21 March 2024
EMA/178796/2024
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: paracetamol/tramadol

Procedure no.: PSUSA/00002310/202308

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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
Algotra 75 mg/650 mg comprimés enrobés	BE/H/0299/003	BE660419	LABORATOIRES SMB S.A.	BE
Algotra 75 mg/650 mg comprimés enrobés	BE/H/0299/003	2022090206	LABORATOIRES SMB S.A.	LU
Algotra 75 mg/650 mg omhulde tabletten	BE/H/0299/003	BE660419	LABORATOIRES SMB S.A.	BE
Algotra 75 mg/650 mg Überzogene Tabletten	BE/H/0299/003	BE660419	LABORATOIRES SMB S.A.	BE
Algotra 75 mg/650 mg Überzogene Tabletten	BE/H/0299/003	2022090206	LABORATOIRES SMB S.A.	LU
Captor 37,5 mg/325 mgcomprimidos EFG	not available	75629	FERRER INTERNACIONAL, S.A.	ES
Captor 37,5 mg/325 mgcomprimidos EFG	not available	75629	FERRER INTERNACIONAL, S.A.	ES
Captor 75 mg/650 mg comprimidos	not available	75630	FERRER INTERNACIONAL, S.A.	ES
Captor 75 mg/650 mg comprimidos	not available	75630	FERRER INTERNACIONAL, S.A.	ES
Clanderon 75 mg / 650 mg comprimidos recubiertos con pelicula	PT/H/1315/002	81005	ARISTO PHARMA IBERIA, S.L.	ES
Clanderon 75 mg/650 mg comprimidos efervescentes	PT/H/1210/002	80416	ARISTO PHARMA IBERIA, S.L.	ES
Diliban 75 mg/650 mg comprimidos efervescentes	not available	84811	LABORATORIOS GEBRO PHARMA, S.A.	ES
Diliban Retard 75 mg/650 mg comprimidos de liberación prolongada	ES/H/0613/001	73638	LABORATORIOS GEBRO PHARMA, S.A.	ES
Diliban75 mg/650 mg comprimidos	not available	75633	LABORATORIOS GEBRO PHARMA, S.A.	ES
Doreta 75 mg/650 mg apvalkotās tabletes	HU/H/0190/002	12-0062	KRKA, D.D., NOVO MESTO	LV
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/11	KRKA, D.D., NOVO MESTO	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
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/12	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/13	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/14	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/15	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/16	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/17	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/18	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/19	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/20	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/01	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/02	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/03	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/04	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/05	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/06	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/07	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/08	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/09	KRKA, D.D., NOVO MESTO	RO

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Doreta 75 mg/650 mg comprimate filmate	HU/H/0190/002	6992/2014/10	KRKA, D.D., NOVO MESTO	RO
Doreta 75 mg/650 mg filmom obalené tablety	HU/H/0190/002	65/0742/11-S	KRKA, D.D., NOVO MESTO	SK
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/012	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/013	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/014	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/015	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/016	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/017	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/018	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/019	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/020	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/021	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/022	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/036	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/038	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/043	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/044	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/034	KRKA, D.D., NOVO MESTO	SI

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Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/035	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/037	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/039	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/040	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/041	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmsko obložene tablete	HU/H/0190/002	H/09/00506/042	KRKA, D.D., NOVO MESTO	SI
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/12	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/13	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/14	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/15	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/16	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/17	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/18	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/19	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/20	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/21	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/33	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/34	KRKA, D.D., NOVO MESTO	HU

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Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/35	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/36	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/37	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/38	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/39	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/40	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/41	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg filmtabletta	HU/H/0190/002	OGYI-T-21059/42	KRKA, D.D., NOVO MESTO	HU
Doreta 75 mg/650 mg õhukese polümeerikattega tabletid	HU/H/0190/002	772412	KRKA, D.D., NOVO MESTO	EE
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/001	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/002	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/003	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/004	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/005	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg pailginto atpalaidavimo	HU/H/0190/003	LT/1/23/5225/006	KRKA, D.D., NOVO MESTO	LT

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tabletės				
