17 March 2016
EMA/245349/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: trimetazidine

Procedure no.: PSUSA/00003043/201508

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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
Adexor MR 35 mg módosított hatóanyagleadású filmtabletta		OGYI-T-9067/01	LES LABORATOIRES SERVIER (SURESNES)	HU
Angitrim		41/0395/10-S	Pro.Med.CS. PRAHA A.S.	SK
Apo-Trimetazidin 35 mg		83/202/12-C	Apotex Europe B.V.	CZ
Apo-Trimet PR		19918	Apotex Europe B.V.	PL
Apstar 35 mg	DE/H/2652/001/DC	20110184	GLENMARK PHARMACEUTICALS S.R.O.	BG
Apstar 35 mg	DE/H/2652/001/DC	3417/2011/11	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/27	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/28	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/29	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/30	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/32	GLENMARK PHARMACEUTICALS S.R.O.	RO
Apstar 35 mg comprimate cu eliberare prelungită	DE/H/2652/001	3417/2011/31	GLENMARK PHARMACEUTICALS S.R.O.	RO
APSTAR 35 mg retard tabletta	DE/H/2652/001/DC	OGYI-T-21715/01-24	Glenmark Pharmaceuticals s.r.o.	HU
Apstar 35 mg tablety s predl&#xFFFD;en&#xFFFD;m	DE/H/2652/001/DC	41/0070/11-S	GLENMARK PHARMACEUTICALS	SK

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uvolnovanm			S.R.O.	
Cyto-protectin MR		10845	Ethifarm Sp.z.o.o.	PL
Dilatan 20mg comprimate filmate		5372/2005/01	Terapia SA	RO
Dilatan MR 35mg comprimate filmate cu eliberare modificata		5845/2005/01	Terapia SA	RO
Dimesar		18422	Sandoz GmbH	PL
IDAPTAN 20 mg comprimidos recubiertos con película		56.704	DANVAL S.A.	ES
Lupamadazin 35 mg Retardtabletten	DE/H/2652/001	80108.00.00	LUPIN (EUROPE) LIMITED	DE
Lutrazine 35mg Retardtabletten	DE/H/2653/001	80109.00.00	Mylan dura GmbH	DE
METAZYDYNA		8613	PABIANICKIE ZAKŁADY FARMACEUTYCZNE POLFA S.A.	PL
Metazydyna MR	PL/H/0358/001/DC	19813	Adamed SP.Z.o.o.	PL
MEZITAN		OGYI-T-21279/01	ARAMIS PHARMA KFT	HU
Moduxin	HU/H/0310/001/MR	LT/1/11/2616/001-003	Gedeon Richter	LT
Moduxin	HU/H/0310/001/MR	11-0503	Gedeon Richter	LV
Moduxin	HU/H/0310/001/MR	4940/2004/01	Gedeon Richter Romania	RO
Moduxin MR	HU/H/0310/001/MR	20110539	Gedeon Richter	BG
Moduxin MR	HU/H/0310/001/MR	OGYI-T-20603/02-05	Gedeon Richter	HU
Moduxin MR	HU/H/0310/001/MR	7061/2014/01-04	Gedeon Richter Romania	RO
Portora 35 mg tablety s prodlouženým uvolňováním	DE/H/2652/001	83/251/11-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Preductal 20 mg comprimate filmate		2161/2009/01-02-03	LES LABORATOIRES SERVIER (SURESNES)	RO
PREDUCTAL MR		83/328/01-C	LES LABORATOIRES	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
			SERVIER (SURESNES)	
PREDUCTAL MR		8461	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	PL
PREDUCTAL MR		41/0355/01-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Preductal MR 35 mg módosított hatóanyagleadású filmtabletta		OGYI-T-8844/01	LES LABORATOIRES SERVIER (SURESNES)	HU
PREDUCTAL MR 35 mg comprimate cu eliberare modificată		670/2008/01-02-03	LES LABORATOIRES SERVIER (SURESNES)	RO
PREDUCTAL MR 35 mg filmsko obložene tablete s prirejenim sproščanjem		5363-I-2411/12	SERVIER PHARMA D.O.O	SI
PREDUCTAL MR 35 mg modifikuoto atpalaidavimo plėvele dengtos tabletės		LT/1/02/3176/001	LES LABORATOIRES SERVIER (SURESNES)	LT
PREDUCTAL MR 35 mg modifikuoto atpalaidavimo plėvele dengtos tablets		LT/1/02/3176/002	LES LABORATOIRES SERVIER (SURESNES)	LT
PREDUCTAL MR 35 mg tablete s prilagođenim oslobađanjem		UP/I-530-09/13-02/206	SERVIER PHARMA D.O.O -CROATIA	CR
Preductal MR 35mg ilgstošās darbības apvalkotās tabletes		03-0064	LES LABORATOIRES SERVIER (SURESNES)	LV
Preductal MR, 35 mg toimeainet modifitseeritult vabastavad tabletid		412803	LES LABORATOIRES SERVIER (SURESNES)	ET

