/1	GALEN PHARMA IRELAND LIMITED	IE
Tramake 50mg Tablets	not available	PA 22680/3/1	GALEN PHARMA IRELAND LIMITED	IE
Tramal 100 mg Ampullen	not available	17.690	GRÜNENTHAL GES. M.B.H.	AT
TRAMAL 100 mg čípky	not available	65/076/91-S/C	STADA ARZNEIMITTEL AG	CZ
TRAMAL 100 mg ilgstošās darbības tabletes	not available	99-0530	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg otopina za injekciju/infuziju	not available	HR-H-286762123	STADA D.O.O.	HR
TRAMAL 100 mg supozitoriji	not available	99-0529	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg tablete s podaljšanim sproščanjem	not available	H/92/01556/010	STADA ARZNEIMITTEL AG	SI
Tramal 100 mg tablete s produljenim oslobadanjem	not available	HR-H-610358597	STADA D.O.O.	HR
Tramal 100 mg tablete s produljenim	not available	HR-H-610358597	STADA 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
oslobadanjem				
Tramal 100 mg, Injektionslösung	not available	1116.00.02	GRÜNENTHAL GMBH	DE
Tramal 100 mg, Injektionslösung	not available	120/88/06/0250	GRÜNENTHAL GMBH	LU
Tramal 100 mg/1 ml perorální roztok	not available	65/077/91-S/C	STADA ARZNEIMITTEL AG	CZ
TRAMAL 100 mg/2 ml injekční roztok	not available	65/079/91-S/C	STADA ARZNEIMITTEL AG	CZ
Tramal 100 mg/2 ml solução injetável	not available	8565002	GRÜNENTHAL S.A.	PT
Tramal 100 mg/ml gotas orais, solução	not available	3081288	GRÜNENTHAL S.A.	PT
Tramal 100 mg/ml gotas orais, solução	not available	8740605	GRÜNENTHAL S.A.	PT
Tramal 100 mg/ml gotas orais, solução	not available	2710085	GRÜNENTHAL S.A.	PT
Tramal 100 mg/ml Lösung zum Einnehmen	not available	1116.00.01	GRÜNENTHAL GMBH	DE
Tramal 100 mg/ml orala droppar, lösning i doseringspumpflaskan	not available	12688	ORION OYJ	FI
Tramal 100 mg/ml oralne kapi, otopina	not available	HR-H-296852819	STADA D.O.O.	HR
Tramal 100 mg/ml oralne kapi, otopina	not available	HR-H-296852819	STADA D.O.O.	HR
Tramal 100 mg/ml oralne kapi, otopina	not available	HR-H-296852819	STADA D.O.O.	HR
Tramal 100 mg/ml peroralne kapljice, raztopina v steklenici z odmerno črpalko	not available	H/92/01556/006	STADA ARZNEIMITTEL AG	SI
TRAMAL 100 mg/ml pilieni iekšķīgai	not available	13-0057	STADA ARZNEIMITTEL AG	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
lietošanai, šķīdums pudelē ar dozēšanas sūkni				
TRAMAL 100 mg/ml pilieni iekšķīgai lietošanai, šķīdums pudelē ar pilinātāju	not available	99-0528	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg/ml tipat, liuos	not available	12688	ORION OYJ	FI
Tramal 100, oplossing voor injectie 100 mg/2 ml	not available	RVG 15510	GRÜNENTHAL B.V.	NL
TRAMAL 150 mg ilgstošās darbības tabletes	not available	99-0531	STADA ARZNEIMITTEL AG	LV
Tramal 150 mg tablete s podaljšanim sproščanjem	not available	H/92/01556/002	STADA ARZNEIMITTEL AG	SI
Tramal 150 mg tablete s podaljšanim sproščanjem	not available	H/92/01556/011	STADA ARZNEIMITTEL AG	SI
Tramal 150 mg tablete s produljenim oslobadanjem	not available	HR-H-170945872	STADA D.O.O.	HR
Tramal 150 mg tablete s produljenim oslobadanjem	not available	HR-H-170945872	STADA D.O.O.	HR
TRAMAL 200 mg ilgstošās darbības tabletes	not available	99-0532	STADA ARZNEIMITTEL AG	LV
Tramal 200 mg tablete s podaljšanim sproščanjem	not available	H/92/01556/003	STADA ARZNEIMITTEL AG	SI
Tramal 200 mg tablete s podaljšanim sproščanjem	not available	H/92/01556/012	STADA ARZNEIMITTEL AG	SI
Tramal 200 mg tablete s produljenim	not available	HR-H-505350700	STADA 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
oslobadanjem				
Tramal 200 mg tablete s produljenim oslobadanjem	not available	HR-H-505350700	STADA D.O.O.	HR
Tramal 50 mg Ampullen	not available	17692	GRÜNENTHAL GES. M.B.H.	AT
Tramal 50 mg cápsulas	not available	8565119	GRÜNENTHAL S.A.	PT
Tramal 50 mg cápsulas	not available	8565101	GRÜNENTHAL S.A.	PT
TRAMAL 50 mg cietās kapsulas	not available	99-0527	STADA ARZNEIMITTEL AG	LV
Tramal 50 mg hårda kapslar	not available	11453	ORION OYJ	FI
Tramal 50 mg kapseli, kova	not available	11453	ORION OYJ	FI
Tramal 50 mg Kapseln	not available	17.688	GRÜNENTHAL GES. M.B.H.	AT
Tramal 50 mg kapsule	not available	HR-H-722830373	STADA D.O.O.	HR
Tramal 50 mg kapsule	not available	HR-H-722830373	STADA D.O.O.	HR
Tramal 50 mg liukeneva tabletti	not available	13245	ORION OYJ	FI
