26 January 2018
EMA/270645/2015
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: tramadol

Procedure No.: PSUSA/00003002/201705

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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	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 100 mg retard kemény kapszula	not available	OGYI-T-6970/05	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 100 mg retard kemény kapszula	not available	OGYI-T-6970/06	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561338	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561340	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561011	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561023	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561353	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561035	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561365	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561050	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561391	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561377	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561047	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561389	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561403	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561062	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561415	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561074	MEDA PHARMA S.P.A.	IT

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ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561427	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561441	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561439	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561086	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561466	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg compresse a rilascio prolungato	UK/H/0306/001	034561454	MEDA PHARMA S.P.A.	IT
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/08	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/07	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 150 mg retard kemény kapszula	not available	OGYI-T-6970/09	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561478	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561480	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561098	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561100	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561112	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561492	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561504	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561528	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561124	MEDA PHARMA S.P.A.	IT

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ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561516	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561136	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561530	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561542	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561567	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561555	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561148	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561151	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561579	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561581	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561593	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561163	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg compresse a rilascio prolungato	UK/H/0306/002	034561605	MEDA PHARMA S.P.A.	IT
ADAMON 200 mg retard kemény kapszula	not available	OGYI-T-6970/10	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 200 mg retard kemény kapszula	not available	OGYI-T-6970/11	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561617	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561175	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561629	MEDA PHARMA S.P.A.	IT

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ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561187	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561631	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561199	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561643	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561656	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561201	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561670	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561668	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561682	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561213	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561225	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561706	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561694	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561237	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561718	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561720	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561732	MEDA PHARMA S.P.A.	IT
ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561249	MEDA PHARMA S.P.A.	IT

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ADAMON 300 mg compresse a rilascio prolungato	UK/H/0306/003	034561744	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561757	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561252	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561769	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561264	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561771	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561276	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561783	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561795	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561288	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561807	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561821	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561819	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561833	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561290	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561302	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561845	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561314	MEDA PHARMA S.P.A.	IT

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ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561858	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561326	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561860	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561872	MEDA PHARMA S.P.A.	IT
ADAMON 400 mg compresse a rilascio prolungato	UK/H/0306/004	034561884	MEDA PHARMA S.P.A.	IT
Adamon 50 mg - Schmelztabletten	UK/H/0640/001	1-25733	MEDA PHARMA GMBH	AT
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/01	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/02	MEDA PHARMA HUNGARY KFT.	HU
ADAMON 50 mg retard kemény kapszula	not available	OGYI-T-6970/03	MEDA PHARMA HUNGARY 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 200 mg-Filmtabletten	UK/H/0306/002	1-23431	MEDA PHARMA GMBH	AT
Adamon long retard 300 mg-Filmtabletten	UK/H/0306/003	1-23432	MEDA PHARMA GMBH	AT
Adamon long retard 400 mg-Filmtabletten	UK/H/0306/004	1-23433	MEDA PHARMA GMBH	AT
Adamon SR 100 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/002	65/0182/06-S	MEDA PHARMA GMBH & CO. KG	SK
Adamon SR 100, 100 mg, kapsułki o przedłużonym uwalnianiu	not available	9361	MEDA PHARMA GMBH & CO. KG	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
Adamon SR 150 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/003	65/0183/06-S	MEDA PHARMA GMBH & CO. KG	SK
Adamon SR 150, 150 mg, kapsułki o przedłużonym uwalnianiu	not available	9362	MEDA PHARMA GMBH & CO. KG	PL
Adamon SR 200 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/004	65/0185/06-S	MEDA PHARMA GMBH & CO. KG	SK
Adamon SR 200, 200 mg, kapsułki o przedłużonym uwalnianiu	not available	9363	MEDA PHARMA GMBH & CO. KG	PL
Adamon SR 50 mg, tvrdé kapsuly s predĺženým uvoľňovaním	UK/H/0301/001	65/0184/06-S	MEDA PHARMA GMBH & CO. KG	SK
Adamon SR 50, 50 mg, kapsułki o przedłużonym uwalnianiu	not available	9360	MEDA PHARMA GMBH & CO. KG	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/ 2 ml solución inyectable	not available	59.086	GRÜNENTHAL PHARMA S.A.	ES
Adolonta 100 mg/ml gotas orales en solución	not available	61.617	GRÜNENTHAL PHARMA S.A.	ES
Adolonta 50 mg cápsulas duras	not available	59.088	GRÜNENTHAL PHARMA S.A.	ES
Adolonta 50 mg cápsulas duras	not available	59.088	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

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Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 100 mg comprimidos de liberación prolongada	DE/H/0108/001	61784	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 150 mg comprimidos de liberación prolongada	DE/H/0108/002	61785	GRÜNENTHAL PHARMA S.A.	ES
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

Product 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 200 mg comprimidos de liberación prolongada	DE/H/0108/003	61786	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 200 mg comprimidos de liberación prolongada	DE/H/0108/003	61786	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 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
Adolonta retard 50 mg comprimidos de liberación prolongada	DE/H/0108/004	68570	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 50 mg comprimidos de liberación prolongada	DE/H/0108/004	68570	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 50 mg comprimidos de liberación prolongada	DE/H/0108/004	68570	GRÜNENTHAL PHARMA S.A.	ES
Adolonta retard 50 mg comprimidos de liberación prolongada	DE/H/0108/004	68570	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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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® 100 mg Retardtabletten	not available	42878.00.01	TAD PHARMA GMBH	DE
Amadol® 150 mg Retardtabletten	not available	42878.01.01	TAD PHARMA GMBH	DE
Amadol® 200 mg Retardtabletten	not available	42878.02.01	TAD PHARMA GMBH	DE
Amadol® Kapseln	not available	32283.00.00	TAD PHARMA GMBH	DE
BIODALGIC 50 mg, comprimé effervescent	not available	561 772-2	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	350 673-4	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	561 772-2	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	350 673-4	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	561 772-2	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	350 673-4	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	561 772-2	BIOCODEX	FR
BIODALGIC 50 mg, comprimé effervescent	not available	350 673-4	BIOCODEX	FR
Brimisol PR 100 mg Prolonged-Release Tablet	not available	17907/ 0134	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Brimisol PR 200 mg Prolonged-Release Tablet	not available	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

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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	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
Ceparidin 50 mg capsulas EFG	not available	64.448	ARAFARMA GROUP, S.A	ES
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-4975/08	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg retard filmtabletta	not available	OGYI-T-4975/07	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg retard filmtabletta	not available	OGYI-T-4975/08	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg retard filmtabletta	not available	OGYI-T-4975/07	TEVA GYÓGYSZERGYÁR ZRT	HU
CONTRAMAL 100 mg suppositoires	not available	BE 163064	SA GRÜNENTHAL N.V.	BE
Contramal 100 mg végbélkúp	not available	OGYI-T-4975/06	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg végbélkúp	not available	OGYI-T-4975/06	TEVA GYÓGYSZERGYÁR ZRT	HU
CONTRAMAL 100 mg zetpillen	not available	BE 163064	SA GRÜNENTHAL N.V.	BE
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

Product 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 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	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg/ml belsőleges oldatos cseppek	not available	OGYI-T-4975/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg/ml belsőleges oldatos cseppek adagolópumpával	not available	OGYI-T-4975/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 100 mg/ml belsőleges oldatos cseppek adagolópumpával	not available	OGYI-T-4975/05	TEVA GYÓGYSZERGYÁR ZRT	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 soluzione orale con erogatore	not available	028853101	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL 100 mg/ml, solution buvable	not available	34009 362 268 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

Product 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 150 mg compresse a rilascio prolungato	DE/H/0108/002	028853 075	GRÜNENTHAL ITALIA S.R.L.	IT
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	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 150 mg retard filmtabletta	not available	OGYI-T-4975/09	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 150 mg retard filmtabletta	not available	OGYI-T-4975/10	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 150 mg retard filmtabletta	not available	OGYI-T-4975/09	TEVA GYÓGYSZERGYÁR ZRT	HU
CONTRAMAL 200 mg compresse a rilascio prolungato	DE/H/0108/003	028853 087	GRÜNENTHAL ITALIA S.R.L.	IT
CONTRAMAL 200 mg compresse a rilascio prolungato	DE/H/0108/003	028853 087	GRÜNENTHAL ITALIA S.R.L.	IT
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-4975/11	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 200 mg retard filmtabletta	not available	OGYI-T-4975/12	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 200 mg retard filmtabletta	not available	OGYI-T-4975/11	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 200 mg retard filmtabletta	not available	OGYI-T-4975/12	TEVA GYÓGYSZERGYÁR ZRT	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

Product 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 kemény kapszula	not available	OGYI-T-4975/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 50 mg kemény kapszula	not available	OGYI-T-4975/02	TEVA GYÓGYSZERGYÁR ZRT	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
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/13	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/13	TEVA GYÓGYSZERGYÁR ZRT	HU
Contramal 50 mg/ml oldatos injekció	not available	OGYI-T-4975/03	TEVA GYÓGYSZERGYÁR ZRT	HU
CONTRAMAL 50 mg/ml soluzione iniettabile	not available	028853051	GRÜNENTHAL ITALIA S.R.L.	IT
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	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	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
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

Product 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	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	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	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	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

Product 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	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	2003070150	SA GRÜNENTHAL N.V.	LU
Contramal Retard 100 mg comprimés à libération prolongée	DE/H/0108/001	2003070150	SA GRÜNENTHAL N.V.	LU
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

Product 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 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	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	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

Product 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 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 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	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	BE188483	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 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 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	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

Product 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 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 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	BE364752	SA GRÜNENTHAL N.V.	BE
Contramal Retard 150 mg comprimés à libération prolongée	DE/H/0108/002	BE364752	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 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	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 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

Product 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 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 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 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

Product 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 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	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 comprimés à libération prolongée	DE/H/0108/002	2003070152	SA GRÜNENTHAL N.V.	LU
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

Product 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 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 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

Product 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 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 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

Product 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 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
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

Product 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 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 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	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

Product 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 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	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	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	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

Product 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 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	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	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	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

Product 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 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	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	2003070151	SA GRÜNENTHAL N.V.	LU
Contramal Retard 200 mg comprimés à libération prolongée	DE/H/0108/003	2003070151	SA GRÜNENTHAL N.V.	LU
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

Product 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	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
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

Product 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	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
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	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
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 comprimés à libération prolongée	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
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	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

Product 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 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	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	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

Product 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 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	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

Product 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 50 mg comprimés à libération prolongée	DE/H/0108/004	2007040035	SA GRÜNENTHAL N.V.	LU
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 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	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

Product 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 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
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	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

Product 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 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	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	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	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

Product 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 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	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 Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281653	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 Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg comprimés à libération prolongée	FR/H/0272/001	0477/08040052	SA GRÜNENTHAL N.V.	LU
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	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 Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281653	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281671	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	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 Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 100 mg tabletten met verlengde afgifte	FR/H/0272/001	BE281662	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	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 Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	BE281696	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg comprimés à libération prolongée	FR/H/0272/002	0477/08040053	SA GRÜNENTHAL N.V.	LU
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	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 Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281687	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281696	SA GRÜNENTHAL N.V.	BE
Contramal Uno 200 mg tabletten met verlengde afgifte	FR/H/0272/002	BE281705	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281714	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	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 Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg comprimés à libération prolongée	FR/H/0272/003	0477/08040054	SA GRÜNENTHAL N.V.	LU
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281714	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	SA GRÜNENTHAL N.V.	BE
Contramal Uno 300 mg tabletten met verlengde afgifte	FR/H/0272/003	BE281723	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 100 mg Depottabletter	UK/H/4160/001	48179	MYLAN AB	DK
Dolatramyl 100 mg depottabletter	UK/H/4160/001	10-8115	MYLAN AB	NO
Dolatramyl 100 mg depottabletter	UK/H/4160/001	45573	MYLAN AB	SE
Dolatramyl 100 mg depottabletti	UK/H/4160/001	29450	MYLAN AB	FI
Dolatramyl 150 mg Depottabletter	UK/H/4160/002	48180	MYLAN AB	DK
Dolatramyl 150 mg depottabletter	UK/H/4160/002	10-8116	MYLAN AB	NO
Dolatramyl 150 mg depottabletter	UK/H/4160/002	45574	MYLAN AB	SE
Dolatramyl 150 mg depottabletti	UK/H/4160/002	29451	MYLAN AB	FI
Dolatramyl 200 mg Depottabletter	UK/H/4160/002	48181	MYLAN AB	DK
Dolatramyl 200 mg depottabletter	UK/H/4160/003	10-8117	MYLAN AB	NO
Dolatramyl 200 mg depottabletter	UK/H/4160/003	45575	MYLAN AB	SE
Dolatramyl 200 mg depottabletti	UK/H/4160/003	29452	MYLAN AB	FI
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
DOLPAR 100 mg comprimidos de liberación prolongada	FR/H/0272/001	67.585	LABORATORIOS DR. ESTEVE 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
DOLPAR 200 mg comprimidos de liberación prolongada	FR/H/0272/002	67.586	LABORATORIOS DR. ESTEVE S.A.	ES
DOLPAR 300 mg comprimidos de liberación prolongada	FR/H/0272/003	67.587	LABORATORIOS DR. ESTEVE S.A.	ES
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 ampoules, solution injectable	not available	BE168603	ZAMBON NV	BE
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	0411/11071217	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