Doreta 75 mg/650 mg pailginto atpalaidavimo tabletės	HU/H/0190/003	LT/1/23/5225/007	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/012	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/013	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/014	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/015	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/016	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/017	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/018	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/019	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/020	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg plėvele dengtos tabletės	HU/H/0190/002	LT/1/09/1638/021	KRKA, D.D., NOVO MESTO	LT
Doreta 75 mg/650 mg potahované tablety	HU/H/0190/002	65/185/12-C	KRKA, D.D., NOVO MESTO	CZ
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/01	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/02	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/03	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg	HU/H/0190/003	14092/2021/04	KRKA, D.D., NOVO MESTO	RO

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comprimate cu eliberare prelungită				
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/05	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/06	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/07	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/08	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/09	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/10	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/11	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/12	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/13	KRKA, D.D., NOVO MESTO	RO
Doreta EP 75 mg/650 mg comprimate cu eliberare prelungită	HU/H/0190/003	14092/2021/14	KRKA, D.D., NOVO MESTO	RO
Doreta Prolong 75 mg/650 mg tablety s prodlouženým uvolňováním	HU/H/0190/003	65/575/15-C	KRKA, D.D., NOVO MESTO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/43	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/44	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/45	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/46	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/47	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/48	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/49	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/50	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/51	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg retard tabletta	HU/H/0190/003	OGYI-T-21059/52	KRKA, D.D., NOVO MESTO	HU
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/045	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/046	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/047	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/048	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/049	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg	HU/H/0190/003	H/09/00506/050	KRKA, D.D., NOVO MESTO	SI

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tablete s podaljšanim sproščanjem				
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/051	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/052	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/053	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/054	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/055	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/056	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/057	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablete s podaljšanim sproščanjem	HU/H/0190/003	H/09/00506/058	KRKA, D.D., NOVO MESTO	SI
Doreta SR 75 mg/650 mg tablety s predĺženým uvoľňovaním	HU/H/0190/003	65/0452/15-S	KRKA, D.D., NOVO MESTO	SK
Doreta SR, 75 mg + 650 mg, tabletki o przedłużonym uwalnianiu	HU/H/0190/003	22895	KRKA, D.D., NOVO MESTO	PL
Doreta, 75 mg + 650 mg, tabletki powlekane	HU/H/0190/002	19609	KRKA, D.D., NOVO MESTO	PL
EBYNDO 75 mg/650 mg compresse rivestite con	PT/H/2482/002	049599071	ITALFARMACO S.P.A.	IT

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film				
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599083	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599095	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599107	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599071	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599083	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599095	ITALFARMACO S.P.A.	IT
EBYNDO 75 mg/650 mg compresse rivestite con film	PT/H/2482/002	049599107	ITALFARMACO S.P.A.	IT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822135	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822143	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822168	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822150	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos	PT/H/2482/002	5822135	ITALFARMACO S.P.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
por película				
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822143	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822168	ITALFARMACO S.P.A	PT
Ebyndo 75 mg/650 mg comprimidos revestidos por película	PT/H/2482/002	5822150	ITALFARMACO S.P.A	PT
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 809 7 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 391 900 5 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 802 2 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 805 1 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 815 7 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 391 912 3 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 816 3 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 818 6 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 808 0 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 812 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 806 8 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 391 911 7 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 813 4 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 391 898 0 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 804 5 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 814 0 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 391 899 7 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 810 5 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 803 9 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 811 1 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé effervescent	FR/H/0211/002	34009 574 807 4 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 358 573 9 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 605 6 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 599 6 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 606 2 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 615 1 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37,5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 359 228 3 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37.5 mg/325 mg film coated-tablets	FR/H/0212/001	PA 2242/6/1	GRÜNENTHAL PHARMA LTD.	IE