Tramal 50 mg otopina za injekciju/infuziju	not available	HR-H-065187023	STADA D.O.O.	HR
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/004	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/001	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/011	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/009	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/002	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/005	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s	DE/H/0108/004	H/09/02034/013	STADA ARZNEIMITTEL AG	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
podaljšanim sproščanjem				
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/008	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/010	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/012	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/006	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/014	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/003	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg tablete s podaljšanim sproščanjem	DE/H/0108/004	H/09/02034/007	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg trde kapsule	not available	H/92/01556/004	STADA ARZNEIMITTEL AG	SI
TRAMAL 50 mg tvrdé tobolky	not available	65/075/91-S/C	STADA ARZNEIMITTEL AG	CZ
Tramal 50 mg upplösliga tabletter	not available	13245	ORION OYJ	FI
Tramal 50 mg, Injektionslösung	not available	1116.01.02	GRÜNENTHAL GMBH	DE
TRAMAL 50 mg/1 ml injekční roztok	not available	65/078/91-S/C	STADA ARZNEIMITTEL AG	CZ
Tramal 50 mg/ml injektioneste, liuos	not available	11452	ORION OYJ	FI
Tramal 50 mg/ml injektionsvätska, lösning	not available	11452	ORION OYJ	FI
Tramal 50 mg/ml raztopina za injiciranje/infundiranje	not available	H/92/01556/008	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg/ml raztopina za	not available	H/92/01556/009	STADA ARZNEIMITTEL AG	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
injiciranje/infundiranje				
TRAMAL 50 mg/ml šķīdums injekcijām	not available	96-0105	STADA ARZNEIMITTEL AG	LV
TRAMAL čapíky 100 mg	not available	65/0076/91-S	STADA ARZNEIMITTEL AG	SK
TRAMAL injekčný roztok 100 mg/2 ml	not available	65/0079/91-S	STADA ARZNEIMITTEL AG	SK
TRAMAL injekčný roztok 50 mg/ml	not available	65/0078/91-S	STADA ARZNEIMITTEL AG	SK
Tramal Kapseln, 50 mg, Hartkapseln	not available	1116.00.00	GRÜNENTHAL GMBH	DE
Tramal Kapseln, 50 mg, Hartkapseln	not available	120/88/06/0249	GRÜNENTHAL GMBH	LU
TRAMAL kvapky 100 mg/ml	not available	65/0077/91-S	STADA ARZNEIMITTEL AG	SK
Tramal long 100 mg Retardtabletten	not available	120/95/12/0413	GRÜNENTHAL GMBH	LU
Tramal long 100 mg Retardtabletten	DE/H/0798/002	37294.00.00	GRÜNENTHAL GMBH	DE
Tramal long 150 mg Retardtabletten	not available	0120/97/04/0280	GRÜNENTHAL GMBH	LU
Tramal long 150 mg Retardtabletten	DE/H/0798/003	37294.01.00	GRÜNENTHAL GMBH	DE
Tramal long 200 mg Retardtabletten	DE/H/0798/004	37294.02.00	GRÜNENTHAL GMBH	DE
Tramal long 200 mg Retardtabletten	not available	0120/97/04/0282	GRÜNENTHAL GMBH	LU
Tramal long 50 mg Retardtabletten	DE/H/0798/001	60444.00.00	GRÜNENTHAL GMBH	DE
TRAMAL ORAL DROPS, SOLUTION 100MG/ML	not available	30122/15/10-11-2016	VIANEX S.A.	GR
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2971984	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg	DE/H/0108/001	2511186	GRÜNENTHAL 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
comprimidos de libertação prolongada				
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	5419486	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg depottabletti	DE/H/0108/001	12497	GRÜNENTHAL GMBH	FI
Tramal retard 100 mg Filmtabletten	not available	1-21219	GRÜNENTHAL GES. M.B.H.	AT
TRAMAL RETARD 100 mg tablety s prodlouženým uvolňováním	not available	65/873/97-C	STADA ARZNEIMITTEL AG	CZ
TRAMAL RETARD 100 mg toimeainet prolongeeritult vabastavad tabletid	not available	263099	STADA ARZNEIMITTEL AG	EE
TRAMAL RETARD 100 mg, comprimate cu eliberare prelungită, 100 mg	not available	12323/2019/01	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 100 mg, comprimate cu eliberare prelungită, 100 mg	not available	12323/2019/02	STADA ARZNEIMITTEL AG	RO
Tramal retard 100 mg, tabletten met verlengde afgifte	DE/H/0108/001	RVG 22361	GRÜNENTHAL B.V.	NL