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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	0411/10120898	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 afgifte	UK/H/0330/003	BE370203	ZAMBON NV	BE
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	0411/10120899	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	0411/10120896	ZAMBON NV	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
DOLZAM UNO 150 mg comprimés à libération prolongée	UK/H/0306/001	BE207916	ZAMBON NV	BE
DOLZAM UNO 150 mg comprimés à libération prolongée	UK/H/0306/001	BE370221	ZAMBON NV	BE
DOLZAM UNO 150 mg comprimés à libération prolongée	UK/H/0306/001	0411/10120894	ZAMBON NV	LU
DOLZAM UNO 200 mg comprimés à libération prolongée	UK/H/0306/002	BE207925	ZAMBON NV	BE
DOLZAM UNO 200 mg comprimés à libération prolongée	UK/H/0306/002	BE370237	ZAMBON NV	BE
Dolzam Uno 200 mg comprimés à libération prolongée	UK/H/0306/002	0411/10120893	ZAMBON NV	LU
Dolzam Uno 200 mg tabletten met verlengde afgifte	UK/H/0306/002	BE207925	ZAMBON NV	BE
DOLZAM UNO 200 mg tabletten met verlengde afgifte	UK/H/0306/002	BE370237	ZAMBON NV	BE
DOLZAM UNO 300 mg comprimés à libération prolongée	UK/H/0306/003	BE207934	ZAMBON NV	BE
DOLZAM UNO 300 mg comprimés à libération prolongée	UK/H/0306/003	BE370246	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
Dolzam Uno 300 mg comprimés à libération prolongée	UK/H/0306/003	0411/10120892	ZAMBON NV	LU
Dolzam Uno 300 mg tabletten met verlengde afgifte	UK/H/0306/003	BE207934	ZAMBON NV	BE
DOLZAM UNO 300 mg tabletten met verlengde afgifte	UK/H/0306/003	BE370246	ZAMBON NV	BE
DOLZAM UNO 400 mg comprimés à libération prolongée	UK/H/0306/004	BE370255	ZAMBON NV	BE
Dolzam Uno 400 mg comprimés à libération prolongée	UK/H/0306/004	0411/10120895	ZAMBON NV	LU
Dolzam Uno 400 mg tabletten met verlengde afgifte	UK/H/0306/004	BE370255	ZAMBON NV	BE
FORTRADOL 100 mg compresse a rilascio prolungato	not available	028878092	ALFA WASSERMANN SPA	IT
FORTRADOL 100 mg/2 ml soluzione iniettabile	not available	028878128	ALFA WASSERMANN SPA	IT
FORTRADOL 100 mg/ml gocce orali soluzione con contagocce	not available	028878080	ALFA WASSERMANN SPA	IT
FORTRADOL 150 mg compresse a rilascio prolungato	not available	028878142	ALFA WASSERMANN SPA	IT
FORTRADOL 200 mg compresse a rilascio prolungato	not available	028878155	ALFA WASSERMANN SPA	IT
FORTRADOL 50 mg capsule rigide	not available	028878078	ALFA WASSERMANN SPA	IT
FORTRADOL 50 mg/ml soluzione iniettabile	not available	028878116	ALFA WASSERMANN 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
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 cápsulas de libertação prolongada	DE/H/0639/002	5010756	FERRER INTERNACIONAL, S.A.	PT
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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 depotkapsel, hård	UK/H/0225/004	14066	MEDA AB	SE
Gemadol 50 mg depotkapsel, hård	UK/H/0225/001	14063	MEDA AB	SE
Gemadol Retard	UK/H/0225/002	19326	MEDA AS	DK
Gemadol Retard	UK/H/0225/003	19327	MEDA AS	DK
Gemadol Retard	UK/H/0225/001	19325	MEDA AS	DK
Gemadol Retard	UK/H/0225/004	19328	MEDA AS	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
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/2 ml otopina za injekcije	not available	UP/I-530-09/12-02/89	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 100 mg/ml oralne kapi	not available	UP/I-530-09/12-02/92	BELUPO D.D.	HR
LUMIDOL 50 mg tvrde kapsule	not available	UP/I-530-09/12-02/91	BELUPO D.D.	HR
LUMIDOL 50 mg/ml otopina za injekcije	not available	UP/I-530-09/12-02/88	BELUPO D.D.	HR
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	not available	65/0459/94-S	MEDOCHEMIE LTD.	SK
MABRON 100 mg/2 ml, injekcní roztok	not available	65/788/94-C	MEDOCHEMIE LTD.	CZ
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	not available	65/0076/93-S	MEDOCHEMIE LTD.	SK
MABRON 50 mg capsule	not available	8728/2016/01	MEDOCHEMIE LTD.	RO
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 tobolky	not available	65/1000/93-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
MABRON 50 mg/ml injekcinis 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 prelungita	not available	7244/2014/01	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/02	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/03	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/04	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/05	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/06	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/07	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/08	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/09	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/10	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/11	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 prelungita	not available	7244/2014/12	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/13	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/14	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/15	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/16	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/17	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/18	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/19	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/20	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/21	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/22	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/23	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 prelungita	not available	7244/2014/24	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/25	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/26	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/27	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/28	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/29	MEDOCHEMIE LTD.	RO
Mabron retard 100 mg comprimate cu eliberare prelungita	not available	7244/2014/30	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

Product 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/1/07/0754/006	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 atpalaidavimo tabletes	NL/H/0889/001	LT/1/07/0754/010	MEDOCHEMIE LTD.	LT
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

Product 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/1/07/0754/018	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
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

Product 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/1/07/0754/030	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 predlženým uvolnovaním	NL/H/0538/001	65/0275/05-S	MEDOCHEMIE LTD.	SK
Mabron retard 100 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/001	20240	MEDOCHEMIE LTD.	CY
MABRON RETARD 100 tablety s prodlouženým uvolňováním	NL/H/0538/001	65/369/05-C	MEDOCHEMIE LTD.	CZ
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/01	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/02	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/03	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/04	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/05	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/06	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/07	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 prelungita	not available	7245/2014/08	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/09	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/10	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/11	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/12	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/13	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/14	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/15	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/16	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/17	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/18	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/19	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 prelungita	not available	7245/2014/20	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/21	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/22	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/23	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/24	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/25	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/26	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/27	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/28	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/29	MEDOCHEMIE LTD.	RO
Mabron retard 150 mg comprimate cu eliberare prelungita	not available	7245/2014/30	MEDOCHEMIE LTD.	RO
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/031	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/1/07/0754/032	MEDOCHEMIE LTD.	LT
MABRON RETARD 150 mg pailginto atpalaidavimo tabletes	NL/H/0889/002	LT/1/07/0754/033	MEDOCHEMIE LTD.	LT
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

Product 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/1/07/0754/044	MEDOCHEMIE LTD.	LT
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 predlženým uvolnovaním	NL/H/0538/002	65/0285/05-S	MEDOCHEMIE LTD.	SK
Mabron retard 150 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/002	20241	MEDOCHEMIE LTD.	CY
MABRON RETARD 150 tablety s prodlouženým uvolňováním	NL/H/0538/002	65/370/05-C	MEDOCHEMIE LTD.	CZ
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/01	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/02	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/03	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
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/04	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/05	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/06	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/07	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/08	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/09	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/10	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/11	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/12	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/13	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/14	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/15	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
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/16	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/17	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/18	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/19	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/20	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/21	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/22	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/23	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/24	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/25	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/26	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/27	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 prelungita	not available	7246/2014/28	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/29	MEDOCHEMIE LTD.	RO
Mabron retard 200 mg comprimate cu eliberare prelungita	not available	7246/2014/30	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 mg pailginto atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/069	MEDOCHEMIE LTD.	LT
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

Product 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 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 atpalaidavimo tabletes	NL/H/0889/003	LT/1/07/0754/080	MEDOCHEMIE LTD.	LT
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

Product 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 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
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 predlženým uvolnovaním	NL/H/0538/003	65/0286/05-S	MEDOCHEMIE LTD.	SK
Mabron retard 200 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0889/003	20239	MEDOCHEMIE LTD.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MABRON RETARD 200 tablety s prodlouženým uvolnováním	NL/H/0538/003	65/371/05-C	MEDOCHEMIE LTD.	CZ
Mabron retard, 100 mg toimeainet modifitseeritult vabastavad tabletid	NL/H/0538/001	493405	MEDOCHEMIE LTD.	EE
Mabron retard, 150 mg toimeainet modifitseeritult vabastavad tabletid	NL/H/0538/002	493605	MEDOCHEMIE LTD.	EE
Mabron retard, 200 mg toimeainet modifitseeritult vabastavad tabletid	NL/H/0538/003	493505	MEDOCHEMIE LTD.	EE
MABRON soluţie injectabilă	not available	7314/2006/01	MEDOCHEMIE LTD.	RO
MABRON soluţie injectabilă	not available	7314/2006/02	MEDOCHEMIE LTD.	RO
MABRON soluţie injectabilă	not available	7314/2006/03	MEDOCHEMIE LTD.	RO
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 ilgstošās darbības tabletes	NL/H/0538/003	05-0502	MEDOCHEMIE LTD.	LV
Mabron, 50 mg/ml süste- või infusioonilahus	not available	340701	MEDOCHEMIE LTD.	EE
Mandolgin	not available	18666	SANDOZ A/S	DK
Mandolgin	not available	18665	SANDOZ A/S	DK
Mandolgin	not available	19352	SANDOZ A/S	DK
Mandolgin Retard	not available	19423	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
Mandolgin Uno	not available	19085	SANDOZ A/S	DK
Mandolgin Uno	not available	19086	SANDOZ A/S	DK
Mandolgin Uno	not available	19087	SANDOZ A/S	DK
Mandolgin Uno	not available	19088	SANDOZ A/S	DK
Maneo 100 mg Prolonged-release tablets	UK/H/4160/001	PL 04569/1406	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 150 mg Prolonged-release tablets	UK/H/4160/002	PL 04569/1407	GENERICS [UK] LIMITED	UK
Maneo 200 mg Prolonged-release tablets	UK/H/4160/003	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 08829/0163	CHIESI LIMITED	UK
MAXITRAM SR 150 mg prolonged-release capsule, hard	DE/H/0639/003	PL 08829/0164	CHIESI LIMITED	UK
MAXITRAM SR 200 mg prolonged-release capsule, hard	DE/H/0639/004	PL 08829/0165	CHIESI LIMITED	UK
MAXITRAM SR 50 mg prolonged-release capsule, hard	DE/H/0639/001	PL 08829/0162	CHIESI LIMITED	UK
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 870-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 872-5	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 873-1	SANOFI-AVENTIS 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
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 871-9	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 383-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 869-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 385-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 877-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 384-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 875-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 878-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 876-0	SANOFI-AVENTIS 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
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 874-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 870-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 872-5	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 873-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 871-9	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 383-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 869-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 385-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 877-7	SANOFI-AVENTIS 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
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 384-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 875-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 878-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 876-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 100 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 874-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 945-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 520-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 944-6	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 942-3	SANOFI-AVENTIS 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
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 941-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 940-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 947-5	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 948-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 521-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 522-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 950-6	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 949-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 946-9	SANOFI-AVENTIS 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
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 945-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 520-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 944-6	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 942-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 941-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 940-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 947-5	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 948-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 521-8	SANOFI-AVENTIS 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
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 522-4	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 950-6	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 949-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 200 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 946-9	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 953-5	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 952-9	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 951-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 954-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 955-8	SANOFI-AVENTIS 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
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 523-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 089-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 092-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 525-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 524-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 091-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 090-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 094-6	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 953-5	SANOFI-AVENTIS 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
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 952-9	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 951-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 954-1	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	367 955-8	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 523-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 089-2	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 092-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 525-3	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	566 524-7	SANOFI-AVENTIS 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
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 091-7	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 090-0	SANOFI-AVENTIS FRANCE	FR
MONOALGIC LP 300 MG, COMPRIME A LIBERATION PROLONGÉE (UNE PRISE QUOTIDIENNE)	not available	368 094-6	SANOFI-AVENTIS FRANCE	FR
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 470 6 2	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 100 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/001	34009 362 471 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