IXPRIM 37.5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 618 0 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37.5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 616 8 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37.5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 614 5 8	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
IXPRIM 37.5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 563 617 4 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
IXPRIM 37.5 mg/325 mg, comprimé pelliculé	FR/H/0211/001	34009 359 230 8 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Ixprim effervescent 37.5 mg/325 mg effervescent tablets	FR/H/0212/002	PA 2242/6/2	GRÜNENTHAL PHARMA LTD.	IE
KOLIBRI 37,5 mg / 325 mg, compresse effervescenti	not available	036993071	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse effervescenti	not available	036993069	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse effervescenti	not available	036993095	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse effervescenti	not available	036993083	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse rivestite con film	not available	036993044	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse rivestite con film	not available	036993020	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse rivestite con film	not available	036993032	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse rivestite con film	not available	036993018	ALFASIGMA S.P.A.	IT
KOLIBRI 37,5 mg / 325 mg, compresse rivestite con film	not available	036993057	ALFASIGMA S.P.A.	IT
Palgotal 75 mg/650 mg potahované tablety	CZ/H/0688/001	65/252/14-C	ZENTIVA, K.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Palgotal, 75 mg + 650 mg, tabletki powlekane	CZ/H/0688/001	22121	ZENTIVA, K.S.	PL
PATROL 37,5 mg / 325 mg, compresse effervescenti	not available	036996066	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse effervescenti	not available	036996078	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse effervescenti	not available	036996080	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse effervescenti	not available	036996092	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse rivestite con film	not available	036996041	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse rivestite con film	not available	036996027	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse rivestite con film	not available	036996039	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse rivestite con film	not available	036996015	ALFASIGMA S.P.A.	IT
PATROL 37,5 mg / 325 mg, compresse rivestite con film	not available	036996054	ALFASIGMA S.P.A.	IT
Pazital 37,5 mg/325 mg comprimidos recubiertos con película	not available	66853	GRÜNENTHAL PHARMA S.A.	ES
Poltram Combo Forte, 75 mg + 650 mg, tabletki powlekane	not available	23297	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Pontalsic 37,5 mg/325 mg	FR/H/0211/001	65.158	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
comprimidos recubiertos con película				
Tramabian® 75 mg/650 mg Filmtabletten	HU/H/0190/002	93508.00.00	TAD PHARMA GMBH	DE
TRAMACET 37.5 mg/325 mg effervescent tablets	FR/H/0211/002	PL 50414/0025	GRÜNENTHAL PHARMA LTD.	XI
TRAMACET 37.5mg/325mg Film coated tablets	FR/H/0211/001	PL 50414/0024	GRÜNENTHAL PHARMA LTD.	XI
Tramadol + Paracetamol Aristo 75 mg + 650 mg comprimidos efervescentes	PT/H/1210/002	5669809	ARISTO PHARMA IBERIA, S.L.	PT
Tramadol + Paracetamol Aristo 75 mg + 650 mg comprimidos efervescentes	PT/H/1210/002	5669775	ARISTO PHARMA IBERIA, S.L.	PT
Tramadol + Paracetamol Aristo 75 mg + 650 mg comprimidos revestidos por película	PT/H/1315/002	5659222	ARISTO PHARMA IBERIA, S.L.	PT
Tramadol + Paracetamol Aristo 75 mg + 650 mg comprimidos revestidos por película	PT/H/1315/002	5659214	ARISTO PHARMA IBERIA, S.L.	PT
Tramadol + Paracetamol Aristo 75 mg + 650 mg comprimidos revestidos por película	PT/H/1315/002	5659206	ARISTO PHARMA IBERIA, S.L.	PT
Tramadol + Paracetamol Bluepharma 75 mg + 650 mg comprimidos	not available	5493564	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Tramadol + Paracetamol Bluepharma 75 mg + 650 mg comprimidos	not available	5493556	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	5488655	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 + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	5488663	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/633/02	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/633/02	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/633/02	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/633/02	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 + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Generis 75 mg + 650 mg comprimidos	PT/H/0633/002	PT/H/0633/002	GENERIS FARMACÊUTICA, S.A.	PT
Tramadol + Paracetamol Helm	PT/H/2716/001	PT/H/2716/001	HELM AG	PT
Tramadol + Paracetamol Krka 75 mg + 650 mg comprimidos de libertação prolongada	HU/H/0190/003	5667019	KRKA, D.D., NOVO MESTO	PT
Tramadol + Paracetamol Krka 75 mg + 650 mg comprimidos de libertação prolongada	HU/H/0190/003	5667027	KRKA, D.D., NOVO MESTO	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 + Paracetamol Krka 75 mg + 650 mg comprimidos de libertação prolongada	HU/H/0190/003	5667035	KRKA, D.D., NOVO MESTO	PT
Tramadol + Paracetamol Krka 75 mg + 650 mg comprimidos de libertação prolongada	HU/H/0190/003	5679469	KRKA, D.D., NOVO MESTO	PT
Tramadol + Paracetamol Krka 75 mg + 650 mg comprimidos revestidos por película	not available	5678909	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Tramadol + Paracetamol Pharmakern 75 mg + 650 mg comprimidos	not available	5834973	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Tramadol + Paracetamol ratiopharm 75 mg + 650 mg comprimidos	HR/H/0232/001	5729561	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Tramadol + Paracetamol ratiopharm 75 mg + 650 mg comprimidos	HR/H/0232/001	5729553	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Tramadol + Paracetamol ratiopharm 75 mg + 650 mg comprimidos	HR/H/0232/001	5729579	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Tramadol + Paracetamol toLife 75 mg + 650 mg comprimidos	PT/H/0632/002	5488721	TOWA PHARMACEUTICAL S.A.	PT
Tramadol + Paracetamol toLife 75 mg + 650 mg comprimidos	PT/H/0632/002	5488713	TOWA PHARMACEUTICAL S.A.	PT
Tramadol/ Paracetamol Aristo 75 mg/650 mg comprimidos efervescentes.	not available	86603	ARISTO PHARMA IBERIA, S.L.	ES
Tramadol/ Paracetamol Aristo 75 mg / 650 mg	not available	86589	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
comprimidos recubiertos con película.				