Tramal retard 100 mg Retardtabletten	DE/H/0108/001	37297.00.00	GRÜNENTHAL GMBH	DE
Tramal Retard 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	7862	STADA ARZNEIMITTEL AG	PL
Tramal retard 150 mg comprimidos de	DE/H/0108/002	5419585	GRÜNENTHAL 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
libertação prolongada				
Tramal retard 150 mg comprimidos de libertação prolongada	DE/H/0108/002	2972081	GRÜNENTHAL S.A.	PT
Tramal retard 150 mg comprimidos de libertação prolongada	DE/H/0108/002	2511285	GRÜNENTHAL S.A.	PT
Tramal retard 150 mg depottabletti	DE/H/0108/002	12498	GRÜNENTHAL GMBH	FI
Tramal retard 150 mg Filmtabletten	not available	1-21218	GRÜNENTHAL GES. M.B.H.	AT
TRAMAL RETARD 150 mg tablety s prodlouženým uvolňováním	not available	65/539/99-C	STADA ARZNEIMITTEL AG	CZ
TRAMAL RETARD 150 mg toimeainet prolongeeritult vabastavad tabletid	not available	263199	STADA ARZNEIMITTEL AG	EE
TRAMAL RETARD 150 mg, comprimate cu eliberare prelungită, 150 mg	not available	12324/2019/02	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 150 mg, comprimate cu eliberare prelungită, 150 mg	not available	12324/2019/01	STADA ARZNEIMITTEL AG	RO
Tramal retard 150 mg, tabletten met verlengde afgifte	DE/H/0108/002	RVG 22362	GRÜNENTHAL B.V.	NL
Tramal retard 150 mg Retardtabletten	DE/H/0108/002	37297.01.00	GRÜNENTHAL GMBH	DE
Tramal Retard 150, 150 mg, tabletki o przedłużonym uwalnianiu	not available	7863	STADA ARZNEIMITTEL AG	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
Tramal retard 200 mg comprimidos de libertação prolongada	DE/H/0108/003	5419684	GRÜNENTHAL S.A.	PT
Tramal retard 200 mg comprimidos de libertação prolongada	DE/H/0108/003	2972180	GRÜNENTHAL S.A.	PT
Tramal retard 200 mg comprimidos de libertação prolongada	DE/H/0108/003	2511384	GRÜNENTHAL S.A.	PT
Tramal retard 200 mg depottabletti	DE/H/0108/003	12499	GRÜNENTHAL GMBH	FI
Tramal retard 200 mg Filmtabletten	not available	1-21217	GRÜNENTHAL GES. M.B.H.	AT
TRAMAL RETARD 200 mg tablety s prodlouženým uvolňováním	not available	65/540/99-C	STADA ARZNEIMITTEL AG	CZ
TRAMAL RETARD 200 mg toimeainet prolongeeritult vabastavad tabletid	not available	263299	STADA ARZNEIMITTEL AG	EE
TRAMAL RETARD 200 mg, comprimate cu eliberare prelungită, 200 mg	not available	12325/2019/02	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 200 mg, comprimate cu eliberare prelungită, 200 mg	not available	12325/2019/01	STADA ARZNEIMITTEL AG	RO
Tramal retard 200 mg, tabletten met verlengde afgifte	DE/H/0108/003	RVG 22363	GRÜNENTHAL B.V.	NL
Tramal retard 200 mg, Retardtabletten	DE/H/0108/003	37297.02.00	GRÜNENTHAL GMBH	DE
Tramal Retard 200, 200	not available	7864	STADA ARZNEIMITTEL AG	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
mg, tabletki o przedłużonym uwalnianiu				
Tramal retard 50 mg depottabletti	DE/H/0108/004	22364	GRÜNENTHAL GMBH	FI
Tramal retard 50 mg Retardtabletten	DE/H/0108/004/MR	60442.00.00	GRÜNENTHAL GMBH	DE
Tramal Retard 50, 50 mg, tabletki o przedłużonym uwalnianiu	DE/H/0108/004	16764	STADA ARZNEIMITTEL AG	PL
TRAMAL retard tablety 100 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL retard tablety 150 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL retard tablety 200 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL SOLUTION FOR INJECTION 100MG/2ML AMP	not available	61457/19-08-2014	VIANEX S.A.	GR
TRAMAL SUPPOSITORY 100MG	not available	54399/06-02-2013	VIANEX S.A.	GR
Tramal Tabletten, 50 mg	not available	28705.00.00	GRÜNENTHAL GMBH	DE
Tramal Tropfen	not available	17.689	GRÜNENTHAL GES. M.B.H.	AT
Tramal Tropfen, 100 mg/ml, Lösung zum Einnehmen	not available	120/88/06/0248	GRÜNENTHAL GMBH	LU
TRAMAL tvrdé kapsuly 50 mg	not available	65/0075/91-S	STADA ARZNEIMITTEL AG	SK
Tramal Zäpfchen, 100 mg	not available	13143.00.00	GRÜNENTHAL GMBH	DE
Tramal Zäpfchen, 100 mg	not available	120/95/12/0414	GRÜNENTHAL GMBH	LU
TRAMAL, 100 mg rektaalsuposiidid	not available	142096	STADA ARZNEIMITTEL AG	EE
Tramal, 100 mg, czopki	not available	2537	STADA ARZNEIMITTEL AG	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
Tramal, 100 mg/ml, krople doustne, roztwór	not available	2539	STADA ARZNEIMITTEL AG	PL
TRAMAL, 50 mg kõvakapslid	not available	141996	STADA ARZNEIMITTEL AG	EE