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 478 7 1	THERABEL LUCIEN PHARMA S.A.	FR
MONOCRIXO L.P. 150 mg, gélule à libération prolongée (UNE PRISE QUOTIDIENNE)	FI/H/0164/002	34009 362 479 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
MONOTRAMAL L.P. 100 mg, comprimé à libération prolongée (UNE PRISE QUOTIDIENNE)	FR/H/0272/001	NL28699	ENDO VENTURES LIMITED	FR
MONOTRAMAL L.P. 200 mg, comprimé à libération prolongée (UNE PRISE QUOTIDIENNE)	FR/H/0272/002	NL28700	ENDO VENTURES LIMITED	FR
MONOTRAMAL L.P. 300 mg, comprimé à libération prolongée (UNE PRISE QUOTIDIENNE)	FR/H/0272/003	NL28701	ENDO VENTURES LIMITED	FR
Noax uno 100 mg Retardtabletten	FR/H/0272/001	1-26327	ANGELINI PHARMA ÖSTERREICH GMBH	AT
Noax Uno 100 mg, tabletki o przedluzonym uwalnianiu	FR/H/0272/001	12067	ANGELINI PHARMA ÖSTERREICH GMBH	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
Noax uno 200 mg Retardtabletten	FR/H/0272/002	1-26329	ANGELINI PHARMA ÖSTERREICH GMBH	AT
Noax Uno 200 mg, tabletki o przedłużonym uwalnianiu	FR/H/0272/002	12064	ANGELINI PHARMA ÖSTERREICH GMBH	PL
Noax uno 300 mg Retardtabletten	FR/H/0272/003	1-26331	ANGELINI PHARMA ÖSTERREICH GMBH	AT
Noax Uno 300 mg, tabletki o przedluzonym uwalnianiu	FR/H/0272/003	12063	ANGELINI PHARMA ÖSTERREICH GMBH	PL
Nobligan 100 mg/ml orala droppar, lösning	not available	14305	GRÜNENTHAL GMBH	SE
Nobligan 50 mg hörð hylki	not available	910181	GRÜNENTHAL GMBH	IS
Nobligan 50 mg kapslar, hårda	not available	12845	GRÜNENTHAL GMBH	SE
Nobligan 50 mg kapsler, harde	not available	8268	GRÜNENTHAL GMBH	NO
Nobligan Retard 100 mg depottabletter	not available	98-2970	GRÜNENTHAL GMBH	NO
Nobligan retard 100 mg depottabletter	DE/H/0108/001	13287	GRÜNENTHAL GMBH	SE
Nobligan retard 100 mg depottabletter	DE/H/0108/001	13287	GRÜNENTHAL GMBH	SE
Nobligan retard 100 mg depottabletter	DE/H/0108/001	13287	GRÜNENTHAL GMBH	SE
Nobligan Retard 100 mg forðatöflur	not available	980423	GRÜNENTHAL GMBH	IS
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 depottabletter	DE/H/0108/002	13288	GRÜNENTHAL GMBH	SE
Nobligan retard 150 mg depottabletter	DE/H/0108/002	13288	GRÜNENTHAL GMBH	SE
Nobligan Retard 150 mg forðatöflur	not available	980424	GRÜNENTHAL 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
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 depottabletter	DE/H/0108/003	13289	GRÜNENTHAL GMBH	SE
Nobligan retard 200 mg depottabletter	DE/H/0108/003	13289	GRÜNENTHAL GMBH	SE
Nobligan Retard 200 mg forðatöflur	not available	980425	GRÜNENTHAL GMBH	IS
Nobligan Retard, depottabletter 100 mg	DE/H/0108/001	18540	GRÜNENTHAL GMBH	DK
Nobligan Retard, depottabletter 100 mg	DE/H/0108/001	18540	GRÜNENTHAL GMBH	DK
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 150 mg	DE/H/0108/002	18541	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 Retard, depottabletter 200 mg	DE/H/0108/003	18542	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
ORATRAM 100 100 mg, tabletki o przedluzonym uwalnianiu	NL/H/0888/001	14313	MOLTENI FARMACEUTICI POLSKA SP. Z O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ORATRAM 150 150 mg, tabletki o przedluzonym uwalnianiu	NL/H/0888/002	14315	MOLTENI FARMACEUTICI POLSKA SP. Z O.O.	PL
ORATRAM 200 200 mg, tabletki o przedłużonym uwalnianiu	NL/H/0888/003	14314	MOLTENI FARMACEUTICI POLSKA SP. Z O.O.	PL
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 004.3	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 003.7	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 002.0	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 001.4	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 000.8	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 515.4	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 514.8	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 513.1	MEDA PHARMA SAS	FR
OROZAMUDOL 50 mg, comprimé orodispersible	UK/H/0640/001	366 512.5	MEDA PHARMA SAS	FR
Paxilfar 100 mg/ 2 ml solution for injection	not available	9515411	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
Paxilfar 100mg tablets	not available	6/13/84	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
Poltram, 100 mg/ml, krople doustne, roztwór	not available	9690	MEDANA PHARMA SPOLKA AKCYJNA	PL
Poltram, 100 mg/ml, krople doustne, roztwór	not available	9690	MEDANA PHARMA SPOLKA AKCYJNA	PL
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074042	THERABEL GIENNE 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
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074055	THERABEL GIENNE PHARMA S.P.A.	IT
PRONTALGIN 50 mg Compresse effervescenti	NL/H/0113/005	033074067	THERABEL GIENNE PHARMA S.P.A.	IT
PRONTALGIN 50 mg/ml soluzione iniettabile	NL/H/0113/001	033074028	THERABEL GIENNE PHARMA S.P.A.	IT
PRONTALGIN Capsule, 50 mg capsule rigide	NL/H/0113/002	033074030	THERABEL GIENNE PHARMA S.P.A.	IT
PRONTALGIN Gocce, 100 mg/ml gocce orali, soluzione	NL/H/0113/003	033074016	THERABEL GIENNE PHARMA S.P.A.	IT
Ralgen 50 mg kemény kapszula	not available	OGYI-T-20310/15	ZENTIVA, A.S.	HU
Ralgen 50 mg kemény kapszula	not available	OGYI-T-20310/14	ZENTIVA, A.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/04	ZENTIVA, A.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/05	ZENTIVA, A.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/03	ZENTIVA, A.S.	HU
Ralgen SR 100 mg retard tabletta	not available	OGYI-T-20310/02	ZENTIVA, A.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/07	ZENTIVA, A.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/09	ZENTIVA, A.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/08	ZENTIVA, A.S.	HU
Ralgen SR 150 mg retard tabletta	not available	OGYI-T-20310/06	ZENTIVA, A.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/12	ZENTIVA, A.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/11	ZENTIVA, A.S.	HU
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/13	ZENTIVA, A.S.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ralgen SR 200 mg retard tabletta	not available	OGYI-T-20310/10	ZENTIVA, A.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	NP/H/0828/006	H/93/01481/005	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg svečke	NP/H/0828/006	H/93/01481/005	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg tablete s podaljšanim sproščanjem	NP/H/0828/002	H/93/01481/001	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg/ml peroralne kapljice, raztopina	NP/H/0828/005	H/93/01481/006	KRKA, D.D., NOVO MESTO	SI
Tadol 100 mg/ml peroralne kapljice, raztopina	NP/H/0828/005	H/93/01481/007	KRKA, D.D., NOVO MESTO	SI
Tadol 150 mg tablete s podaljšanim sproščanjem	NP/H/0828/003	H/93/01481/002	KRKA, D.D., NOVO MESTO	SI
Tadol 200 mg tablete s podaljšanim sproščanjem	NP/H/0828/004	H/93/01481/003	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg trde kapsule	NP/H/0828/001	H/93/01481/004	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	NP/H/0829/001	H/93/01481/010	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	NP/H/0829/001	H/93/01481/011	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	NP/H/0829/001	H/93/01481/008	KRKA, D.D., NOVO MESTO	SI
Tadol 50 mg/ml raztopina za injiciranje/infundiranje	NP/H/0829/001	H/93/01481/009	KRKA, D.D., NOVO MESTO	SI
Tadol, kapsler, hårde	DE/H/0282/001	32972	PHARMACODANE APS	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
Tadol, suppositorier	DE/H/0282/003	32974	PHARMACODANE APS	DK
TAKADOL® 100 mg, comprimé effervescent sécable	not available	355 314-2	LABORATOIRES EXPANSCIENCE	FR
TAKADOL® 100 mg, comprimé effervescent sécable	not available	355 315-9	LABORATOIRES EXPANSCIENCE	FR
TAKADOL® 100 mg, comprimé effervescent sécable	not available	355 316-5	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, druppels 100 mg/ml	NL/H/0113/003	RVG 17780	THERABEL PHARMA N.V.	NL
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 gotas orales en solución	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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Tiparol 50 mg hård kapsel	not available	13347	BLUEFISH PHARMACEUTICALS AB	SE
Tiparol 50 mg hård kapsel	not available	13347	BLUEFISH PHARMACEUTICALS AB	SE
T-long ® 100 mg Retardkapseln Hartkapsel, retardiert Wirkstoff: Tramadolhydrochlorid	FI/H/0164/001	54607.00.00	LABORATOIRES SMB S.A.	DE
T-long ® 150 mg Retardkapseln Hartkapsel, retardiert Wirkstoff: Tramadolhydrochlorid	FI/H/0164/002	54607.01.00	LABORATOIRES SMB S.A.	DE
T-long ® 200 mg Retardkapseln Hartkapsel, retardiert Wirkstoff: Tramadolhydrochlorid	FI/H/0164/003	54607.02.00	LABORATOIRES SMB S.A.	DE
TOPALGIC 100 mg/2 ml, solution injectable en ampoule	not available	558 881-9	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 100 mg/2 ml, solution injectable en ampoule	not available	558 881-9	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 100 MG/ML SOLUTION BUVABLE	not available	362 067-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 100 MG/ML SOLUTION BUVABLE	not available	362 067-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 50 MG, GÉLULE	not available	558 878-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 50 MG, GÉLULE	not available	339 118-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 50 MG, GÉLULE	not available	34009 300 022 7 8	SANOFI-AVENTIS 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
TOPALGIC 50 MG, GÉLULE	not available	558 878-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 50 MG, GÉLULE	not available	339 118-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC 50 MG, GÉLULE	not available	34009 300 022 7 8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 621-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	560 421-1	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 141-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 mg, comprimé à libération prolongée	DE/H/0108/001	560 422-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 142-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 622-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 621-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	560 421-1	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 141-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 mg, comprimé à libération prolongée	DE/H/0108/001	560 422-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 142-7	SANOFI-AVENTIS 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
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 622-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 621-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	560 421-1	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 141-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 mg, comprimé à libération prolongée	DE/H/0108/001	560 422-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 142-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 622-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 621-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	560 421-1	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 141-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 mg, comprimé à libération prolongée	DE/H/0108/001	560 422-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 142-7	SANOFI-AVENTIS 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
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 622-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 621-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	560 421-1	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 141-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 mg, comprimé à libération prolongée	DE/H/0108/001	560 422-8	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	343 142-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 100 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/001	351 622-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	560 423-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 623-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 143-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 145-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 mg, comprimé à libération prolongée	DE/H/0108/002	560 424-0	SANOFI-AVENTIS 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
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 624-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	560 423-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 623-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 143-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 145-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 mg, comprimé à libération prolongée	DE/H/0108/002	560 424-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 624-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	560 423-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 623-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 143-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 145-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 mg, comprimé à libération prolongée	DE/H/0108/002	560 424-0	SANOFI-AVENTIS 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
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 624-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	560 423-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 623-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 143-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 145-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 mg, comprimé à libération prolongée	DE/H/0108/002	560 424-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 624-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	560 423-4	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 623-0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 143-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	343 145-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 150 mg, comprimé à libération prolongée	DE/H/0108/002	560 424-0	SANOFI-AVENTIS 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
TOPALGIC LP 150 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/002	351 624-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 625-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	560 425-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	343 146-2	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 627-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	34009 343 147 9 7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 mg, comprimé à libération prolongée	DE/H/0108/003	34009 5604263 0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 625-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	560 425-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	343 146-2	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 627-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	34009 343 147 9 7	SANOFI-AVENTIS 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
TOPALGIC LP 200 mg, comprimé à libération prolongée	DE/H/0108/003	34009 5604263 0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 625-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	560 425-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	343 146-2	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 627-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	34009 343 147 9 7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 mg, comprimé à libération prolongée	DE/H/0108/003	34009 5604263 0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 625-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	560 425-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	343 146-2	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 627-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	34009 343 147 9 7	SANOFI-AVENTIS 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
TOPALGIC LP 200 mg, comprimé à libération prolongée	DE/H/0108/003	34009 5604263 0	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 625-3	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	560 425-7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	343 146-2	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	351 627-6	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 MG, COMPRIMÉ À LIBÉRATION PROLONGÉE	DE/H/0108/003	34009 343 147 9 7	SANOFI-AVENTIS FRANCE	FR
TOPALGIC LP 200 mg, comprimé à libération prolongée	DE/H/0108/003	34009 5604263 0	SANOFI-AVENTIS FRANCE	FR
Toram 50 mg kapsule	not available	UP/I-530-09/12-01/164	FARMAL DD.	HR
Tradol 50 mg Effervescent Tablets	not available	PA 711/29/2	ROWEX LTD	IE
Tradol 50mg Hard Capsules	not available	PA 711/29/1	ROWEX LTD	IE
Tradol 50mg/ml Solution for Injection or Infusion	not available	PA 711/29/3	ROWEX LTD	IE
Tradol SR 100 mg Prolonged Release Tablets	not available	PA 711/29/5	ROWEX LTD	IE
Tradol SR 150 mg Prolonged Release Tablets	not available	PA 711/29/6	ROWEX LTD	IE
Tradol SR 200 mg Prolonged Release Tablets	not available	PA 711/29/7	ROWEX LTD	IE
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

Product 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 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, comprimate filmate	not available	8300/2006/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
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 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 prelungita	not available	4765/2012/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Tradolan Retard 150 mg comprimate cu eliberare prelungita	not available	4765/2012/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
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 prelungita	not available	4766/2012/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Tradolan Retard 200 mg comprimate cu eliberare prelungita	not available	4766/2012/02	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 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-Filmtabletten	not available	1-21239	G.L. PHARMA GMBH	AT
Tradolan Retard, depottabletter	AT/H/0117/003	32518	G.L. PHARMA GMBH	DK
Tradolan Retard, depottabletter	AT/H/0117/001	32516	G.L. PHARMA GMBH	DK
Tradolan Retard, depottabletter	AT/H/0117/002	32517	G.L. PHARMA GMBH	DK
Tradolan, tabletter, filmovertrukne	not available	18233	G.L. PHARMA GMBH	DK
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	MEDA PHARMA S.L.	ES
Tradonal retard 150 mg cápsulas duras de liberación prolongada	UK/H/0225/003	62.112	MEDA PHARMA S.L.	ES
Tradonal retard 200 mg cápsulas duras de liberación prolongada	UK/H/0225/004	62.113	MEDA PHARMA S.L.	ES
Tradonal retard 50 mg cápsulas duras de liberación prolongada	UK/H/0225/001	62.110	MEDA PHARMA S.L.	ES
TRADONAL ODIS 50 mg comprimés orodispersibles	UK/H/0640/001	BE267032	S.A. MEDA PHARMA N.V.	BE
TRADONAL ODIS 50 mg comprimés orodispersibles.	UK/H/0640/001	2004/110026	S.A. MEDA PHARMA N.V.	LU
TRADONAL ODIS 50 mg orodispergeerbare tabletten	UK/H/0640/001	BE 267032	S.A. MEDA PHARMA N.V.	BE
TRADONAL ODIS 50 mg Schmelztabletten	UK/H/0640/001	BE267032	S.A. MEDA PHARMA N.V.	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRADONAL ODIS 50 mg Schmelztabletten	UK/H/0640/001	2004/110026	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	BE195447	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	2011/010940	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 100 mg, capsules met verlengde afgifte, hard	UK/H/0225/002	BE195447	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 100 mg, gélules à libération prolongée	UK/H/0225/002	BE195447	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 100 mg, gélules à libération prolongée	UK/H/0225/002	2011/010940	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	BE195456	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	2011/010941	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 150 mg, capsules met verlengde afgifte, hard	UK/H/0225/003	BE195456	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 150 mg, gélules à libération prolongée	UK/H/0225/003	BE195456	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 150 mg, gélules à libération prolongée	UK/H/0225/003	2011/010941	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 200 mg Hartkapseln, retardiert	UK/H/0225/004	BE195465	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 200 mg Hartkapseln, retardiert	UK/H/0225/004	2011/010942	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 200 mg, capsules met verlengde afgifte, hard	UK/H/0225/004	BE195465	S.A. MEDA PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRADONAL Retard 200 mg, gélules à libération prolongée	UK/H/0225/004	BE195465	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 200 mg, gélules à libération prolongée	UK/H/0225/004	2011/010942	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 50 mg Hartkapseln, retardiert	UK/H/0225/001	BE195377	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 50 mg Hartkapseln, retardiert	UK/H/0225/001	2011/010939	S.A. MEDA PHARMA N.V.	LU
TRADONAL Retard 50 mg, capsules met verlengde afgifte, hard	UK/H/0225/001	BE195377	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 50 mg, gélules à libération prolongée	UK/H/0225/001	BE195377	S.A. MEDA PHARMA N.V.	BE
TRADONAL Retard 50 mg, gélules à libération prolongée	UK/H/0225/001	2011/010939	S.A. MEDA PHARMA N.V.	LU
Tradonal Smelt, 50 mg smelttabletten	UK/H/0640/001	RVG 30752	MEDA PHARMA B.V.	NL
TRADONAL, 100 mg, Injektionslösung	not available	BE 177615	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 100 mg, Injektionslösung	not available	2003/087561	S.A. MEDA PHARMA N.V.	LU
TRADONAL, 100 mg, oplossing voor injectie	not available	BE177615	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 100 mg, solution injectable	not available	BE177615	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 100 mg, solution injectable	not available	2003/087561	S.A. MEDA PHARMA N.V.	LU
TRADONAL, 100 mg/ml, druppels voor oraal gebruik	not available	BE177633	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 100 mg/ml, solution buvable en gouttes	not available	BE177633	S.A. MEDA PHARMA N.V.	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRADONAL, 100 mg/ml, solution buvable en gouttes	not available	2003/087560	S.A. MEDA PHARMA N.V.	LU
TRADONAL, 100 mg/ml, Tropfen zum Einnehmen	not available	BE 177633	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 100 mg/ml, Tropfen zum Einnehmen	not available	2003/087560	S.A. MEDA PHARMA N.V.	LU
TRADONAL, 50 mg, capsules	not available	BE 177642	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 50 mg, gélules	not available	BE 177642	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 50 mg, gélules	not available	2003/087559	S.A. MEDA PHARMA N.V.	LU
TRADONAL, 50 mg, Kapseln	not available	BE 177642	S.A. MEDA PHARMA N.V.	BE
TRADONAL, 50 mg, Kapseln	not available	2003/087559	S.A. MEDA PHARMA N.V.	LU
TRADONAL™ S.R. 100 mg Capsule	UK/H/0225/002	034233027	MEDA PHARMA S.P.A.	IT
TRADONAL™ S.R. 100 mg Capsule	UK/H/0225/002	034233066	MEDA PHARMA S.P.A.	IT
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
TRADONAL™ S.R. 50 mg Capsule	UK/H/0225/001	034233054	MEDA PHARMA S.P.A.	IT
TRADONAL™ S.R. 50 mg Capsule	UK/H/0225/001	034233015	MEDA PHARMA S.P.A.	IT
Tradorec XL 100 mg prolonged-release tablets	FR/H/0272/001	PL 43808/0001	ENDO VENTURES LIMITED	UK
Tradorec XL 200 mg prolonged-release tablets	FR/H/0272/002	PL 43808/0002	ENDO VENTURES LIMITED	UK
Tradorec XL 300 mg prolonged-release tablets	FR/H/0272/003	PL 43808/0003	ENDO VENTURES LIMITED	UK
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672018	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
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672020	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672032	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672044	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672057	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672069	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672083	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672095	MEDA PHARMA S.P.A.	IT
TRAFLASH 50 mg compressa orodispersibile	UK/H/0640/001	036672071	MEDA PHARMA S.P.A.	IT
TRALGIT 100 INJ injekční roztok tramadoli hydrochloridum	not available	65/028/04-C	ZENTIVA, A.S.	CZ
TRALGIT 100 INJ injekční roztok tramadoli hydrochloridum	not available	65/028/04-C	ZENTIVA, A.S.	CZ
TRALGIT 100 INJ injekční roztok tramadoli hydrochloridum	not available	65/028/04-C	ZENTIVA, A.S.	CZ
TRALGIT 100 inj. injekčný roztok	not available	65/0172/03-S	ZENTIVA, A.S.	SK
TRALGIT 100 inj. injekčný roztok	not available	65/0172/03-S	ZENTIVA, A.S.	SK
TRALGIT 100 inj. injekčný roztok	not available	65/0172/03-S	ZENTIVA, A.S.	SK
TRALGIT 50 INJ injekční roztok tramadoli hydrochloridum	not available	65/027/04-C	ZENTIVA, A.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRALGIT 50 INJ injekční roztok tramadoli hydrochloridum	not available	65/027/04-C	ZENTIVA, A.S.	CZ
TRALGIT 50 INJ injekční roztok tramadoli hydrochloridum	not available	65/027/04-C	ZENTIVA, A.S.	CZ
TRALGIT 50 mg kapsuly	not available	65/0169/03-S	ZENTIVA, A.S.	SK
TRALGIT 50 mg kapsuly	not available	65/0169/03-S	ZENTIVA, A.S.	SK
TRALGIT gtt 100 mg/ml perorálne roztokové kvapky	not available	65/0170/03-S	ZENTIVA, A.S.	SK
TRALGIT gtt 100 mg/ml perorálne roztokové kvapky	not available	65/0170/03-S	ZENTIVA, A.S.	SK
TRALGIT gtt. perorální kapky, roztok	not available	65/195/03-C	ZENTIVA, A.S.	CZ
TRALGIT gtt. perorální kapky, roztok	not available	65/195/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 100 100 mg tablety s predĺženým uvoľňovaním	not available	65/0258/02-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 100 100 mg tablety s predĺženým uvoľňovaním	not available	65/0258/02-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 tablety s prodlouženým uvolňováním	not available	65/357/01-C	ZENTIVA, A.S.	CZ
TRALGIT SR 100 tablety s prodlouženým uvolňováním	not available	65/357/01-C	ZENTIVA, A.S.	CZ
TRALGIT SR 100 tablety s prodlouženým uvolňováním	not available	65/357/01-C	ZENTIVA, A.S.	CZ
TRALGIT SR 100 tablety s prodlouženým uvolňováním	not available	65/357/01-C	ZENTIVA, A.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRALGIT SR 150 150 mg tablety s predĺženým uvoľňovaním	not available	65/0259/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 150 150 mg tablety s predĺženým uvoľňovaním	not available	65/0259/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 150 150 mg tablety s predĺženým uvoľňovaním	not available	65/0259/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 150 150 mg tablety s predĺženým uvoľňovaním	not available	65/0259/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 150 tablety s prodlouženým uvolňováním	not available	65/048/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 150 tablety s prodlouženým uvolňováním	not available	65/048/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 150 tablety s prodlouženým uvolňováním	not available	65/048/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 150 tablety s prodlouženým uvolňováním	not available	65/048/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 200 200 mg tablety s predĺženým uvoľňovaním	not available	65/0260/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 200 200 mg tablety s predĺženým uvoľňovaním	not available	65/0260/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 200 200 mg tablety s predĺženým uvoľňovaním	not available	65/0260/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 200 200 mg tablety s predĺženým uvoľňovaním	not available	65/0260/02-S	ZENTIVA, A.S.	SK
TRALGIT SR 200 tablety s prodlouženým uvolňováním	not available	65/049/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 200 tablety s prodlouženým uvolňováním	not available	65/049/03-C	ZENTIVA, A.S.	CZ