Tramadol/Paracetamol 75 mg / 650 mg tablets	not available	PL35533/0043	ASPIRE PHARMA LIMITED	XI
Tramadol/Paracetamol 75 mg / 650 mg tablets	not available	PL35533/0043	ASPIRE PHARMA LIMITED	XI
Tramadol/Paracetamol 75 mg / 650 mg tablets	not available	PL35533/0043	ASPIRE PHARMA LIMITED	XI
Tramadol/Paracetamol 75 mg / 650 mg tablets	not available	PL35533/0043	ASPIRE PHARMA LIMITED	XI
Tramadol/Paracetamol Aristo 75 mg/650 mg Filmtabletten	PT/H/1315/002	93453.00.00	ARISTO PHARMA GMBH (ART 57)	DE
tramadol/paracetamol cinfa 75mg/650mg comprimidos recubiertos con película	not available	78.174	LABORATORIOS CINFA, S.A.	ES
Tramadol/Paracetamol Kern Pharma 75 mg/650 mg comprimidos	not available	88012	KERN PHARMA, S.L.	ES
Tramadol/Paracetamol Krka 75 mg/650 mg comprimidos recubiertos con película	HU/H/0190/002	76781	KRKA, D.D., NOVO MESTO	ES
Tramadol/Paracetamol Krka 75 mg/650 mg filmomhulde tabletten	HU/H/0463/002	BE511377	KRKA, D.D., NOVO MESTO	BE
Tramadol/Paracetamol Krka 75 mg/650 mg filmomhulde tabletten	HU/H/0463/002	BE511377	KRKA, D.D., NOVO MESTO	BE
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/01	KRKA, D.D., NOVO MESTO	HU
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/02	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/03	KRKA, D.D., NOVO MESTO	HU
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/01	KRKA, D.D., NOVO MESTO	HU
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/02	KRKA, D.D., NOVO MESTO	HU
Tramadol/Paracetamol Krka 75 mg/650 mg filmtabletta	HU/H/0463/002	OGYI-T-23201/03	KRKA, D.D., NOVO MESTO	HU
Tramadol/Paracetamol Krka 75 mg/650 mg Filmtabletten	HU/H/0190/002	136152	KRKA, D.D., NOVO MESTO	AT
Tramadol/paracetamol Normon 75 mg/650 mg comprimidos	ES/H/0519/001	84429	LABORATORIOS NORMON, S.A.	ES
Tramadol/Paracetamol Qualigen 75 mg / 650 mg comprimidos recubiertos con película.	not available	84579	NEURAXPHARM SPAIN, S.L.U.	ES
Tramadol/Paracetamol ratiopharm 75mg/650mg comprimidos recubiertos con película	not available	77385	TEVA PHARMA S.L.U.,	ES
Tramadol/Paracetamol Stada 75 mg/650 mg comprimidos	not available	75.632	LABORATORIO STADA, S.L.	ES
Tramadol/Paracetamol Teva 75mg/650mg comprimidos recubiertos con película	not available	77382	TEVA PHARMA S.L.U.,	ES
Tramadol/Paracetamol Zentiva 75 mg/650 mg filmom obalené tablety	CZ/H/0688/001	65/0195/14-S	ZENTIVA, K.S.	SK
Tramadolo e Paracetamolo	PT/H/1315/002	043580048	ARISTO PHARMA GMBH (ART	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
Aristo 75 mg/650 mg compresse rivestite con film			57)	
Tramadolo e Paracetamolo Aristo 75 mg/650 mg compresse rivestite con film	PT/H/1315/002	043580051	ARISTO PHARMA GMBH (ART 57)	IT
Tramadolo e Paracetamolo Aristo 75 mg/650 mg compresse rivestite con film	PT/H/1315/002	043580063	ARISTO PHARMA GMBH (ART 57)	IT
Tramadolo e Paracetamolo Aristo, 75 mg/650 mg compresse rivestite con film	PT/H/1315/002	043580087	ARISTO PHARMA GMBH (ART 57)	IT
TRAMAPAR FORTE, 75 mg + 650 mg, tabletki powlekane	not available	22834	POLFARMEX S.A.	PL
ZALDIAR	not available	65/0152/04-S	STADA ARZNEIMITTEL AG	SK
ZALDIAR 37,5 mg / 325 mg, comprimés pelliculés	FR/H/0212/001	BE 244553	SA GRÜNENTHAL N.V.	BE
ZALDIAR 37,5 mg / 325 mg, comprimés pelliculés	FR/H/0212/001	2003010017	SA GRÜNENTHAL N.V.	LU
Zaldiar 37,5 mg/325 mg apvalkotās tabletes	not available	04-0185	STADA ARZNEIMITTEL AG	LV
ZALDIAR 37,5 mg/325 mg comprimate filmate	not available	9695/2017/01	STADA ARZNEIMITTEL AG	RO
ZALDIAR 37,5 mg/325 mg comprimate filmate	not available	9695/2017/03	STADA ARZNEIMITTEL AG	RO
ZALDIAR 37,5 mg/325 mg comprimate filmate	not available	9695/2017/02	STADA ARZNEIMITTEL AG	RO
ZALDIAR 37,5 mg/325 mg comprimate filmate	not available	9695/2017/04	STADA ARZNEIMITTEL AG	RO
Zaldiar 37,5 mg/325 mg comprimidos efervescentes	FR/H/0212/002	70.584	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
Zaldiar 37,5 mg/325 mg comprimidos recubiertos con película	FR/H/0212/001	65149	GRÜNENTHAL PHARMA S.A.	ES
ZALDIAR 37,5 mg/325 mg filmom obložene tablete	not available	HR-H-252809369	STADA D.O.O.	HR