Tramal, 50 mg, kapsułki, twarde	not available	2536	STADA ARZNEIMITTEL AG	PL
Tramal, 50 mg/1 ml, roztwór do wstrzykiwań	not available	2538	STADA ARZNEIMITTEL AG	PL
Tramal, capsules, hard 50 mg	not available	RVG 15511	GRÜNENTHAL B.V.	NL
Tramal, druppels voor oraal gebruik, oplossing 100 mg/ml	not available	RVG 15513	GRÜNENTHAL B.V.	NL
Tramalgic 50 mg kemény kapszula	not available	OGYI-T-6565/01	TAKEDA PHARMA KFT.	HU
Tramalgic 50 mg kemény kapszula	not available	OGYI-T-6565/02	TAKEDA PHARMA KFT.	HU
Tramalgic 50 mg kemény kapszula	not available	OGYI-T-6565/03	TAKEDA PHARMA KFT.	HU
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846017	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846029	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846031	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846043	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846056	S.F. GROUP SRL	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
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846068	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846070	S.F. GROUP SRL	IT
TRAMALIN 100 mg compresse a rilascio prolungato	AT/H/0118/001	035846082	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846094	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846106	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846118	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846120	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846132	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846144	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846157	S.F. GROUP SRL	IT
TRAMALIN 150 mg compresse a rilascio prolungato	AT/H/0118/002	035846169	S.F. GROUP SRL	IT
TRAMALIN 200 mg	AT/H/0118/003	035846171	S.F. GROUP SRL	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
compresse a rilascio prolungato				
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846183	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846195	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846207	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846219	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846245	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846221	S.F. GROUP SRL	IT
TRAMALIN 200 mg compresse a rilascio prolungato	AT/H/0118/003	035846233	S.F. GROUP SRL	IT
Tramamed 100 mg Retardtabletten	DE/H/0288/001	44607.00.00	HEXAL AG	DE
Tramamed 150 mg Retardtabletten	DE/H/0288/002	44607.01.00	HEXAL AG	DE
Tramamed 200 mg Retardtabletten	DE/H/0288/003	44607.02.00	HEXAL AG	DE
Tramastad 100 mg/2 ml Ampullen	not available	1-23281	STADA ARZNEIMITTEL GMBH	AT
Tramastad 100 mg/ml Tropfen	not available	1-23282	STADA ARZNEIMITTEL GMBH	AT
Tramastad 50 mg	not available	1-23280	STADA ARZNEIMITTEL	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
Kapseln			GMBH	
TRAMIUM 100 mg depotkapseli, kova	FI/H/0164/001	16710	LABORATOIRES SMB S.A.	FI
Tramium 100 mg gélules à libération prolongée.	FI/H/0164/001	BE249103	LABORATOIRES SMB S.A.	BE
Tramium 100 mg gélules à libération prolongée.	FI/H/0164/001	2003070040	LABORATOIRES SMB S.A.	LU
Tramium 100 mg hård depotkapsel	FI/H/0164/001	16710	LABORATOIRES SMB S.A.	FI
Tramium 100 mg Harde capsules met verlengde afgifte.	FI/H/0164/001	BE249103	LABORATOIRES SMB S.A.	BE
Tramium 100 mg Hartkapseln, retardiert	FI/H/0164/001	BE249103	LABORATOIRES SMB S.A.	BE
Tramium 100 mg Hartkapseln, retardiert	FI/H/0164/001	2003070040	LABORATOIRES SMB S.A.	LU
TRAMIUM 150 mg depotkapseli, kova	FI/H/0164/002	16711	LABORATOIRES SMB S.A.	FI
Tramium 150 mg gélules à libération prolongée.	FI/H/0164/002	BE249112	LABORATOIRES SMB S.A.	BE
Tramium 150 mg gélules à libération prolongée.	FI/H/0164/002	2003070039	LABORATOIRES SMB S.A.	LU
Tramium 150 mg hård depotkapsel	FI/H/0164/002	16711	LABORATOIRES SMB S.A.	FI
Tramium 150 mg Harde capsules met verlengde afgifte.	FI/H/0164/002	BE249112	LABORATOIRES SMB S.A.	BE
Tramium 150 mg Hartkapseln, retardiert	FI/H/0164/002	BE249112	LABORATOIRES SMB S.A.	BE
Tramium 150 mg Hartkapseln, retardiert	FI/H/0164/002	2003070039	LABORATOIRES SMB S.A.	LU
TRAMIUM 200 mg depotkapseli, kova	FI/H/0164/003	16712	LABORATOIRES SMB S.A.	FI
Tramium 200 mg gélules	FI/H/0164/003	BE249121	LABORATOIRES SMB 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
à libération prolongée.				