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRALGIT SR 200 tablety s prodlouženým uvolňováním	not available	65/049/03-C	ZENTIVA, A.S.	CZ
TRALGIT SR 200 tablety s prodlouženým uvolňováním	not available	65/049/03-C	ZENTIVA, A.S.	CZ
Tralodie® 100 mg Capsule rigide a rilascio prolungato	FI/H/0164/001	035986013	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 100 mg Capsule rigide a rilascio prolungato	FI/H/0164/001	035986025	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 100 mg Capsule rigide a rilascio prolungato	FI/H/0164/001	035986037	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 100 mg Capsule rigide a rilascio prolungato	FI/H/0164/001	035986049	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato	FI/H/0164/002	035986052	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato	FI/H/0164/002	035986064	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato	FI/H/0164/002	035986076	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 150 mg Capsule rigide a rilascio prolungato	FI/H/0164/002	035986088	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato	FI/H/0164/003	035986090	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato	FI/H/0164/003	035986102	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato	FI/H/0164/003	035986114	THERABEL GIENNE PHARMA S.P.A.	IT
Tralodie® 200 mg Capsule rigide a rilascio prolungato	FI/H/0164/003	035986126	THERABEL GIENNE PHARMA S.P.A.	IT
Tramabene - Tropfen	not available	1-21799	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 100 mg - Ampullen	not available	1-21797	RATIOPHARM ARZNEIMITTEL VERTRIEBS-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
Tramabene 100 mg – Retardtabletten	not available	1-23913	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 150 mg Retardtabletten	NL/H/0892/002	1-27076	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 150 mg Retardtabletten	NL/H/0892/002	1-27076	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 200 mg Retardtabletten	NL/H/0892/003	1-27077	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 200 mg Retardtabletten	NL/H/0892/003	1-27077	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 50 mg - Ampullen	not available	1-21796	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 50 mg - Kapseln	not available	1-21798	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Tramabene 50 mg kapsuly	not available	65/0224/98-S	RATIOPHARM GMBH	SK
TRAMABENE 50 tobolky	not available	65/156/99-C	RATIOPHARM GMBH	CZ
TRAMABENE kapky	not available	65/155/99-C	RATIOPHARM GMBH	CZ
Tramabene Perorálne kvapky	not available	65/0120/99-S	RATIOPHARM GMBH	SK
Tramabeta	2125528	25528.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 100 mg	2144604	44604.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 150 mg	2144605	44604.01.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tramabeta long 200 mg	2144606	44604.02.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Tram-ac 50 mg	DE/H/0144/001	37203.00.00	HEXAL AG	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
Tramadex 50 mg Capsules	not available	19207	DELORBIS PHARMACEUTICALS LTD	CY
Tramadin 50 mg kapseli, kova	not available	12346	RATIOPHARM GMBH	FI
Tramadin 50 mg kapseli, kova	not available	12346	RATIOPHARM GMBH	FI
TRAMADOL	not available	5495/2005/01	KRKA, D.D., NOVO MESTO	RO
TRAMADOL MEDA 50mg CÁPSULAS	not available	2830982	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Tramadol "Actavis", kapsler, hårde	UK/H/0380/001	31829	ACTAVIS GROUP PTC EHF.	DK
Tramadol "Aurobindo", hårde kapsler	NL/H/2480/001	49659	AUROBINDO PHARMA (MALTA) LIMITED	DK
Tramadol "Vitabalans", tabletter	FI/H/0779/001	49370	VITABALANS OY	DK
Tramadol 100 injekt - 1A-Pharma®	not available	32751.00.00	1 A PHARMA GMBH	DE
TRAMADOL 100 mg comprimate cu eliberare prelungita	not available	4146/2011/01	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 MG-ROTEXMEDICA	not available	38543.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
TRAMADOL 100 MG-ROTEXMEDICA	not available	38543.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
TRAMADOL 100 MG-ROTEXMEDICA	not available	38543.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
TRAMADOL 100 MG-ROTEXMEDICA	not available	38543.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
Tramadol 100 ret - 1A Pharma®	not available	49470.00.00	1 A PHARMA GMBH	DE
Tramadol 150 ret - 1A Pharma®	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

Product 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 1A Pharma 50 mg - Kapseln	not available	1-24617	1A PHARMA GMBH	AT
Tramadol 200 ret - 1A Pharma®	not available	49470.02.00	1 A PHARMA GMBH	DE
Tramadol 50 Heumann	-	25538.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Tramadol 50 Kapseln - 1A-Pharma, Hartkapseln	not available	32751.00.02	1 A PHARMA GMBH	DE
Tramadol 50 mg Capsules	not available	PL 17907/0110	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Tramadol 50 mg Capsules	UK/H/5829/001	PL 33414/0127	CHELONIA HEALTHCARE LIMITED	UK
Tramadol 50 mg capsules, hard	NL/H/2480/001	PL 16363/0335	MILPHARM LIMITED	UK
TRAMADOL 50 MG-ROTEXMEDICA	not available	38449.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
TRAMADOL 50 MG-ROTEXMEDICA	not available	38449.00.00	ROTEXMEDICA GMBH ARZNEIMITTELWERK	DE
Tramadol 50 tabs - 1 A Pharma, 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	PL 20395 / 0065	RELON CHEM LIMITED	UK
Tramadol 50mg Capsules	not available	PL 00289/1603	TEVA UK 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 AbZ 50 mg/ml Injektionslösung	not available	32307.00.00	ABZ-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 acis Tropfen	not available	35169.00.00	ACIS ARZNEIMITTEL GMBH	DE
Tramadol actas 100 mg Retardtabletten	not available	49469.00.00	ARISTO PHARMA GMBH (ART 57)	DE
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	FR/H/0403/001/DC	IS/1/09/064/01	ACTAVIS GROUP PTC EHF.	IS
TRAMADOL ACTAVIS 100 MG/ML BELSŐLEGES OLDATOS CSEPPEK	not available	OGYI-T-7724/01	ACTAVIS GROUP PTC EHF.	HU
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 kapslar, hårda	UK/H/0380/001	16600	ACTAVIS GROUP PTC EHF.	SE
Tramadol Actavis 50 mg kemény kapszula	OGYI-T-7724/05-07	OGYI-T-7724/05-07	ACTAVIS GROUP PTC EHF.	HU
TRAMADOL ACTAVIS 50 mg, gélule	FR/H/0403/001	NL 37429	ACTAVIS GROUP PTC EHF.	FR
Tramadol Actavis 50 mg/ml oldatos injekció	OGYI-T-7724/03-04	OGYI-T-7724/03-04	ACTAVIS GROUP PTC EHF.	HU
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 Brausetabletten	not available	37019.00.00	ALIUD PHARMA GMBH	DE
Tramadol AL 50 Kapseln	not available	30903.00.02	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
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/03	ALIUD PHARMA GMBH	HU
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/02	ALIUD PHARMA GMBH	HU
Tramadol AL 50 mg kemény kapszula	not available	OGYI-T-7869/01	ALIUD PHARMA GMBH	HU
Tramadol AL Tropfen	not available	30903.00.01	ALIUD PHARMA GMBH	DE
Tramadol Arcana 100 mg Retardtabletten	UK/H/4160/001	1-31515	ARCANA ARZNEIMITTEL GMBH	AT
Tramadol Arcana 150 mg Retardtabletten	UK/H/4160/002	1-31516	ARCANA ARZNEIMITTEL GMBH	AT
Tramadol Arcana 200 mg Retardtabletten	UK/H/4160/003	1-31517	ARCANA ARZNEIMITTEL GMBH	AT
TRAMADOL ARENA 50 mg capsule	not available	9254/2016/02	ARENA GROUP S.A	RO
Tramadol Arena 50 mg capsule	not available	9254/2016/01	ARENA GROUP S.A	RO
Tramadol Aristo 150 mg comprimidos de liberación prolongada EFG	not available	65834	ARISTO PHARMA IBERIA, S.L.	ES
Tramadol Aristo 200 mg comprimidos de liberación prolongada EFG	not available	65835	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 Aristo100 mg comprimidos de liberación prolongada EFG	not available	65833	ARISTO PHARMA IBERIA, S.L.	ES
TRAMADOL ARROW 50 mg, comprimé	not available	26860	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. 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	32723	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	32724	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 200 mg, comprimé à libération prolongée	not available	NL 36435	ARROW GENERIQUES	FR
TRAMADOL ARROW L.P. 200 mg, gélule à libération prolongée	not available	32725	ARROW GENERIQUES	FR
TRAMADOL ASTA Medica 100 mg Solución Inyectable EFG	not available	63.465	MEDA PHARMA S.L.	ES
TRAMADOL ASTA Medica EFG Cápsulas	not available	61.849	MEDA PHARMA S.L.	ES
TRAMADOL ASTA Medica Gotas EFG	not available	62.925	MEDA PHARMA S.L.	ES
Tramadol Aurobindo 50 mg cápsulas duras EFG	NL/H/2480/001	77.656	LABORATORIOS AUROBINDO S.L.U.	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	5032024	AUROVITAS 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 Aurovitas 150 mg comprimidos de libertação prolongada	NL/H/0890/002	5032057	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 200 mg comprimidos de libertação prolongada	NL/H/0890/003	5032107	AUROVITAS UNIPESSOAL, LDA.	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 UNIPESSOAL, LDA.	CZ
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	5703368	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS 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 Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	5703376	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Aurovitas 50 mg/ml solução injetável ou para perfusão	PT/H/1548/001	PT/H/1548/001	AUROVITAS UNIPESSOAL, LDA.	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 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 mg cápsulas	not available	5012331	LABORATÓRIOS AZEVEDOS - INDÚSTRIA 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
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 BIOGARAN 50 mg, gélule	not available	3400935248442	BIOGARAN	FR
TRAMADOL BIOGARAN 50 mg, gélule	not available	3400935248442	BIOGARAN	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 300 141 0 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 550 049 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 128 1 5	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 BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 125 2 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 129 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 121 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 130 6 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 571 534 7 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 127 5 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 131 2 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 122 3 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 120 0 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 124 6 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 132 9 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 571 533 0 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 BIOGARAN LP 100 mg, comprimé à libération prolongée	DE/H/0798/002	34009 382 126 9 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 300 141 1 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 550 049 6 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 223 4 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 225 7 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 228 6 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 220 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 230 0 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 229 2 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 222 8 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 571 539 9 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 231 7 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
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 224 0 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 571 538 2 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 226 3 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 221 1 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 150 mg, comprimé à libération prolongée	DE/H/0798/003	34009 382 232 3 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 300 141 2 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 550 049 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 259 9 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 258 2 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 243 5 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 241 2 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 247 0 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 BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 257 6 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 249 3 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 253 0 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 248 7 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 571 540 7 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 571 541 3 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 245 8 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 254 7 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL BIOGARAN LP 200 mg, comprimé à libération prolongée	DE/H/0798/004	34009 382 255 3 2	LABORATOIRES GRÜNENTHAL S.A.S.	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
Tramadol Ciclum 50 mg cápsulas	not available	2679686	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	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 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
TRAMADOL CRISTERS LP 150 mg, comprimé pelliculé à libération prolongée	not available	34009 578 149 1 5	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 578 151 6 5	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
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

Product 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 200 mg, comprimé pelliculé à libération prolongée	not available	34009 578 170 0 8	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
Tramadol Denk 50 mg Brause	not available	34728.00.00	DENK PHARMA GMBH & CO. KG	DE
Tramadol Denk 50 mg Brause	not available	34728.00.00	DENK PHARMA GMBH & CO. KG	DE
Tramadol Dermogen 50 mg cápsulas duras EFG	not available	65724	DERMOGEN FARMA, S.A.	ES
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/2ml solution injectable	DE/H/0282/004	BE245786	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 EG 100 mg/2ml solution injectable	DE/H/0282/004	0019/08039713	EUROGENERICS N.V./S.A.	LU
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 Brausetabletten	not available	BE232802	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg bruistabletten	not available	BE232802	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg comprimés	UK/H/0305/001	BE216474	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg comprimés effervescents	not available	BE232802	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg tabletten	UK/H/0305/001	BE216474	EUROGENERICS N.V./S.A.	BE
Tramadol EG 50 mg Tabletten	UK/H/0305/001	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 mg, comprimé à libération prolongée	not available	NL37921	EG LABO - LABORATOIRES EUROGENERICS	FR
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

Product 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 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 50 mg, gélule	not available	34009 266 907 8 6	EVOLUPHARM	FR
TRAMADOL EVOLUGEN 50 mg, gélule	not available	34009 266 908 4 7	EVOLUPHARM	FR
TRAMADOL EVOLUGEN 50 mg, gélule	not available	34009 266 910 9 7	EVOLUPHARM	FR
TRAMADOL EVOLUGEN 50 mg, gélule	not available	34009 583 516 9 3	EVOLUPHARM	FR
Tramadol Farmal 100 mg tablete s produljenim oslobađanjem	not available	HR-H-671676641	FARMAL DD.	HR
Tramadol Farmal 150 mg tablete s produljenim oslobađanjem	not available	HR-H-918620020	FARMAL DD.	HR
Tramadol Farmal 200 mg tablete s produljenim oslobađanjem	not available	HR-H-086392384	FARMAL DD.	HR
TRAMADOL FARMALIDER 100 mg/ml, solución oral .E.F.G	not available	69404	FARMALIDER, S.A.	ES
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