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PREDUTRIM		4479/2012/01	ROMPHARM COMPANY S.R.L., ROMANIA	RO
Pretimectal 35mg modified-release tablets		II-30198/16.07.2015	Tchaikapharma High Quality Medicines Inc.	BG
Protevasc	HU/H/0310/001/MR	83/686/11-C	Gedeon Richter	CZ
Protevasc	HU/H/0310/001/MR	41/0438/11-S	Gedeon Richter	SK
Protevasc SR	HU/H/0310/001/MR	18820	Gedeon Richter Polska Sp. Z.o.o.	PL
TACIREL LM		5205588	LES LABORATOIRES SERVIER (SURESNES)	PT
TevaTrim 35 mg prolonged-release tablets	not available	20110058	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Triazyt MR		2194	Biofarm Sp.zoo	PL
Trimeductan MR		11518	PRZEDSIĘBIORSTWO FARMACEUTYCZNE LEK-AM SP. Z O.O.	PL
TRIMELUZINE	DE/H/2654/01/DC	3564/2011/01-24	SANDOZ S.R.L., ROMANIA	RO
Trimeluzine 35 mg tablete s podaljšanim sproščanjem	HU/H/0347/001	5363-I-868/12	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Trimepect 35 mg	SK/H/0164/001/DC	41/0773/11-S	Pharmaswiss Česká Republika S.R.O.	SK
Trimepect 35 mg tablete s prilagođenim oslobađanjem		UP/I-530-09/11-01/479	Pharmaswiss Česká Republika S.R.O.	CR
Trimetacor		MA No. II-16000 Reg. No. 20110715	Alvogen IPCo S.ar.l, Luxembourg	BG
Trimetaratio	not available	9077	RATIOPHARM GMBH	PL
Trimetazidin Actavis	EE/H/0134/001	83/578/10-C	Actavis Group PTC ehf.	CZ
Trimetazidin Mylan 35mg tablety s	DE/H/2653/001	83/199/11-C	Generics [UK] Ltd	CZ