Tramium 200 mg gélules à libération prolongée.	FI/H/0164/003	2003070041	LABORATOIRES SMB S.A.	LU
Tramium 200 mg hård depotkapsel	FI/H/0164/003	16712	LABORATOIRES SMB S.A.	FI
Tramium 200 mg Harde capsules met verlengde afgifte.	FI/H/0164/003	BE249121	LABORATOIRES SMB S.A.	BE
Tramium 200 mg Hartkapseln, retardiert	FI/H/0164/003	BE249121	LABORATOIRES SMB S.A.	BE
Tramium 200 mg Hartkapseln, retardiert	FI/H/0164/003	2003070041	LABORATOIRES SMB S.A.	LU
Tramól-L 100 mg forðatöflur	AT/H/0117/001	IS/1/01/037/01	ACTAVIS GROUP PTC EHF.	IS
Tramól-L 150 mg forðatöflur	AT/H/0117/002	IS/1/01/037/02	ACTAVIS GROUP PTC EHF.	IS
Tramól-L 200 mg forðatöflur	AT/H/0117/003	IS/1/01/037/03	ACTAVIS GROUP PTC EHF.	IS
Tramól-L 200 mg forðatöflur	AT/H/0117/003	IS/1/01/037/03	ACTAVIS GROUP PTC EHF.	IS
Tramól-L 200 mg forðatöflur	AT/H/0117/003	IS/1/01/037/03	ACTAVIS GROUP PTC EHF.	IS
Tramól-L 200 mg forðatöflur	AT/H/0117/003	IS/1/01/037/03	ACTAVIS GROUP PTC EHF.	IS
Tramquel SR 100 mg prolonged-release hard capsules	UK/H/0301/002	PL 04569/1775	GENERICS [UK] LIMITED	UK
Tramquel SR 150 mg prolonged-release hard capsules	UK/H/0301/003	PL 04569/1776	GENERICS [UK] LIMITED	UK
Tramquel SR 200 mg prolonged-release hard capsules	UK/H/0301/004	PL 04569/1777	GENERICS [UK] LIMITED	UK
Tramquel SR 50 mg	UK/H/0301/001	PL 04569/1774	GENERICS [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
prolonged-release hard capsules				
Tramulief SR 100 mg prolonged-release tablets	NL/H/0889/001	PL 20072/0235	AMDIPHARM UK LIMITED	UK
Tramulief SR 150 mg prolonged-release tablets	NL/H/0889/002	PL 20072/0236	AMDIPHARM UK LIMITED	UK
Tramulief SR 200 mg prolonged-release tablets	NL/H/0889/003	PL 20072/0237	AMDIPHARM UK LIMITED	UK
Tramundal 50 mg Filmtabletten	not available	1-21777	MUNDIPHARMA GES.M.B.H	AT
Tramundal retard 100 mg Filmtabletten	not available	1-22187	MUNDIPHARMA GES.M.B.H	AT
Tramundal retard 150 mg Filmtabletten Wirkstoff: Tramadolhydrochlorid	AT/H/0120/001	1-22188	MUNDIPHARMA GES.M.B.H	AT
Tramundal retard 200 mg Filmtabletten Wirkstoff: Tramadolhydrochlorid	AT/H/0120/002	1-22189	MUNDIPHARMA GES.M.B.H	AT
Tramundal Tropfen 100 mg	not available	1-22186	MUNDIPHARMA GES.M.B.H	AT
Tramundin 100 mg tablete s podaljšanim sproščanjem	not available	H/00/01557/001	MUNDIPHARMA GES.M.B.H	SI
Tramundin 100mg/ml Lösung zum Einnehmen	not available	25526.00.00	MUNDIPHARMA GMBH	DE
Tramundin retard 100 mg tablete s produljenim oslobađanjem	not available	HR-H-002424661	MUNDIPHARMA GES.M.B.H	HR
TRAMUNDIN RETARD 100mg	not available	65/540/00-C	MUNDIPHARMA GES.M.B.H	CZ
Tramundin retard 150 mg Retardtabletten	AT/H/0120/001	34105.01.00	MUNDIPHARMA 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
Wirkstoff: Tramadolhydrochlorid				
Tramundin retard 200 mg Retardtabletten Wirkstoff: Tramadolhydrochlorid	AT/H/0120/002	34105.02.00	MUNDIPHARMA GMBH	DE
Tramundin, 100 mg, tabletki powlekane o przedłużonym uwalnianiu	not available	9474	MUNDIPHARMA A/S	PL
Tramundin® 50 mg Filmtabletten Wirkstoff: Tramadolhydrochlorid	not available	6391213.00.00	MUNDIPHARMA GMBH	DE
Tramundin® retard 100 mg Retardtabletten Wirkstoff: Tramadolhydrochlorid	not available	34105.00.00	MUNDIPHARMA GMBH	DE
TRAM-U-RON OD 100 mg cápsulas de libertação prolongada	FI/H/0164/001	4392882	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cápsulas de libertação prolongada	FI/H/0164/001	4392981	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cápsulas de libertação prolongada	FI/H/0164/001	4393088	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cápsulas de libertação prolongada	FI/H/0164/001	4393187	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg Cápsulas de libertação prolongada	FI/H/0164/002	4393286	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg Cápsulas de libertação prolongada	FI/H/0164/002	4393385	BENE FARMACÊUTICA, 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
TRAM-U-RON OD 150 mg Cápsulas de libertação prolongada	FI/H/0164/002	4393484	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg Cápsulas de libertação prolongada	FI/H/0164/002	4393583	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg Cápsulas de libertação prolongada	FI/H/0164/003	4393682	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg Cápsulas de libertação prolongada	FI/H/0164/003	4393781	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg Cápsulas de libertação prolongada	FI/H/0164/003	4393880	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg Cápsulas de libertação prolongada	FI/H/0164/003	4393989	BENE FARMACÊUTICA, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3381688	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3381787	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3381886	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382082	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3381985	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg	UK/H/0306/001	3768587	BGP PRODUCTS	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
Comprimido de libertação prolongada			UNIPESSOAL, LDA.	