Product 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 Phar 50 mg, cápsulas	not available	3759495	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Generis Phar 50 mg, cápsulas	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 100 mg/ml PCH, druppels voor oraal gebruik	not available	RVG 27177	PHARMACHEMIE B.V	NL
Tramadol HCl Actavis Retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0890/001	RVG 33553	ACTAVIS GROUP PTC EHF.	NL
Tramadol HCl Actavis Retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0890/001	RVG 33553	ACTAVIS GROUP PTC EHF.	NL
Tramadol HCl Actavis Retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0890/002	RVG 33554	ACTAVIS GROUP PTC EHF.	NL
Tramadol HCl Actavis Retard 150 mg, tabletten met gereguleerde afgifte	NL/H/0890/002	RVG 33554	ACTAVIS GROUP PTC EHF.	NL
Tramadol HCl Actavis Retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0890/003	RVG 33555	ACTAVIS GROUP PTC EHF.	NL
Tramadol HCl Actavis Retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0890/003	RVG 33555	ACTAVIS GROUP PTC EHF.	NL
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 bruis 50 PCH, bruistabletten 50 mg	not available	RVG 25343	PHARMACHEMIE 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 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
Tramadol HCl Duiven retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0538/001	RVG 30562	ICC B.V.	NL
Tramadol HCl Duiven retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0539/001	RVG 30577	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 150 mg, tabletten met gereguleerde afgifte	NL/H/0539/002	RVG 30578	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 Duiven retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0539/003	RVG 30579	ICC B.V.	NL
Tramadol HCl Mylan 50 mg, capsules	not available	RVG 26280	MYLAN B.V.	NL
Tramadol HCl ratiopharm 50 mg, capsules.	not available	RVG 21682	RATIOPHARM NEDERLAND B.V	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

Product 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 retard 100 mg, tabletten met gereguleerde afgifte	NL/H/0889/001	RVG 32924	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 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/0889/002	RVG 32925	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 200 mg Teva, tabletten met gereguleerde afgifte	not available	RVG 33753	TEVA NEDERLAND B.V.	NL
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/0889/003	RVG 32926	ICC B.V.	NL
Tramadol HCl retard 200 mg, tabletten met gereguleerde afgifte	NL/H/0892/003	RVG 33756	ICC 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 retard CF 100 mg, tabletten met gereguleerde afgifte	not available	RVG 34128	CENTRAFARM B.V.	NL
Tramadol HCl retard CF 100 mg, tabletten met gereguleerde afgifte	not available	RVG 34128	CENTRAFARM B.V.	NL
Tramadol HCl retard CF 200 mg, tabletten met gereguleerde afgifte	not available	RVG 34130	CENTRAFARM B.V.	NL
Tramadol HCl retard CF 200 mg, tabletten met gereguleerde afgifte	not available	RVG 34130	CENTRAFARM 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 verlengde afgifte	UK/H/4160/001	RVG 109064	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 verlengde afgifte	UK/H/4160/002	RVG 109065	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

Product 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 Retard Mylan 200 mg, tabletten met verlengde afgifte	UK/H/4160/003	RVG 109066	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 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 Heumann Tropfen	-	25530.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Tramadol Hexal 100 mg kapslar, hårda	not available	13466	HEXAL A/S	SE
Tramadol HEXAL 100 mg, kapseli, kova	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, kapseli, kova	not available	12729	HEXAL A/S	FI
Tramadol Hikma 100 mg Injektionslösung	not available	30181.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Tramadol Hikma 100 mg Injektionslösung	not available	30181.00.00	HIKMA FARMACÊUTICA (PORTUGAL), 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
Tramadol Hydrochloride 50 mg Capsules	not available	PL 44041/0037	NOUMED LIFE SCIENCES	UK
Tramadol hydrochloride 50 mg capsules	not available	PL 17780/0139	WINTHROP PHARMACEUTICALS UK LTD	UK
Tramadol hydrochloride 50 mg capsules	not available	PL 17780/0139	WINTHROP PHARMACEUTICALS UK LTD	UK
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 PHARMACEUTICALS LTD	UK
Tramadol Hydrochloride 50mg Capsules	not available	PL 20117/0086	MORNINGSIDE HEALTHCARE 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	not available	PL 11311/0084	TILLOMED LABORATORIES LTD	UK
TRAMADOL HYDROCHLORIDE 50mg CAPSULES	UK/H/0380/001	PL 0142/0484	ACTAVIS UK LTD.	UK
Tramadol hydrochloride 50mg Capsules	not available	PL 10622/0050	PLIVA PHARMA LIMITED	UK
Tramadol Hydrochloride 50mg Effervescent Tablets	not available	PL 28444/0217	ACTIVASE PHARMACEUTICALS LIMITED	UK
Tramadol Hydrochloride 50mg Effervescent Tablets	not available	PL 36722/0119	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Tramadol Hydrochloride 50mg Effervescent Tablets	not available	PL 36722/0119	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
TRAMADOL HYDROCHLORIDE 50MG HARD CAPSULES	FR/H/0403/001	PA 1380/96/1	ACTAVIS GROUP PTC EHF.	IE
Tramadol Hydrochloride 50mg Hard Capsules	FR/H/0403/001/DC	PA 1380/96/1	ACTAVIS GROUP PTC EHF.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramadol Hydrochloride Capsules 50mg	not available	AA054/06901	ACCORD HEALTHCARE LIMITED	MT
Tramadol Hydrochloride Capsules 50mg	not available	PL 20075/0290	ACCORD HEALTHCARE LIMITED	UK
Tramadol Kalceks 100 mg/2 ml šķīdums injekcijām	not available	16-0115	KALCEKS	LV
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 ilgstošās darbības tabletes	not available	00-0666	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 100 mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1055/007	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 100 mg žvakutės	not available	LT/1/94/1055/002	KRKA, D.D., NOVO MESTO	LT
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 otopina za injekcije	not available	UP/I-530-09/11-02/292	KRKA-FARMA D.O.O.	HR
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 oralne kapi	not available	UP/I-530-09/11-02/290	KRKA-FARMA D.O.O.	HR
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 200 mg ilgstošās darbības tabletes	not available	03-0207	KRKA, D.D., NOVO MESTO	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
Tramadol Krka 200mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1055/009	KRKA, D.D., NOVO MESTO	LT
Tramadol Krka 50 mg cietās kapsulas	not available	96-0110	KRKA, D.D., NOVO MESTO	LV
Tramadol Krka 50 mg kapsule	not available	UP/I-530-09/11-02/289	KRKA-FARMA D.O.O.	HR
Tramadol Krka 50 mg kietosios kapsulės	not available	LT/1/94/1055/001	KRKA, D.D., NOVO MESTO	LT
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 otopina za injekcije	not available	UP/I-530-09/11-02/291	KRKA-FARMA D.O.O.	HR
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, 100 mg rektaalsuposiidid	not available	139696	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 100 mg toimeainet prolongeeritult vabastavad tabletid	not available	331200	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 100 mg/ml suukaudsed tilgad, lahus	not available	148696	KRKA, D.D., NOVO MESTO	EE
Tramadol Krka, 50 mg kõvakapslid	not available	139596	KRKA, D.D., NOVO MESTO	EE
Tramadol Labesfal, 100 mg, comprimidos de libertação prolongada	not available	4348587	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Labesfal, 100 mg/2 ml, solução injectável	not available	3210994	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 100 mg/2 ml, solução injectável	not available	4842399	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 50 mg cápsulas	not available	3827599	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
Tramadol Labesfal, 50 mg cápsulas	not available	3827698	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol Labesfal, 50 mg/1 ml, solução injectável	not available	3210796	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Tramadol Labesfal, 50 mg/1 ml, solução injectável	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 suposiidid	not available	321800	G.L. PHARMA GMBH	EE
Tramadol Lannacher 100 mg supozitoriji	not available	99-0232	G.L. PHARMA GMBH	LV
Tramadol Lannacher 100 mg žvakutes	not available	LT/1/97/1431/009	G.L. PHARMA GMBH	LT
Tramadol Lannacher 100 mg, toimeainet prolongeeritult vabastavad tabletid	not available	345101	G.L. PHARMA GMBH	EE
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

Product 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/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 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

Product 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, 50 mg õhukese polümeerikilega kaetud tabletid	not available	274199	G.L. PHARMA GMBH	EE
TRAMADOL LAVOISIER® 50 mg/ ml, solution injectable	not available	VNL35853-3400957486020	LABORATOIRES CHAIX ET DU MARAIS	FR
TRAMADOL LAVOISIER® 50 mg/ ml, solution injectable	not available	VNL35853-3400957486020	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 Einnehmen	not available	13144.00.02	LIBRA-PHARM GMBH	DE
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	not available	32751.00.01	1 A PHARMA GMBH	DE
TRAMADOL MABO 50 mg cápsulas E.F.G.	not available	63.758	MABO-FARMA, 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 MEDA 100mg/2ml SOLUÇÃO INJECTÁVEL	not available	2831188	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAMADOL MEDA 100mg/ml GOTAS ORAIS SOLUÇÃO	not available	3081080	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAMADOL MEDA 100mg/ml GOTAS ORAIS SOLUÇÃO	not available	2831089	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463179	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463211	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463328	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463302	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463203	MYLAN, LDA	PT
Tramadol Mylan 100 mg comprimidos de libertação prolongada	UK/H/4160/001	5463310	MYLAN, LDA	PT
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN DURA GMBH	DE
Tramadol Mylan 100 mg Retardtabletten	not available	39923.00.00	MYLAN DURA GMBH	DE
Tramadol Mylan 100 mg Tablety s predĺženým uvoľňovaním	UK/H/4160/001	65/0556/12-S	GENERICS [UK] LIMITED	SK
Tramadol Mylan 100 mg, tablety s prodlouženým uvolňováním	UK/H/4160/001	65/120/13-C	GENERICS [UK] 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
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 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463245	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463229	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463336	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463237	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463344	MYLAN, LDA	PT
Tramadol Mylan 150 mg comprimidos de libertação prolongada	UK/H/4160/002	5463351	MYLAN, LDA	PT
Tramadol Mylan 150 mg Tablety s predĺženým uvoľňovaním	UK/H/4160/002	65/0557/12-S	GENERICS [UK] LIMITED	SK
Tramadol Mylan 150 mg tablety s prodlouženým uvolňováním	UK/H/4160/002	65/121/13-C	GENERICS [UK] LIMITED	CZ
Tramadol Mylan 200 mg comprimidos de libertação prolongada	UK/H/4160/003	5463260	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	UK/H/4160/003	5463252	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	UK/H/4160/003	5463369	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	UK/H/4160/003	5463377	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	UK/H/4160/003	5463401	MYLAN, LDA	PT
Tramadol Mylan 200 mg comprimidos de libertação prolongada	UK/H/4160/003	5463278	MYLAN, LDA	PT
Tramadol Mylan 200 mg Tablety s predĺženým uvoľňovaním	UK/H/4160/003	65/0558/12-S	GENERICS [UK] LIMITED	SK
Tramadol Mylan 200 mg tablety s prodlouženým uvolňováním	UK/H/4160/003	65/122/13-C	GENERICS [UK] LIMITED	CZ
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé pelliculé à libération prolongée	not available	NL 39453	MYLAN S.A.S	FR
TRAMADOL MYLAN PHARMA LP 100 mg, comprimé pelliculé à libération prolongée	not available	NL 39453	MYLAN S.A.S	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé pelliculé à libération prolongée	not available	NL 39454	MYLAN S.A.S	FR
TRAMADOL MYLAN PHARMA LP 150 mg, comprimé pelliculé à libération prolongée	not available	NL 39454	MYLAN S.A.S	FR
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé pelliculé à libération prolongée	not available	NL 39455	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
TRAMADOL MYLAN PHARMA LP 200 mg, comprimé pelliculé à libération prolongée	not available	NL 39455	MYLAN S.A.S	FR
TRAMADOL NORMON 100 mg/2 ml Solución inyectable EFG	not available	63.734	LABORATORIOS NORMON, S.A.	ES
TRAMADOL NORMON 50 mg Cápsulas EFG	not available	63910	LABORATORIOS NORMON, S.A.	ES
TRAMADOL RANBAXY 50 MG CAPSULAS EFG	not available	64.305	LABORATORIOS RANBAXY S.L.	ES
TRAMADOL RANBAXY L.P. 100 mg, comprimé à libération prolongée	not available	NL 34291	RANBAXY PHARMACIE GENERIQUES	FR
TRAMADOL RANBAXY L.P. 150 mg, comprimé à libération prolongée	not available	NL 34292	RANBAXY PHARMACIE GENERIQUES	FR
TRAMADOL RANBAXY L.P. 200 mg, comprimé à libération prolongée	not available	NL 34 293	RANBAXY PHARMACIE GENERIQUES	FR
Tramadol ratiopharm 50 mg cápsulas duras EFG	not available	63.472	RATIOPHARM ESPANA SA	ES
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 "Hexal"	DE/H/0448/001	36837	HEXAL A/S	DK
Tramadol Retard "Actavis", depottabletter	NL/H/0890/001	40152	ACTAVIS GROUP HF.	DK
Tramadol Retard "Actavis", depottabletter	NL/H/0890/002	40153	ACTAVIS GROUP HF.	DK
Tramadol Retard "Actavis", depottabletter	NL/H/0890/003	40154	ACTAVIS GROUP HF.	DK
Tramadol Retard 100 mg, capsules met verlengde afgifte, hard	UK/H/0225/002	RVG 22326	MEDA PHARMA B.V.	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramadol Retard 100 mg, capsules met verlengde afgifte, hard*	not available	RVG 23007 = RVG 22326	MEDA PHARMA B.V.	NL
Tramadol Retard 150 mg comprimate cu eliberare prelungită	not available	6137/2014/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Retard 150 mg, capsules met verlengde afgifte, hard	UK/H/0225/003	RVG 22327	MEDA PHARMA B.V.	NL
Tramadol Retard 150 mg, capsules met verlengde afgifte, hard*	not available	RVG 23008=22327	MEDA PHARMA B.V.	NL
Tramadol Retard 200 mg comprimate cu eliberare prelungita	not available	6138/2014/01	KRKA, D.D., NOVO MESTO	RO
Tramadol Retard 200 mg, capsules met verlengde afgifte, hard	UK/H/0225/004	RVG 22328	MEDA PHARMA B.V.	NL
Tramadol Retard 200 mg, capsules met verlengde afgifte, hard*	not available	RVG 23009=22328	MEDA PHARMA B.V.	NL
Tramadol Retard 50 mg, capsules met verlengde afgifte, hard	UK/H/0225/001	RVG 22325	MEDA PHARMA B.V.	NL
Tramadol Retard 50 mg, capsules met verlengde afgifte, hard*	not available	RVG 23006=22325	MEDA PHARMA B.V.	NL
Tramadol Retard Actavis 100 mg	NL/H/0890/001	65/199/07-C	ACTAVIS GROUP HF.	CZ
Tramadol Retard Actavis 100 mg depottabletter	NL/H/0890/001	24560	ACTAVIS GROUP PTC EHF.	SE
Tramadol Retard Actavis 100 mg tablety s predlženým uvolnovaním	NL/H/0890/001	65/0216/07-S	ACTAVIS GROUP HF.	SK
Tramadol Retard Actavis 150 mg depottabletter	NL/H/0890/002	24561	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
Tramadol Retard Actavis 150 mg tablety s predĺženým uvoľňovaním	NL/H/0890/002	65/0217/07-S	ACTAVIS GROUP HF.	SK
Tramadol Retard Actavis 200 mg depottabletter	NL/H/0890/003	24562	ACTAVIS GROUP PTC EHF.	SE
Tramadol Retard Actavis 200 mg tablety s predĺženým uvoľňovaním	NL/H/0890/003	65/0218/07-S	ACTAVIS GROUP HF.	SK
Tramadol Retard Apotex 100 mg comprimidos de liberación prolongada EFG	NL/H/0539/001	72427	APOTEX EUROPE B.V.	ES
Tramadol Retard Apotex 150 mg comprimidos de liberación prolongada EFG	NL/H/0539/002	71710	APOTEX EUROPE B.V.	ES
Tramadol Retard Apotex 200 mg comprimidos de liberación prolongada EFG.	NL/H/0539/003	72.430.	APOTEX EUROPE B.V.	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 Retardtabletten	NL/H/0888/001	BE300212	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 100 mg Retardtabletten	NL/H/0888/001	BE300221	EUROGENERICS N.V./S.A.	BE
Tramadol Retard EG 100 mg Retardtabletten	NL/H/0888/001	BE300203	EUROGENERICS N.V./S.A.	BE
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 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
Tramadol Retard EG 150 mg tabletten met verlengde afgifte	NL/H/0888/002	BE300255	EUROGENERICS N.V./S.A.	BE
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