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/003	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/002	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/001	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/009	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/006	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/008	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/007	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/005	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/010	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/011	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmsko obložene tablete	FR/H/0212/001	H/05/01692/004	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg filmtabletta	FR/H/0212/001	OGYI-T-20557/01	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg filmtabletta	FR/H/0212/001	OGYI-T-20557/02	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg filmtabletta	FR/H/0212/001	OGYI-T-20557/05	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg filmtabletta	FR/H/0212/001	OGYI-T-20557/06	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg	FR/H/0212/001	OGYI-T-20557/03	STADA ARZNEIMITTEL AG	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
filmtabletta				
Zaldiar 37,5 mg/325 mg Filmtabletten	FR/H/0212/001	1-25827	GRÜNENTHAL GES. M.B.H.	AT
ZALDIAR 37,5 mg/325 mg Filmtabletten	FR/H/0212/001	55310.00.00	GRÜNENTHAL GMBH	DE
Zaldiar 37,5 mg/325 mg pezsgőtabletta	FR/H/0212/002	OGYI-T-20557/09	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg pezsgőtabletta	FR/H/0212/002	OGYI-T-20557/07	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg pezsgőtabletta	FR/H/0212/002	OGYI-T-20557/08	STADA ARZNEIMITTEL AG	HU
Zaldiar 37,5 mg/325 mg pezsgotabletta.	FR/H/0212/002	OGYI-T-20557/10	STADA ARZNEIMITTEL AG	HU
ZALDIAR 37,5 mg/325 mg potahované tablety	not available	65/237/02-C	STADA ARZNEIMITTEL AG	CZ
ZALDIAR 37,5 mg/325 mg šumeće tablete	not available	HR-H-676041058	STADA D.O.O.	HR
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/018	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/015	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/013	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/016	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/014	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/022	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/024	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/025	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/027	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg	FR/H/0212/002	H/05/01692/026	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
šumeče tablete				
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/020	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/017	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/032	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/028	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/029	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/030	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/019	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/012	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/021	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/031	STADA ARZNEIMITTEL AG	SI
Zaldiar 37,5 mg/325 mg šumeče tablete	FR/H/0212/002	H/05/01692/023	STADA ARZNEIMITTEL AG	SI
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 819 2 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 827 5 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 391 876 7 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 822 3 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325	FR/H/0212/002	34009 574 833 5 7	LABORATOIRES GRÜNENTHAL	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimé effervescent			S.A.S.	
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 832 9 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 820 0 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 835 8 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 830 6 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 831 2 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 391 874 4 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 391 888 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 829 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 825 2 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 828 1 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 834 1 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325	FR/H/0212/002	34009 391 889 1 1	LABORATOIRES GRÜNENTHAL	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimé effervescent			S.A.S.	