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382181	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382280	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382389	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382488	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382587	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382686	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382785	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382884	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3382983	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383189	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de	UK/H/0306/001	3383080	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
libertação prolongada				
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3827086	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383288	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383387	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383486	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383585	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383783	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 150 mg Comprimido de libertação prolongada	UK/H/0306/001	3383882	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/002	3383981	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/002	3384088	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/002	3384187	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃｯﾃ｣	UK/H/0306/002	3384286	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
｣o prolongada				
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384385	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3768686	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384484	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3678182	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384682	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384880	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384781	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3384989	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385085	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385184	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ	UK/H/0306/002	3385283	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
｣o prolongada				
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385580	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385382	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3827185	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385689	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385580	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385887	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385986	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3385788	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 200 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/002	3386083	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3386281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ	UK/H/0306/003	3386182	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
｣o prolongada				
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386984	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3387081	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386687	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386786	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386588	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3387289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3387180	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386885	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3386489	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ｣o prolongada	UK/H/0306/003	3768785	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃｧﾃ	UK/H/0306/003	3386380	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
｣o prolongada				
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387487	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3388287	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387586	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387685	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387388	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3827284	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387883	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3387982	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ ｣o prolongada	UK/H/0306/003	3388089	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 300 mg Comprimido de libertaﾃﾃﾃ	UK/H/0306/003	3388188	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
⌋o prolongada				
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388485	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388683	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388386	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388584	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388782	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388881	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3388980	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3768884	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389285	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389483	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389384	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389582	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389681	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389780	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3390085	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3827383	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertação prolongada	UK/H/0306/004	3389988	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/004	3389889	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/004	3390184	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/004	3390283	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/004	3390382	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaﾃｯﾃ｣o prolongada	UK/H/0306/004	3390481	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 400 mg Comprimido de libertaテァテ」o prolongada	UK/H/0306/004	3389087	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX Long 400 mg Comprimido de libertaテァテ」o prolongada	UK/H/0306/004	3389186	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Travex retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	42429.02.00	MEDA PHARMA GMBH & CO. KG	DE
Travex retard 50 mg Hartkapseln, retardiert	UK/H/0225/001	42429.00.00	MEDA PHARMA GMBH & CO. KG	DE
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	2632784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376588	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376687	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376786	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376885	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376984	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	2632883	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377180	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377081	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377388	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377487	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377586	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	2632982	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377685	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377883	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377982	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376083	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 50 mg	UK/H/0225/001	2632685	BGP PRODUCTS	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
Cápsula dura de libertação prolongada			UNIPESSOAL, LDA.	