Product 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 Retardtabletten	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 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
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

Product 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 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	UK/H/4160/001	76572	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 150 mg comprimidos de liberación prolongada EFG	UK/H/4160/002	76573	MYLAN PHARMACEUTICALS S.L.	ES
Tramadol Retard MYLAN 200 mg comprimidos de liberación prolongada EFG	UK/H/4160/003	76574	MYLAN PHARMACEUTICALS S.L.	ES
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 ESPANA SA	ES
Tramadol retard ratiopharm 150 mg comprimidos de liberación prolongada EFG	not available	77218	RATIOPHARM ESPANA SA	ES
Tramadol retard ratiopharm 200 mg comprimidos de liberación prolongada EFG	not available	77219	RATIOPHARM ESPANA SA	ES
Tramadol Retard Sandoz 150 mg depottabletti	NL/H/0483/002	19304	SANDOZ 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 STADA Genéricos 100 mg comprimidos de liberación prolongada EFG	not available	77.200	STADA GENÉRICOS, S.L.	ES
Tramadol retard STADA Genéricos 100 mg comprimidos de liberación prolongada EFG	not available	77.200	STADA GENÉRICOS, S.L.	ES
Tramadol retard STADA Genéricos 150 mg comprimidos de liberación prolongada EFG	not available	77.198	STADA GENÉRICOS, S.L.	ES
Tramadol retard STADA Genéricos 150 mg comprimidos de liberación prolongada EFG	not available	77.198	STADA GENÉRICOS, S.L.	ES
Tramadol retard STADA Genéricos 200 mg comprimidos de liberación prolongada EFG	not available	77.202	STADA GENÉRICOS, S.L.	ES
Tramadol retard STADA Genéricos 200 mg comprimidos de liberación prolongada EFG	not available	77.202	STADA GENÉRICOS, 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 Ritisca 50 mg cápsulas	PT/H/1576/001	5700257	AUROVITAS UNIPESSOAL, LDA.	PT
Tramadol Sandoz 100 mg Brausetabletten	not available	39935.00.00	HEXAL AG	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 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	BE 279325	SANDOZ N.V.	BE
Tramadol Sandoz 100 mg/ml druppels voor oraal gebruik, oplossing	not available	BE 279343	SANDOZ N.V.	BE
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 cápsulas duras EFG	NL/H/0113/002	61.948	SANDOZ FARMACÉUTICA, S.A.	ES
TRAMADOL SANDOZ 50 mg cápsulas duras EFG	NL/H/0113/002	61.948	SANDOZ FARMACÉUTICA, S.A.	ES
Tramadol Sandoz 50 mg capsules, hard	not available	BE254344	SANDOZ N.V.	BE
TRAMADOL SANDOZ 50 mg, comprimé	not available	351 065-8	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	368 193-4	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	351 066-4	SANDOZ	FR
TRAMADOL SANDOZ 50 mg, comprimé	not available	351 068-7	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	383 988-4	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	383 989-0	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
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	383 986-1	SANDOZ	FR
TRAMADOL SANDOZ L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	383 987-8	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	385 524-5	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	385 525-1	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	572 785-3	SANDOZ	FR
TRAMADOL SANDOZ L.P. 150 mg, comprimé pelliculé à libération prolongée	not available	572 787-6	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	383 991-5	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	572 175-0	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	572 176-7	SANDOZ	FR
TRAMADOL SANDOZ L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	383 990-9	SANDOZ	FR
Tramadol Sandoz Retard 200 mg tablety s prodlouženým uvolňováním	DE/H/0576/001	65/392/06-C	SANDOZ GMBH	CZ
Tramadol Sandoz UNO 150 mg Retardtabletten	not available	44610.01.00	HEXAL AG	DE
Tramadol Sandoz UNO 200 mg Retardtabletten	DE/H/0576/001	44610.02.00	HEXAL AG	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 soluţie injectabilă 100 mg/2 ml	not available	2848/2002/01	KRKA, D.D., NOVO MESTO	RO
Tramadol soluţie injectabilă 50 mg/1 ml	not available	2846/2002/01	KRKA, D.D., NOVO MESTO	RO
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	8763/2016/03	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	8763/2016/04	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 100 mg comprimate cu eliberare prelungită	not available	8763/2016/01	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 150 mg comprimate cu eliberare prelungită	not available	8764/2016/03	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 150 mg comprimate cu eliberare prelungită	not available	8764/2016/02	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 150 mg comprimate cu eliberare prelungită	not available	8764/2016/04	ZENTIVA, A.S.	RO
TRAMADOL SR ZENTIVA 150 mg comprimate cu eliberare prelungită	not available	8764/2016/01	ZENTIVA, A.S.	RO
TRAMADOL STADA 50 mg cápsulas duras EFG	not available	64.215	LABORATORIO STADA, S.L.	ES
TRAMADOL STADA 50 mg 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

Product 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 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 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® 150 mg Retardtabletten	AT/H/0117/002	51845.01.00	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	not available	37814.00.00	STADAPHARM GMBH	DE
Tramadol STADA® 50 mg Tabs	not available	37814.00.00	STADAPHARM GMBH	DE
TRAMADOL supozitoare 100 mg	not available	2875/2002/01	KRKA, D.D., NOVO MESTO	RO
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 Tarbis 100 mg comprimidos de liberación prolongada EFG	not available	65.800	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
Tramadol Tarbis 150 mg comprimidos de liberación prolongada EFG	not available	65.801	TARBIS FARMA, S.L.	ES
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 50 mg, comprimé effervescent	not available	NL24341	TEVA SANTÉ	FR
TRAMADOL TEVA 50 mg, gélule	not available	NL25606	TEVA SANTÉ	FR
TRAMADOL TEVA L.P. 100 mg, comprimé pelliculé à libération prolongée	not available	NL38989	TEVA SANTÉ	FR
TRAMADOL TEVA L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	NL39007	TEVA SANTÉ	FR
Tramadol Teva LP 150 mg, comprimé pelliculé à libération prolongée	not available	NL39006	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 FARMACÉUTICA 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 mg tablete	FI/H/0779/001	H/13/01554/002	VITABALANS OY	SI
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

Product 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 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 mg tabletta	FI/H/0779/001	OGYI-T-22315/01	VITABALANS OY	HU
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

Product 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 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 Xantis 100 mg tablety	not available	65/171/07-C	XANTIS PHARMA LIMITED	CZ
Tramadol Xantis 50 mg kapsuly	not available	65/0346/03-S	XANTIS PHARMA LIMITED	SK
Tramadol Xantis 50 mg tvrdé tobolky	not available	65/657/97-C	XANTIS PHARMA LIMITED	CZ
TRAMADOL ZENTIVA 50 mg, comprimé effervescent	not available	561 779-7	SANOFI-AVENTIS FRANCE	FR
TRAMADOL ZENTIVA 50 mg, comprimé effervescent	not available	561 780-5	SANOFI-AVENTIS FRANCE	FR
TRAMADOL ZENTIVA 50 mg, comprimé effervescent	not available	367 488-0	SANOFI-AVENTIS FRANCE	FR
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

Product 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 571 545 9 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
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

Product 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 150 mg, comprimé à libération prolongée	DE/H/1093/003	34009 382 300 9 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
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 libération prolongée	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
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
TRAMADOL ZENTIVA LP 200 mg, comprimé à libération prolongée	DE/H/1093/004	34009 382 312 7 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 382 310 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 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 261 3 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
TRAMADOL ZENTIVA LP 50 mg, comprimé à libération prolongée	DE/H/1093/001	34009 382 267 1 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
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 mg, gélule	UK/H/0380/001	34009 357 133 5 3	ZYDUS FRANCE	FR
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

Product 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 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. 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

Product 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 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. 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 prolongée.	not available	34009 394 178 9 9	ZYDUS FRANCE	FR
Tramadol-CT 100 mg Retardkapseln	not available	62755.00.00	ABZ-PHARMA GMBH	DE
Tramadol-CT 100 mg/ml Tropfen Tropfen zum Einnehmen, Lösung	not available	25527.00.00	ABZ-PHARMA GMBH	DE
Tramadol-CT 150 mg Retardkapseln	not available	62756.00.00	ABZ-PHARMA GMBH	DE
Tramadol-CT 200 mg Retardkapseln	not available	62756.01.00	ABZ-PHARMA GMBH	DE
Tramadol-CT 50 mg Hartkapseln	not available	25527.00.01	ABZ-PHARMA GMBH	DE
Tramadol-hameln 100 mg	ENR2138540	35342.01.00	HAMELN PHARMA PLUS 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
Tramadolhydrochlorid Actavis 100 mg Retardtabletten	NL/H/0890/001	1-26983	ACTAVIS GROUP PTC EHF.	AT
Tramadolhydrochlorid Actavis 150 mg Retardtabletten	NL/H/0890/002	1-26986	ACTAVIS GROUP PTC EHF.	AT
Tramadolhydrochlorid Actavis 200 mg Retardtabletten	NL/H/0890/003	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	MEDA PHARMA B.V.	NL
Tramadolis SANITAS 50 mg/ml injekcinis/infuzinis tirpalas	not available	LT/1/2000/1577/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Tramadolis SANITAS 50 mg/ml injekcinis/infuzinis tirpalas	not available	LT/1/2000/1577/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
TRAMADOLO HEXAL	not available	033998 030	SANDOZ S.P.A.	IT
TRAMADOLO ABC 100 mg/ml gocce orali, soluzione	not available	037941010	ABC FARMACEUTICI S.P.A.	IT
TRAMADOLO ANGELINI, 100 mg supposte	not available	035918059	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TRAMADOLO ANGELINI, 100 mg/2 ml soluzione iniettabile	not available	035918034	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TRAMADOLO ANGELINI, 100mg/ml gocce orali, soluzione	not available	035918046	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TRAMADOLO ANGELINI, 50 mg capsule rigide	not available	035918010	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TRAMADOLO ANGELINI, 50 mg/ml soluzione iniettabile	not available	035918022	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Tramadolo EG 100 mg soluzione iniettabile	DE/H/0282/004	035847096	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
Tramadolo EG 100 mg soluzione iniettabile	DE/H/0282/004	035847084	EG S.P.A.	IT
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 SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847021	EG SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847058	EG SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847072	EG SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847019	EG SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847033	EG SPA	IT
Tramadolo EG 100 mg/ml gocce orali soluzione	DE/H/0282/002	035847045	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847185	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847197	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847247	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847211	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847223	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847235	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847250	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847110	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847146	EG 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
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847122	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847209	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847159	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847173	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847134	EG SPA	IT
Tramadolo EG 50 mg capsule rigide	DE/H/0282/001	035847161	EG SPA	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 prolungato	NL/H/0538/001	037133081	GERMED PHARMA S.R.L.	IT
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

Product 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	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
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

Product 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 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 mg compresse a rilascio prolungato	NL/H/0538/002	037133206	GERMED PHARMA S.R.L.	IT
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

Product 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 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 prolungato	NL/H/0538/002	037133509	GERMED PHARMA S.R.L.	IT
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

Product 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	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
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

Product 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	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	036697023	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697047	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697011	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697035	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697050	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697074	SANDOZ S.P.A.	IT
Tramadolo HCl Sandoz	NL/H/0483/001	036697062	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 Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976211	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976209	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976173	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976185	MYLAN 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 Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976161	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976197	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976134	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976146	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976096	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976159	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976122	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976108	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976110	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976084	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976072	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976060	MYLAN 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 Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976033	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976019	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976045	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976021	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976058	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976641	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976654	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976704	MYLAN S.P.A.	IT
Tramadolo Mylan 100 mg compresse a rilascio prolungato	UK/H/4160/001	040976716	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976425	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976413	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976399	MYLAN 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 Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976387	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976401	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976375	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976351	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976363	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976348	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976336	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976312	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976247	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976250	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976300	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976298	MYLAN 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 Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976274	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976262	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976223	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976235	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976666	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976678	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976728	MYLAN S.P.A.	IT
Tramadolo Mylan 150 mg compresse a rilascio prolungato	UK/H/4160/002	040976730	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976639	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976615	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976603	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976627	MYLAN 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 Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976591	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976589	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976565	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976577	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976553	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976540	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976538	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976514	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976526	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976502	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976490	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976488	MYLAN 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 Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976476	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976464	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976437	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976452	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976449	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976680	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976692	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976742	MYLAN S.P.A.	IT
Tramadolo Mylan 200 mg compresse a rilascio prolungato	UK/H/4160/003	040976755	MYLAN S.P.A.	IT
TRAMADOLO VIATRIS 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875020	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875018	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875032	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875044	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
TRAMADOLO VIATRIS 100 mg/ml gocce orali, soluzione	DE/H/0306/003	035875057	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg capsule rigide	DE/H/0306/001	035875095	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg capsule rigide	DE/H/0306/001	035875107	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg capsule rigide	DE/H/0306/001	035875121	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg capsule rigide	DE/H/0306/001	035875119	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg/ml soluzione iniettabile	DE/H/0306/002	035875071	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg/ml soluzione iniettabile	DE/H/0306/002	035875069	MEDA PHARMA S.P.A.	IT
TRAMADOLO VIATRIS 50 mg/ml soluzione iniettabile	DE/H/0306/002	035875083	MEDA PHARMA S.P.A.	IT
TRAMADOLOR 100 ID comprimate cu eliberare modificată	not available	3763/2003/01	HEXAL AG	RO
TRAMADOLOR 100 ID comprimate cu eliberare modificată	not available	3763/2003/02	HEXAL AG	RO
Tramadolor 100 mg - Ampullen	not available	1-23700	HEXAL PHARMA GMBH	AT
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 módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/09	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
Tramadolor 150 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-8179/08	SANDOZ HUNGÁRIA KFT	HU
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 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 einmal täglich 100 mg Retardtabletten	FR/H/0272/001	63147.00.00	HEXAL AG	DE
Tramadolor einmal täglich 100 mg Retardtabletten	FR/H/0272/001	63147.00.00	HEXAL AG	DE
Tramadolor einmal täglich 200 mg Retardtabletten	FR/H/0272/002	63148.00.00	HEXAL AG	DE
Tramadolor einmal täglich 200 mg Retardtabletten	FR/H/0272/002	63148.00.00	HEXAL AG	DE
Tramadolor einmal täglich 300 mg Retardtabletten	FR/H/0272/003	63149.00.00	HEXAL AG	DE
Tramadolor einmal täglich 300 mg Retardtabletten	FR/H/0272/003	63149.00.00	HEXAL AG	DE
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