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 824 6 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 826 9 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 574 821 7 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimé effervescent	FR/H/0212/002	34009 391 875 0 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5359583	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5983085	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5319785	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5319884	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5359682	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, comprimidos revestidos por película	FR/H/0212/001	5359484	GRÜNENTHAL S.A.	PT
ZALDIAR 37,5 mg/325 mg, filmomhulde tabletten	FR/H/0212/001	BE 244553	SA GRÜNENTHAL N.V.	BE
ZALDIAR 37,5 mg/325 mg, filmomhulde tabletten	FR/H/0212/001	RVG 28113	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
ZALDIAR 37,5 mg/325 mg, õhukese polümeerikattega tabletid	not available	437704	STADA ARZNEIMITTEL AG	EE
ZALDIAR 37,5 mg/325 mg, επικαλυμμένα με λεπτό υμένιο δισκία	FR/H/0212/001	62964/11-07-2017	GRÜNENTHAL GMBH	GR
ZALDIAR 37.5 mg/325 mg, comprimé pelliculé	FR/H/0212/001	34009 563 603 3 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37.5 mg/325 mg, comprimé pelliculé	FR/H/0212/001	34009 563 597 3 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37.5 mg/325 mg, comprimé pelliculé	FR/H/0212/001	34009 563 602 7 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
ZALDIAR 37.5 mg/325 mg, comprimé pelliculé	FR/H/0212/001	34009 358 569 1 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Zaldiar Efe 37,5 mg/325 mg comprimidos efervescentes	FR/H/0212/002	5190517	GRÜNENTHAL S.A.	PT
ZALDIAR EFE 37,5 mg/325 mg, comprimidos efervescentes	FR/H/0212/002	5190509	GRÜNENTHAL S.A.	PT
ZALDIAR EFE 37,5 mg/325 mg, comprimidos efervescentes	FR/H/0212/002	5190525	GRÜNENTHAL S.A.	PT
Zaldiar effervescens 37,5 mg/325 mg šumivé tab	not available	65/0080/11-S	STADA ARZNEIMITTEL AG	SK
Zaldiar Effervescens 37,5 mg/325 mg šumivé tablety	not available	65/107/11-C	STADA ARZNEIMITTEL AG	CZ
Zaldiar Effervescent, (37,5 mg + 325 mg), tabletki musujące	not available	16461	STADA ARZNEIMITTEL AG	PL
Zaldiar, (37,5 mg + 325 mg), tabletki powlekane	not available	10733	STADA ARZNEIMITTEL AG	PL
Zalramol 75 mg/650 mg tabletta	not available	OGYI-T-24104/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Zalramol 75 mg/650 mg tabletta	not available	OGYI-T-24104/05	TEVA GYÓGYSZERGYÁR ZRT	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
Zalramol 75 mg/650 mg tabletta	not available	OGYI-T-24104/06	TEVA GYÓGYSZERGYÁR ZRT	HU
ZARACET 75 mg/650 mg filmom obalené tablety	not available	65/0382/18-S	BELUPO, S.R.O.	SK
ZARACET 75 mg/650 mg filmom obložene tablete	not available	HR-H-549246288	BELUPO D.D.	HR
ZARACET 75 mg/650 mg filmom obložene tablete	not available	HR-H-549246288	BELUPO D.D.	HR
ZARACET 75 mg/650 mg filmom obložene tablete	not available	HR-H-549246288	BELUPO D.D.	HR
ZARACET 75 mg/650 mg filmsko obložene tablete	not available	H/11/01696/005	BELUPO D.O.O.	SI
ZARACET 75 mg/650 mg filmsko obložene tablete	not available	H/11/01696/006	BELUPO D.O.O.	SI
ZARACET 75 mg/650 mg filmsko obložene tablete	not available	H/11/01696/007	BELUPO D.O.O.	SI
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	5456025	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	5456017	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen 75 mg / 650 mg comprimidos	PT/H/0631/002	PT/H/0631/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de libertação prolongada	not available	5680715	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	5680723	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	5680731	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zilpen LP 75 mg + 650 mg comprimidos de liberta□□□□o prolongada	not available	15/H/0134/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Zotramid 75 mg/650 mg tablete	HR/H/0232/001	HR-H-605800702	PLIVA HRVATSKA D.O.O.	HR
Дорета 75 mg/650 mg филмирани таблетки	HU/H/0190/002	20110723	KRKA, D.D., NOVO MESTO	BG