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376182	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376489	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376380	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TRAVEX® ONE 150 mg Retardtabletten	UK/H/0306/001	46414.00.00	MEDA PHARMA GMBH & CO. KG	DE
TRAVEX® ONE 200 mg Retardtabletten	UK/H/0306/002	46414.01.00	MEDA PHARMA GMBH & CO. KG	DE
TRAVEX® ONE 300 mg Retardtabletten	UK/H/0306/003	46414.02.00	MEDA PHARMA GMBH & CO. KG	DE
TRAVEX® ONE 400 mg Retardtabletten	UK/H/0306/004	46414.03.00	MEDA PHARMA GMBH & CO. KG	DE
Travex® retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	42429.01.00	MEDA PHARMA GMBH & CO. KG	DE
Travex® retard 200 mg Hartkapseln, retardiert	UK/H/0225/004	42429.03.00	MEDA PHARMA GMBH & CO. KG	DE
Tridural 100 mg prolonged-release tablets	FR/H/272/01	5715081	KIRONFARMA, PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Tridural 200 mg prolonged-release tablets	FR/H/272/02	5715180	KIRONFARMA, PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tridural 300 mg prolonged-release tablets	FR/H/272/03	5715289	KIRONFARMA, PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Tropium 100mg/2ml ενέσιμο διάλυμα	not available	23281/17/10-01-2018	MEDOCHEMIE HELLAS SA	GR
VIBRALIS 100mg καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	not available	39659/13/12-12-2014	MEDITRINA PHARMACEUTICAL LIMITED	GR
VIBRALIS 150 MG ΚΑΨΑΚΙΑ ΠΑΡΑΤΕΤΑΜΕΝΗΣ ΑΠΟΔΕΣΜΕΥΣΗΣ, ΣΚΛΗΡΑ	not available	38360/29-5-2012	MEDITRINA PHARMACEUTICAL LIMITED	GR
VIBRALIS 200 mg ΚΑΨΑΚΙΑ ΠΑΡΑΤΕΤΑΜΕΝΗΣ ΑΠΟΔΕΣΜΕΥΣΗΣ, ΣΚΛΗΡΑ	not available	38361/29-5-2012	MEDITRINA PHARMACEUTICAL LIMITED	GR
Xymel 50 mg Capsules	not available	PA0126/099/001	CLONMEL HEALTHCARE LTD.	IE
ZAMADOL 24hr 150mg prolonged release tablets	UK/H/0306/001	PL 16950/0084	NAPP PHARMACEUTICALS LTD	UK
ZAMADOL 24hr 200mg prolonged release tablets.	UK/H/0306/002	PL 16950/0085	NAPP PHARMACEUTICALS LTD	UK
ZAMADOL 24hr 300mg prolonged release tablets.	UK/H/0306/003	PL 16950/0086	NAPP PHARMACEUTICALS LTD	UK
ZAMADOL 24hr 400mg prolonged release tablets.	UK/H/0306/004	PL 16950/0087	NAPP PHARMACEUTICALS LTD	UK
Zamadol Capsules 50 mg	not available	PL 46302/0148	MYLAN PRODUCTS LIMITED	UK
Zamadol Injection	not available	PL 46302/0147	MYLAN PRODUCTS LIMITED	UK
Zamadol Melt 50 mg	not available	PL 46302/0155	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
Tablets				
Zamadol SR 100 mg prolonged-release hard capsules	UK/H/0225/002	PL 46302/0150	MYLAN PRODUCTS LIMITED	UK
Zamadol SR 150 mg prolonged-release hard capsules	UK/H/0225/003	PL 46302/0151	MYLAN PRODUCTS LIMITED	UK
Zamadol SR 200 mg prolonged-release hard capsules	UK/H/0225/004	PL 46302/0152	MYLAN PRODUCTS LIMITED	UK
Zamadol SR 50 mg prolonged-release hard capsules	UK/H/0225/001	PL 46302/0149	MYLAN PRODUCTS LIMITED	UK
ZAMUDOL LP 100 mg, gélule à libération prolongée	UK/H/0225/002	34009 346 544 9 7	MEDA PHARMA SAS	FR
ZAMUDOL LP 100 mg, gélule à libération prolongée	UK/H/0225/002	34009 346 543 2 9	MEDA PHARMA SAS	FR
ZAMUDOL LP 150 mg, gélule à libération prolongée	UK/H/0225/003	34009 346 546 1 9	MEDA PHARMA SAS	FR
ZAMUDOL LP 150 mg, gélule à libération prolongée	UK/H/0225/003	34009 346 545 5 8	MEDA PHARMA SAS	FR
ZAMUDOL LP 200 mg, gélule à libération prolongée	UK/H/0225/004	34009 346 548 4 8	MEDA PHARMA SAS	FR
ZAMUDOL LP 200 mg, gélule à libération prolongée	UK/H/0225/004	34009 346 547 8 7	MEDA PHARMA SAS	FR
ZAMUDOL LP 50 mg, gélule à libération prolongée	UK/H/0225/001	34009 346 542 6 8	MEDA PHARMA SAS	FR

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ZAMUDOL LP 50 mg, gélule à libération prolongée	UK/H/0225/001	34009 346 540 3 9	MEDA PHARMA SAS	FR
Zeridame SR 100mg Prolonged Release Tablets	UK/H/6807/001	PL 20075/1113	ACCORD HEALTHCARE LIMITED	UK
Zeridame SR 150mg Prolonged Release Tablets	UK/H/6807/002	PL 20075/1114	ACCORD HEALTHCARE LIMITED	UK
Zeridame SR 200mg Prolonged Release Tablets	UK/H/6807/003	PL 20075/1115	ACCORD HEALTHCARE LIMITED	UK
ZUMALGIC 100 mg, comprimé effervescent	not available	352 359-5	LABORATOIRE XO	FR
ZUMALGIC 50 mg, comprimé effervescent	not available	34009 561 018 6 3	LABORATOIRE XO	FR