Product 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 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
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 retard 100 mg - Tabletten	DE/H/0288/001	1-24377	HEXAL PHARMA GMBH	AT
Tramadolor retard 100 mg tablete	/	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	/	HR-H-008744405	SANDOZ D.O.O.	HR
Tramadolor retard 200 mg - Tabletten	DE/H/0288/003	1-24383	HEXAL PHARMA GMBH	AT
Tramadolor retard 200 mg tablete	/	HR-H-026332681	SANDOZ D.O.O.	HR
Tramadolor® 100 injekt	not available	6137845.01.00	HEXAL AG	DE
Tramadolor® 100 mg Brause	not available	37201.01.00	HEXAL AG	DE
Tramadolor® 100 mg ID	DE/H/0448/001	11558.00.00	HEXAL AG	DE
Tramadolor® 100 mg/ml Lösung, Tropfen zum Einnehmen, Lösung	not available	6301608.00.00	HEXAL AG	DE
Tramadolor® 150 mg ID	DE/H/0448/002	45017.01.00	HEXAL AG	DE
Tramadolor® 200 mg ID	DE/H/0448/003	45017.02.00	HEXAL AG	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
Tramadolor® 50 injekt	not available	35345.00.00	HEXAL AG	DE
Tramadolor® Kapseln	not available	32747.00.02	HEXAL AG	DE
Tramadolor® long 100 mg Hartkapseln, retardiert	not available	42433.01.00	HEXAL AG	DE
Tramadolor® long 150 mg Hartkapseln, retardiert	not available	42433.02.00	HEXAL AG	DE
Tramadolor® long 200 mg Hartkapseln, retardiert	not available	42433.03.00	HEXAL AG	DE
Tramadolor® long 50 mg Hartkapseln, retardiert	not available	42433.00.00	HEXAL AG	DE
Tramadolor® tabs	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
Tramadol-Sandoz 100 mg Retardtabletten	NL/H/0483/001	60258.00.00	HEXAL AG	DE
Tramadol-Sandoz 150 mg Retardtabletten	NL/H/0483/002	60258.01.00	HEXAL AG	DE
Tramadol-Sandoz 200 mg Retardtabletten	NL/H/0483/003	60258.02.00	HEXAL AG	DE
Tramadura Injekt Wirkstoff: Tramadolhydrochlorid	not available	38502.00.00	MYLAN DURA GMBH	DE
Tramag 50	not available	5892/2005/01	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 100 mg tabletten	not available	RVG 22452	MUNDIPHARMA PHARMACEUTICALS BV	NL
Tramagetic Retard 150 mg depottabletter	not available	99-7783	MUNDIPHARMA AS.	NO
TRAMAGETIC RETARD 150 mg tabletten	not available	RVG 22453	MUNDIPHARMA PHARMACEUTICALS BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramagetic Retard 200 mg depottabletter	not available	99-7784	MUNDIPHARMA AS.	NO
TRAMAGETIC RETARD 200 mg tabletten	not available	RVG 22454	MUNDIPHARMA PHARMACEUTICALS BV	NL
Tramagetic Retard 75 mg depottabletter	not available	99-7781	MUNDIPHARMA AS.	NO
TRAMAGETIC RETARD 75 mg tabletten	not available	RVG 22451	MUNDIPHARMA PHARMACEUTICALS BV	NL
Tramagit 100 mg otopina za injekcije	not available	HR-H-237941754	FARMAL DD.	HR
Tramagit 50 mg otopina za injekcije	not available	HR-H-576551176	FARMAL DD.	HR
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
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® 50 mg/ml Injektionslösung	not available	34981.00.00	KREWEL MEUSELBACH GMBH	DE
Tramagit® Tabletten 50 mg	not available	6140209.00.00	KREWEL MEUSELBACH GMBH	DE
TRAMAKE 100mg Tablets	not available	PA 1329/4/2	GALEN LIMITED	IE
TRAMAKE 100mg Tablets	not available	PA 1329/4/2	GALEN LIMITED	IE
TRAMAKE 50mg Tablets	not available	PA 1329/4/1	GALEN LIMITED	IE
TRAMAKE 50mg Tablets	not available	PA 1329/4/1	GALEN LIMITED	IE
Tramal 100 mg ilgstošās darbības tabletes	not available	99-0530	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg otopina za injekcije	not available	UP/I-530-09/09-02/91	STADA D.O.O.	HR
Tramal 100 mg supozitoriji	not available	99-0529	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg svečke	not available	H/92/01556/007	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
Tramal 100 mg tablete s produljenim oslobadanjem	not available	HR-H-61 0358597	STADA D.O.O.	HR
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/2 ml solução injectável	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	not available	UP/I-530-09/09-02/92	STADA D.O.O	HR
Tramal 100 mg/ml peroralne kapljice, raztopina v kapalnem vsebniku	not available	H/92/01556/005	STADA ARZNEIMITTEL AG	SI
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 iekšķigai lietošanai, šķidums pudele ar dozešanas sukni	not available	13-0057	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg/ml pilieni iekšķigai lietošanai, šķidums pudele ar pilinataju	not available	99-0528	STADA ARZNEIMITTEL AG	LV
Tramal 100 mg/ml tipat, liuos	not available	12688	ORION OYJ	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 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 produljenim oslobadanjem	not available	HR-H-505350700	STADA D.O.O.	HR
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 filmsko obložene tablete s podaljšanim sproščanjem	DE/H/0108/004	5363-I-637/13	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg filmsko obložene tablete s podaljšanim sproščanjem	DE/H/0108/004	5363-I-637/13	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg filmsko obložene tablete s podaljšanim sproščanjem	DE/H/0108/004	5363-I-637/13	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg filmsko obložene tablete s podaljšanim sproščanjem	DE/H/0108/004	5363-I-637/13	STADA ARZNEIMITTEL AG	SI
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 kapsule	not available	UP/I-530-09/09-02/89	STADA D.O.O	HR
Tramal 50 mg liukeneva tabletti	not available	13245	ORION OYJ	FI
Tramal 50 mg otopina za injekcije	not available	UP/I-530-09/09-02/90	STADA D.O.O	HR
Tramal 50 mg tablete s produljenim oslobadanjem	not available	HR-H-21 0396637	STADA D.O.O.	HR
Tramal 50 mg trde kapsule	not available	H/92/01556/004	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
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/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 injiciranje/infundiranje	not available	H/92/01556/009	STADA ARZNEIMITTEL AG	SI
Tramal 50 mg/ml škidums injekcijam	not available	96-0105	STADA ARZNEIMITTEL AG	LV
TRAMAL čapíky 100 mg	not available	65/0076/91-S	STADA ARZNEIMITTEL AG	SK
TRAMAL čípky 100 mg	not available	65/076/91-S/C	STADA ARZNEIMITTEL AG	CZ
TRAMAL injekční roztok 100 mg/2 ml	not available	65/079/91-S/C	STADA ARZNEIMITTEL AG	CZ
TRAMAL injekcní roztok 50 mg/1 ml	not available	65/078/91-S/C	STADA ARZNEIMITTEL AG	CZ
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 kapky 100 mg/1 ml	not available	65/077/91-S/C	STADA ARZNEIMITTEL AG	CZ
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 kapsuly 50 mg	not available	65/0075/91-S	STADA ARZNEIMITTEL AG	SK
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

Product 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 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 retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2971984	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2511186	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	5419486	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2971984	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2511186	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	5419486	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	DE/H/0108/001	2971984	GRÜNENTHAL S.A.	PT
Tramal retard 100 mg comprimidos de libertação prolongada	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
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 depottabletti	DE/H/0108/001	12497	GRÜNENTHAL GMBH	FI
Tramal retard 100 mg depottabletti	DE/H/0108/001	12497	GRÜNENTHAL GMBH	FI
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	7021/2006/01	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 100 mg, comprimate cu eliberare prelungită, 100 mg	not available	7021/2006/02	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, tabletten met verlengde afgifte	DE/H/0108/001	RVG 22361	GRÜNENTHAL B.V.	NL
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 mg Retardtabletten	DE/H/0108/001	37297.00.00	GRÜNENTHAL GMBH	DE
Tramal retard 100 mg Retardtabletten	DE/H/0108/001	37297.00.00	GRÜNENTHAL GMBH	DE
Tramal Retard 100, 100 mg, tabletki o przedluzonym uwalnianiu	not available	7862	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 150 mg comprimidos de libertação prolongada	DE/H/0108/002	5419585	GRÜNENTHAL S.A.	PT
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 comprimidos de libertação prolongada	DE/H/0108/002	5419585	GRÜNENTHAL S.A.	PT
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 comprimidos de libertação prolongada	DE/H/0108/002	5419585	GRÜNENTHAL S.A.	PT
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 depottabletti	DE/H/0108/002	12498	GRÜNENTHAL GMBH	FI
Tramal retard 150 mg depottabletti	DE/H/0108/002	12498	GRÜNENTHAL GMBH	FI
TRAMAL RETARD 150 mg toimeainet prolongeeritult vabastavad tabletid	not available	263199	STADA ARZNEIMITTEL AG	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
TRAMAL RETARD 150 mg, comprimate cu eliberare prelungită, 150 mg	not available	6394/2006/02	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 150 mg, comprimate cu eliberare prelungită, 150 mg	not available	6394/2006/01	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, tabletten met verlengde afgifte	DE/H/0108/002	RVG 22362	GRÜNENTHAL B.V.	NL
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 mg Retardtabletten	DE/H/0108/002	37297.01.00	GRÜNENTHAL GMBH	DE
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
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 comprimidos de libertação prolongada	DE/H/0108/003	5419684	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
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 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 depottabletti	DE/H/0108/003	12499	GRÜNENTHAL GMBH	FI
Tramal retard 200 mg depottabletti	DE/H/0108/003	12499	GRÜNENTHAL GMBH	FI
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	6395/2006/02	STADA ARZNEIMITTEL AG	RO
TRAMAL RETARD 200 mg, comprimate cu eliberare prelungită, 200 mg	not available	6395/2006/01	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, tabletten met verlengde afgifte	DE/H/0108/003	RVG 22363	GRÜNENTHAL 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
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 mg, Retardtabletten	DE/H/0108/003	37297.02.00	GRÜNENTHAL GMBH	DE
Tramal retard 200 mg, Retardtabletten	DE/H/0108/003	37297.02.00	GRÜNENTHAL GMBH	DE
Tramal Retard 200, 200 mg, tabletki o przedłużonym uwalnianiu	not available	7864	STADA ARZNEIMITTEL AG	PL
Tramal retard 50 mg depottabletti	DE/H/0108/004	22364	GRÜNENTHAL GMBH	FI
Tramal retard 50 mg depottabletti	DE/H/0108/004	22364	GRÜNENTHAL GMBH	FI
Tramal retard 50 mg depottabletti	DE/H/0108/004	22364	GRÜNENTHAL GMBH	FI
Tramal retard 50 mg Retardtabletten	DE/H/0108/004	60442.00.00	GRÜNENTHAL GMBH	DE
Tramal retard 50 mg Retardtabletten	DE/H/0108/004	60442.00.00	GRÜNENTHAL GMBH	DE
Tramal retard 50 mg Retardtabletten	DE/H/0108/004	60442.00.00	GRÜNENTHAL GMBH	DE
Tramal Retard 50, 50 mg, tabletki o przedluzonym uwalnianiu	DE/H/0108/004	16764	STADA ARZNEIMITTEL AG	PL
Tramal Retard 50, 50 mg, tabletki o przedluzonym uwalnianiu	DE/H/0108/004	16764	STADA ARZNEIMITTEL AG	PL
Tramal Retard 50, 50 mg, tabletki o przedluzonym uwalnianiu	DE/H/0108/004	16764	STADA ARZNEIMITTEL AG	PL
Tramal Retard 50, 50 mg, tabletki o przedluzonym uwalnianiu	DE/H/0108/004	16764	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 tablety 100 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL RETARD TABLETY 100 mg tablety s prodlouženým uvolňováním	not available	65/873/97-C	STADA ARZNEIMITTEL AG	CZ
TRAMAL retard tablety 150 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL RETARD TABLETY 150 mg tablety s prodlouženým uvolňováním	not available	65/539/99-C	STADA ARZNEIMITTEL AG	CZ
TRAMAL retard tablety 200 mg	not available	65/0359/97-S	STADA ARZNEIMITTEL AG	SK
TRAMAL RETARD TABLETY 200 mg tablety s prodlouženým uvolňováním	not available	65/540/99-C	STADA ARZNEIMITTEL AG	CZ
Tramal Tabletten, 50 mg	not available	28705.00.00	GRÜNENTHAL GMBH	DE
TRAMAL tobolky 50 mg Tvrdé tobolky	not available	65/075/91-S/C	STADA ARZNEIMITTEL AG	CZ
Tramal Tropfen, 100 mg/ml, Lösung zum Einnehmen	not available	120/88/06/0248	GRÜNENTHAL GMBH	LU
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
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

Product 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, druppels voor oraal gebruik, oplossing 100 mg/ml	not available	RVG 15513	GRÜNENTHAL B.V.	NL
Tramal® 100 mg Ampullen	not available	17.690	GRÜNENTHAL GES. M.B.H.	AT
Tramal® 100 mg filmsko obložene tablete s podaljšanim sproščanjem	not available	H/92/01556/001	STADA ARZNEIMITTEL AG	SI
Tramal® 150 mg filmsko obložene tablete s podaljšanim sproščanjem	not available	H/92/01556/002	STADA ARZNEIMITTEL AG	SI
Tramal® 200 mg filmsko obložene tablete s podaljšanim sproščanjem	not available	H/92/01556/003	STADA ARZNEIMITTEL AG	SI
Tramal® 50 mg Ampullen	not available	17692	GRÜNENTHAL GES. M.B.H.	AT
Tramal® 50 mg Kapseln	not available	17.688	GRÜNENTHAL GES. M.B.H.	AT
Tramal® retard 100 mg Filmtabletten	not available	1-21219	GRÜNENTHAL GES. M.B.H.	AT
Tramal® retard 150 mg Filmtabletten	not available	1-21218	GRÜNENTHAL GES. M.B.H.	AT
Tramal® retard 200 mg Filmtabletten	not available	1-21217	GRÜNENTHAL GES. M.B.H.	AT
Tramal® Tropfen	not available	17.689	GRÜNENTHAL GES. M.B.H.	AT
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
Tramalgin, 50 mg/mL, roztwór do wstrzykiwań	not available	21762	SOPHARMA WARSZAWA SP. Z O.O.	PL
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

Product 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	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
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

Product 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 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 compresse a rilascio prolungato	AT/H/0118/003	035846171	S.F. GROUP SRL	IT
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

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramapine 50 mg Capsules	not available	PA0281/100/1	PINEWOOD LABORATORIES LIMITED	IE
Tramapine 50 mg Capsules	not available	PA0281/100/1	PINEWOOD LABORATORIES LIMITED	IE
Tramastad 100 mg/2 ml Ampullen	not available	1-23281	STADA ARZNEIMITTEL GMBH	AT
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 100 mg/ml Tropfen	not available	1-23282	STADA ARZNEIMITTEL GMBH	AT
Tramastad 50 mg Kapseln	not available	1-23280	STADA ARZNEIMITTEL GMBH	AT
Tramastad 50 mg Kapseln	not available	1-23280	STADA ARZNEIMITTEL GMBH	AT
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 516-0 OU 34009 366 516 0 9	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 517-7 OU 34009 366 517 7 7	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 518-3 OU 34009 366 518 3 8	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 520-8 OU 34009 366 520 8 8	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 006-6 OU 34009 566 006 6 3	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 007-2 OU 34009 566 007 2 4	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 008-9 OU 34009 566 008 9 2	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 009-5 OU 34009 566 009 5 3	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 010-3 OU 34009 566 010 3 5	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 516-0 OU 34009 366 516 0 9	ETHYPHARM	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
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 517-7 OU 34009 366 517 7 7	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 518-3 OU 34009 366 518 3 8	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	366 520-8 OU 34009 366 520 8 8	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 006-6 OU 34009 566 006 6 3	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 007-2 OU 34009 566 007 2 4	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 008-9 OU 34009 566 008 9 2	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 009-5 OU 34009 566 009 5 3	ETHYPHARM	FR
TRAMELENE 50 mg, comprimé orodispersible	UK/H/0641/001	566 010-3 OU 34009 566 010 3 5	ETHYPHARM	FR
TRAMELENE FLASHTAB 50 mg orodispersible tablet	UK/H/0641/001	PL 06934/0071	ETHYPHARM	UK
TRAMELENE FLASHTAB 50 mg orodispersible tablet	UK/H/0641/001	PL 06934/0071	ETHYPHARM	UK
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 100 mg capsules met verlengde afgifte, hard	FI/H/0164/001	BE249103	LABORATOIRES SMB S.A.	BE
TRAMIUM 100 mg depotkapseli, kova	FI/H/0164/001	16710	LABORATOIRES SMB S.A.	FI
TRAMIUM 100 mg Gélule à libération prolongée	FI/H/0164/001	BE249103	LABORATOIRES SMB S.A.	BE
TRAMIUM 100 mg Gélule à libération prolongée	FI/H/0164/001	2003070040	LABORATOIRES SMB S.A.	LU
Tramium 150 mg ,Hartkapseln , retardiert	FI/H/0164/002	2003070039	LABORATOIRES SMB S.A.	LU
TRAMIUM 150 mg capsules met verlengde afgifte, hard	FI/H/0164/002	BE249112	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
TRAMIUM 150 mg depotkapseli, kova	FI/H/0164/002	16711	LABORATOIRES SMB S.A.	FI
TRAMIUM 150 mg Gélule à libération prolongée	FI/H/0164/002	BE249112	LABORATOIRES SMB S.A.	BE
TRAMIUM 150 mg Gélule à libération prolongée	FI/H/0164/002	2003070039	LABORATOIRES SMB S.A.	LU
Tramium 150 mg, Hartkapseln, retardiert	FI/H/0164/002	BE249112	LABORATOIRES SMB S.A.	BE
Tramium 200 mg ,Hartkapseln , retardiert	FI/H/0164/003	2003070041	LABORATOIRES SMB S.A.	LU
TRAMIUM 200 mg capsules met verlengde afgifte, hard	FI/H/0164/003	BE249121	LABORATOIRES SMB S.A.	BE
TRAMIUM 200 mg depotkapseli, kova	FI/H/0164/003	16712	LABORATOIRES SMB S.A.	FI
TRAMIUM 200 mg Gélule à libération prolongée	FI/H/0164/003	BE249121	LABORATOIRES SMB S.A.	BE
TRAMIUM 200 mg Gélule à libération prolongée	FI/H/0164/003	2003070041	LABORATOIRES SMB S.A.	LU
Tramium 200 mg, Hartkapseln, retardiert	FI/H/0164/003	BE249121	LABORATOIRES SMB S.A.	BE
Tramól-L 100 mg forðatöflur	AT/H/0117/001	IS/1/01/037/01	ACTAVIS HF.	IS
Tramól-L 150 mg forðatöflur	AT/H/0117/002	IS/1/01/037/02	ACTAVIS HF.	IS
Tramól-L 200 mg forðatöflur	AT/H/0117/003	IS/1/01/037/03	ACTAVIS HF.	IS
Tramquel SR 100 mg prolonged-release hard capsules	UK/H/0301/002	PL 15142/0125	BEECHMERE PHARMACEUTICALS LTD	UK
Tramquel SR 150 mg prolonged-release hard capsules	UK/H/0301/003	PL 15142/0126	BEECHMERE PHARMACEUTICALS LTD	UK
Tramquel SR 200 mg prolonged-release hard capsules	UK/H/0301/004	PL 15142/0127	BEECHMERE PHARMACEUTICALS LTD	UK
Tramquel SR 50 mg prolonged-release hard capsules	UK/H/0301/001	PL 15142/0124	BEECHMERE PHARMACEUTICALS LTD	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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	AT/H/0120/001	1-22188	MUNDIPHARMA GES.M.B.H	AT
Tramundal retard 200 mg Filmtabletten	AT/H/0120/002	1-22189	MUNDIPHARMA GES.M.B.H	AT
Tramundal Tropfen	not available	1-22186	MUNDIPHARMA GES.M.B.H	AT
Tramundin 100 mg tablete s podaljšanim sproščanjem	not available	H/00/01557/001	MEDIS, D.O.O.	SI
Tramundin retard 100 mg Retardtabletten	not available	34105.00.00	MUNDIPHARMA GMBH	DE
Tramundin retard 100 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-02/14	MEDIS ADRIA D.O.O.	HR
Tramundin Retard 100mg tablety s prodlouženým uvolňováním	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
Tramundin retard 200 mg Retardtabletten	AT/H/0120/002	34105.02.00	MUNDIPHARMA GMBH	DE
Tramundin, 100 mg, tabletki powlekane o przedluZonym uwalnianiu	not available	9474	NORPHARMA A/S	PL
Tramundin® 50 mg Filmtabletten	not available	6391213.00.00	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
TRAM-U-RON OD 100 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/001	4392882	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/001	4392981	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/001	4393088	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 100 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/001	4393187	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/002	4393286	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/002	4393385	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/002	4393484	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 150 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/002	4393583	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/003	4393682	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/003	4393781	BENE FARMACÊUTICA, LDA.	PT
TRAM-U-RON OD 200 mg cテ。psulas de libertaテァテ」o prolongada	FI/H/0164/003	4393880	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 200 mg C テ。psulas de liberta テアテ」o prolongada	FI/H/0164/003	4393989	BENE FARMACÊUTICA, LDA.	PT
TRAVEX DIREKT 50 mg	UK/H/0640/001	42429.00.01	MEDA PHARMA GMBH & CO. KG	DE
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3381688	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3381787	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3381886	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3382082	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3381985	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3768587	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3382181	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3382280	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3382389	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/001	3382488	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3382587	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3382686	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3382785	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3382884	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3382983	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383189	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383080	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3827086	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383288	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383387	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383486	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383585	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383783	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 150 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/001	3383882	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3383981	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384088	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384187	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384286	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384385	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3768686	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384484	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3678182	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384682	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384880	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384781	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3384989	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385085	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385184	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385283	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385481	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385382	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3827185	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385689	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385580	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385887	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385986	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3385788	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 200 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/002	3386083	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386281	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386182	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386984	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3387081	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386687	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386786	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386588	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3387289	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3387180	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3386885	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3386489	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3768785	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3386380	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387487	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3388287	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387586	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387685	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387388	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387784	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3827284	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387883	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/003	3387982	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3388089	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 300 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/003	3388188	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388485	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388683	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388386	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388584	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388782	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388881	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3388980	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3768884	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3389285	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テァテ」o prolongada	UK/H/0306/004	3389483	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389384	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389582	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389681	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389780	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3390085	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3827383	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389988	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3389889	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3390184	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3390283	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3390382	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta テアテ」o prolongada	UK/H/0306/004	3390481	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX Long 400 mg Comprimido de liberta ｱｱ｣o prolongada	UK/H/0306/004	3389087	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX Long 400 mg Comprimido de liberta ｱｱ｣o prolongada	UK/H/0306/004	3389186	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Travex Rapid 50 mg comprimidos Orodispersíveis	UK/H/0640/001	5264288	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Travex retard 100 mg Hartkapseln, retardiert	UK/H/0225/002	42429.01.00	MEDA PHARMA GMBH & CO. KG	DE
Travex retard 150 mg Hartkapseln, retardiert	UK/H/0225/003	42429.02.00	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
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	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376588	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376687	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376786	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376885	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 100 mg Cápsula dura de libertação prolongada	UK/H/0225/002	4376984	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	2632883	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377180	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377081	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377289	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377388	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 150 mg Cápsula dura de libertação prolongada	UK/H/0225/003	4377487	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377586	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	2632982	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377685	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377784	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377883	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 200 mg Cápsula dura de libertação prolongada	UK/H/0225/004	4377982	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376083	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	2632685	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376182	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376281	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376489	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
TRAVEX® 50 mg Cápsula dura de libertação prolongada	UK/H/0225/001	4376380	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	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
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	34672/24-05-12	MEDOCHEMIE HELLAS SA	GR
TRAMAL® καψάκιο σκληρό 50 mg	not available	97924	VIANEX S.A.	GR
TRAMAL® πόσιμες σταγόνες, διάλυμα 100 mg/ml	not available	30122	VIANEX S.A.	GR
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003074/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003086/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003098/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003047/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003100/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003353/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003050/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003112/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003062/M	ENDO VENTURES LIMITED	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
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003124/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003252/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003011/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003023/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003035/M	ENDO VENTURES LIMITED	IT
UNITRAMA 100 MG COMPRESSE A RILASCIO PROLUNGATO	FR/H/0272/001	037003340/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003136/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003199/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003148/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	37003201/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003151/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003213/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003365/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003163/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003225/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003377/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003175/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003237/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003187/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003249/M	ENDO VENTURES LIMITED	IT
UNITRAMA 200 MG	FR/H/0272/002	037003264/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003290/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003302/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003314/M	ENDO VENTURES LIMITED	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
UNITRAMA 300 MG	FR/H/0272/003	037003389/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003326/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003391/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003338/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003276/M	ENDO VENTURES LIMITED	IT
UNITRAMA 300 MG	FR/H/0272/003	037003288/M	ENDO VENTURES LIMITED	IT
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
Xymel SR 100 mg prolonged release tablets	not available	PA 126/99/2	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 15142/0119	MEDA PHARMACEUTICALS LTD	UK
Zamadol Melt 50 mg Tablets	not available	PL 15142/0128	MEDA PHARMACEUTICALS LTD	UK
Zamadol SR 100 mg prolonged-release hard capsules	UK/H/0225/002	PL 15142/0121	MEDA PHARMACEUTICALS LTD	UK
Zamadol SR 150 mg prolonged-release hard capsules	UK/H/0225/003	PL 15142/0122	MEDA PHARMACEUTICALS LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zamadol SR 200 mg prolonged-release hard capsules	UK/H/0225/004	PL 15142/0123	MEDA PHARMACEUTICALS LTD	UK
Zamadol SR 50 mg prolonged-release hard capsules	UK/H/0225/001	PL 15142/0120	MEDA PHARMACEUTICALS LTD	UK
Zamadol® Injection	not available	PL 15142/0118	MEDA PHARMACEUTICALS LTD	UK
ZAMUDOL LP 100 mg, gélule à libération prolongée	UK/H/0225/002	346 544.9	MEDA PHARMA SAS	FR
ZAMUDOL LP 100 mg, gélule à libération prolongée	UK/H/0225/002	346 543.2	MEDA PHARMA SAS	FR
ZAMUDOL LP 150 mg, gélule à libération prolongée	UK/H/0225/003	346 546.1	MEDA PHARMA SAS	FR
ZAMUDOL LP 150 mg, gélule à libération prolongée	UK/H/0225/003	346 545.5	MEDA PHARMA SAS	FR
ZAMUDOL LP 200 mg, gélule à libération prolongée	UK/H/0225/004	346 548.4	MEDA PHARMA SAS	FR
ZAMUDOL LP 200 mg, gélule à libération prolongée	UK/H/0225/004	346 547.8	MEDA PHARMA SAS	FR
ZAMUDOL LP 50 mg, gélule à libération prolongée	UK/H/0225/001	346 542.6	MEDA PHARMA SAS	FR
ZAMUDOL LP 50 mg, gélule à libération prolongée	UK/H/0225/001	346 540.3	MEDA PHARMA SAS	FR
Zeridame SR 100mg Prolonged Release Tablets	NL/H/0890/001	PL 30306/0722	ACTAVIS GROUP PTC EHF.	UK
Zeridame SR 150mg Prolonged Release Tablets	NL/H/0890/002	PL 30306/0723	ACTAVIS GROUP PTC EHF.	UK
Zeridame SR 200mg Prolonged Release Tablets	NL/H/0890/003	PL 30306/0724	ACTAVIS GROUP PTC EHF.	UK
ZYDOL 100 mg / 2 ml Solution for Injection	not available	PA 1189/1/4	GRÜNENTHAL LTD.	IE
ZYDOL 50 mg Hard Capsules	not available	PA 1189/1/1	GRÜNENTHAL LTD.	IE
ZYDOL 50mg Capsules	not available	PL 21727/0001	GRÜNENTHAL LTD.	UK
ZYDOL 50mg Soluble Tablets	not available	PL 21727/0006	GRÜNENTHAL LTD.	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYDOL SR 100 mg prolonged-release tablets	not available	PA 1189/1/6	GRÜNENTHAL 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 1189/1/7	GRÜNENTHAL LTD.	IE
ZYDOL SR 150 mg prolonged-release tablets	not available	PL 21727/0004	GRÜNENTHAL LTD.	UK
ZYDOL SR 200 mg prolonged-release tablets	not available	PA 1189/1/8	GRÜNENTHAL LTD.	IE
ZYDOL SR 200 mg prolonged-release tablets	not available	PL 21727/0005	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
ZYDOL® 50 mg/ml Solution for Injection	not available	PL 21727/0002	GRÜNENTHAL LTD.	UK
ZYTRAM	not available	970500 (IS)	NORPHARMA A/S	IS
ZYTRAM	not available	970499 (IS)	NORPHARMA A/S	IS
ZYTRAM	not available	970501 (IS)	NORPHARMA A/S	IS
ZYTRAM	not available	970498 (IS)	NORPHARMA A/S	IS
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/00306/002	63.131	MUNDIPHARMA PHARMACEUTICALS SL	ES
Zytram 300 mg comprimidos de liberación prolongada	UK/H/0306/003	63.132	MUNDIPHARMA PHARMACEUTICALS SL	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
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
ZYTRAM SR 100 mg prolonged release tablets	UK/H/0330/002	PL 16950/0102	NAPP PHARMACEUTICALS LTD	UK
ZYTRAM SR 150 mg prolonged release tablets	UK/H/0330/003	PL 16950/0103	NAPP PHARMACEUTICALS LTD	UK
ZYTRAM SR 200 mg prolonged release tablets	UK/H/0330/004	PL 16950/0104	NAPP PHARMACEUTICALS LTD	UK
ZYTRAM SR 75 mg prolonged release tablets	UK/H/0330/001	PL 16950/0101	NAPP PHARMACEUTICALS LTD	UK
ZYTRAM UNO 150 mg forðatöflur	not available	970442 (IS)	NORPHARMA A/S	IS
ZYTRAM UNO forðatöflur 200 mg	not available	970443	NORPHARMA A/S	IS
ZYTRAM UNO forðatöflur 300 mg	not available	970444	NORPHARMA A/S	IS
ZYTRAM UNO forðatöflur 400 mg	not available	970445	NORPHARMA A/S	IS
ΣRAMAL® Δλέζηκν δηάιπκα 100 mg/2ml AMP	not available	61457	VIANEX S.A.	GR
TRAMAL® SR 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	22043/15/06-03-2017	VIANEX S.A.	GR
TRAMAL® SR 150 mg δισκία παρατεταμένης αποδέσμευσης	not available	22044/15/06-03-2017	VIANEX S.A.	GR
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

Product 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® Υπόθετα 100 mg	not available	54399/12	VIANEX S.A.	GR
Маброн 50 mg капсули	not available	20030082	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 STADA 50 mg капсула, твърда	not available	9600290	STADA ARZNEIMITTEL AG	BG
Трамадол STADA 50 mg капсула, твърда	not available	9600290	STADA ARZNEIMITTEL AG	BG
Трамадол STADA 50 mg/ml инжекционен разтвор	not available	9600289	STADA ARZNEIMITTEL AG	BG
Трамадол STADA 50 mg/ml инжекционен разтвор	not available	9600289	STADA ARZNEIMITTEL AG	BG
Трамалгин 50 mg твърди капсули	not available	20030660	UNIPHARM	BG
Трамалгин 50 mg/ml инжекционен разтвор	not available	20050203	SOPHARMA AD	BG