ZYDOL 100 mg / 2 ml Solution for Injection	not available	PA 2242/5/2	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL 50 mg Hard Capsules	not available	PA 2242/5/1	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL 50 mg/ml Solution for Injection	not available	PL 21727/0002	GRÜNENTHAL LTD.	UK
ZYDOL 50mg Capsules	not available	PL 21727/0001	GRÜNENTHAL LTD.	UK
ZYDOL Soluble Tablets	not available	PL 21727/0006	GRÜNENTHAL LTD.	UK
ZYDOL SR 100 mg prolonged-release tablets	not available	PA 2242/5/3	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL SR 100 mg prolonged-release Tablets	not available	PL 21727/0003	GRÜNENTHAL LTD.	UK
ZYDOL SR 150 mg prolonged-release tablets	not available	PA 2242/5/4	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL SR 150 mg prolonged-release tablets	not available	PL 21727/0004	GRÜNENTHAL LTD.	UK
ZYDOL SR 200 mg	not available	PA 2242/5/5	GRÜNENTHAL PHARMA LTD.	IE

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prolonged-release tablets				
ZYDOL SR 200 mg prolonged-release tablets	not available	PL 21727/0005	GRÜNENTHAL LTD.	UK
ZYDOL SR 50 mg prolonged-release tablets	DE/H/0136/004	PA 2242/5/6	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL SR 50 mg prolonged-release tablets	DE/H/0136/004	PA 2242/5/6	GRÜNENTHAL PHARMA LTD.	IE
ZYDOL SR 50 mg prolonged-release tablets	DE/H/0136/004	PL 21727/0024	GRÜNENTHAL LTD.	UK
ZYDOL XL 150 mg prolonged release tablets	not available	PL 16950/0089	NAPP PHARMACEUTICALS LTD	UK
ZYDOL XL 200 mg prolonged release tablets	not available	PL 16950/0090	NAPP PHARMACEUTICALS LTD	UK
ZYDOL XL 300 mg prolonged release tablets	not available	PL 16950/0091	NAPP PHARMACEUTICALS LTD	UK
ZYDOL XL 400 mg prolonged release tablets	not available	PL 16950/0092	NAPP PHARMACEUTICALS LTD	UK
Zytram 150 mg comprimidos de liberación prolongada	UK/H/0306/001	63.130	MUNDIPHARMA PHARMACEUTICALS SL	ES
Zytram 200 mg comprimidos de liberación prolongada	UK/H/0306/002	63.131	MUNDIPHARMA PHARMACEUTICALS SL	ES
Zytram 400 mg comprimidos de liberación prolongada	UK/H/0306/004	63.133	MUNDIPHARMA PHARMACEUTICALS SL	ES
Zytram BID 75 mg comprimidos de liberación prolongada	UK/H/0330/001	63.134	MUNDIPHARMA PHARMACEUTICALS SL	ES
TRAMAL® SR 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	22043/15/06-03-2017	VIANEX S.A.	GR
TRAMAL® SR 150 mg δισκία παρατεταμένης	not available	22044/15/06-03-2017	VIANEX S.A.	GR

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αποδέσμευσης				
TRAMAL® SR 200 mg δισκία παρατεταμένης αποδέσμευσης	not available	22045/15/06-03-2017	VIANEX S.A.	GR
TRAMAL® SR 50 mg δισκία παρατεταμένης αποδέσμευσης	not available	22042/15/06-03-2017	VIANEX S.A.	GR
TRAMAL® Καψάκιο, σκληρό 50 mg	not available	97924/10-12-2014	VIANEX S.A.	GR
Маброн 50 mg капсули	not available	20030082	MEDOCHEMIE LTD.	BG
Маброн 50 mg капсули	not available	20030082	MEDOCHEMIE LTD.	BG
Маброн 50 mg/ml инжекционен разтвор	not available	20030083	MEDOCHEMIE LTD.	BG
Маброн 50 mg/ml инжекционен разтвор	not available	20030083	MEDOCHEMIE LTD.	BG
Маброн MR 100 mg таблетки с удължено освобождаване	not available	20100447	MEDOCHEMIE LTD.	BG
Маброн MR 150 mg таблетки с удължено освобождаване	not available	20100448	MEDOCHEMIE LTD.	BG
Маброн MR 200 mg таблетки с удължено освобождаване	not available	20100449	MEDOCHEMIE LTD.	BG
Трамадол Крка 100 mg таблетки с удължено освобождаване	EE/H/0262/001	20180342	KRKA, D.D., NOVO MESTO	BG
Трамадол Крка 100 mg/2 ml инжекционен/инфузионен разтвор	HR/H/0104/002	20180022	KRKA, D.D., NOVO MESTO	BG
Трамадол Крка 150 mg таблетки с удължено освобождаване	EE/H/0262/002	20180341	KRKA, D.D., NOVO MESTO	BG

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Трамадол Крка 200 mg таблетки с удължено освобождаване	EE/H/0262/003	20180343	KRKA, D.D., NOVO MESTO	BG
Трамадол Крка 50 mg капсули, твърди	HR/H/0104/003	20180037	KRKA, D.D., NOVO MESTO	BG
Трамадол Крка 50 mg/ml инжекционен/инфузионен разтвор	HR/H/0104/002	20180021	KRKA, D.D., NOVO MESTO	BG
Трамадол СТАДА 50 mg твърди капсули	not available	9600290	STADA ARZNEIMITTEL AG	BG
Трамадол СТАДА 50 mg/ml инжекционен разтвор	not available	9600289	STADA ARZNEIMITTEL AG	BG
Трамалгин 50 mg твърди капсули	not available	20030660	SOPHARMA AD	BG
Трамалгин 50 mg/ml инжекционен разтвор	not available	20050203	SOPHARMA AD	BG