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prodlouženým uvolňováním				
Trimetazidin Mylan 35mg Tablety s predĺženým voľňovaním	DE/H/2653/001	41/0064/11-S	Generics [UK] Ltd	SK
Trimetazidin PharmaS 35 mg tablete sa prilagođenim oslobađanjem		UP/I-530-09/11-01/10	PharmaS d.o.o.	CR
Trimetazidin Teva retard 35 mg	HU/H/0344/001-DC	83/908/10-C	Teva Pharmaceuticals CR, s.r.o.	CZ
Trimetazidin ratiopharm retard 35 mg	HU/H/0344/001-DC	41/0851/10-S	ratiopharm GmbH	SK
Trimetazidina Anox, 35mg, comprimidos de libertação prolongada.	not available	5434808	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA; ED 5A, PISO 2, PORTO SALVO; PORTUGAL	PT
Trimetazidina Anox, 35mg, comprimidos de libertação prolongada.	not available	5434774	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA; ED 5A, PISO 2, PORTO SALVO; PORTUGAL	PT
Trimetazidina Atb 35mg comprimate cu eliberare prelungita		3763/2011/01	Antibiotice SA	RO
Trimetazidina Bluepharma 20 mg Comprimidos		4747283 4747382	Bluepharma Genericos-Comercio de Medicamentos, S.A.	PT
Trimetazidina Bluepharma LP		5297015 5297023	Bluepharma Genericos-Comercio de Medicamentos, 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
Trimetazidina Cinfa 20 mg Comprimidos revestidos por película	not available	5359385	CINFA PORTUGAL, LDA.	PT
Trimetazidina Cinfa 20mg comprimidos recubiertos con pelicula EFG		68.228	Laboratorios Cinfa S.A.	ES
TRIMETAZIDINA FARMOZ 35 MG COMPRIMIDOS DE LIBERTACAO PROLONGADA (MG)		II/H/0094/001	Farmoz, Sociedade Tecnico-Medicinal, S.A.	PT
Trimetazidina Generis 35 mg Comprimidos de libertação prolongada	not available	5170279	GENERIS FARMACÊUTICA, S.A.	PT
Trimetazidina Labesfal 20mg comprimidos revestidos		4746798 4746897	Labesfal Genericos S.A.	PT
Trimetazidina Labesfal 35mg comprimido de liberatacao prolongada		5261953 5261946	Labesfal Genericos S.A.	PT
Trimetazidina Mepha 20 mg Comprimidos Revestidos	not available	4747697	MEPHA – INVESTIGAÇÃO, DESENVOLVIMENTO E FABRICAÇÃO FARMACÊUTICA, LDA. PORTO SALVO	PT
Trimetazidina Mepha 20 mg Comprimidos Revestidos	not available	4747796	MEPHA – INVESTIGAÇÃO, DESENVOLVIMENTO E FABRICAÇÃO FARMACÊUTICA, LDA. PORTO SALVO	PT
Trimetazidina Mylan 35mg Comprimido de libertação prolongada	DE/H/2653/001	5383807, 5383773, 5383823, 5383815	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
Trimetazidina Mylan 35mg comprimate cu eliberare prelungită	DE/H/2653/001	3418/2011/01, 3418/2011/02, 3418/2011/03, 3418/2011/04, 3418/2011/05, 3418/2011/06, 3418/2011/07, 3418/2011/08, 3418/2011/09, 3418/2011/10, 3418/2011/11, 3418/2011/12, 3418/2011/13, 3418/2011/14, 3418/2011/15, 3418/2011/16, 3418/2011/17, 3418/2011/18, 3418/2011/19, 3418/2011/20, 3418/2011/21, 3418/2011/22, 3418/2011/23, 3418/2011/24	Generics [UK] Ltd	RO
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5383914	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5383922	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de	DE/H/2652/01/DC	5383906	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
libertação prolongada			SOCIEDADE UNIPESSOAL, LDA.	
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5383872	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5466453	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5466461	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5466438	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina Pharmakern 35 mg comprimidos de libertação prolongada	DE/H/2652/01/DC	5466446	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Trimetazidina ratiopharm 20mg comprimidos revestidos por pelicula	not available	5399688	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA; ED 5A, PISO 2, PORTO SALVO; PORTUGAL	PT
Trimetazidina ratiopharm 20mg comprimidos revestidos por pelicula	not available	5399589	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA;	PT

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			ED 5A, PISO 2, PORTO SALVO; PORTUGAL	
Trimetazidina Teva 20 mg Comprimidos revestidos	not available	4883187	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Trimetazidina Teva 20 mg Comprimidos revestidos	not available	4883088	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Trimetazidina Teva 35 mg comprimidos de libertação prolongada		5185269 5185517	Teva Pharma Produtos Farmacêuticos Lda	PT
Trimetazidina Davur 20 mg comprimidos recubiertos EFG		64309	Laboratorios Davur S.L	ES
Trimetazidina Generis		5170261 5170279	Generis Farmacêutica, S.A.	PT
Trimetazidina Generis 20 mg Comprimidos Revestidos		4747481 4747580	Generis Farmacêutica, S.A.	PT
Trimetazidina LPH		AA711/00201	Labormed Pharma SA, RO	MT
Trimetazidina LPH		6001/2005/01-02-03-04	Labormed Pharma SA, RO	RO
Trimetazidina LPH		5371/2005/01-02-03	Labormed Pharma SA, RO	RO
Trimetazidina Mepha 20 mg Comprimidos Revestidos		4747697 4747796	Mepha 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
Trimetazidina Pensa 20 mg comprimidos recubiertos con película EFG		68227	Pensa Pharma SA	ES
Trimetazidina ratiopharm 20 mg comprimidos recubiertos con película EFG		68578	ratiopharm Espana SA;	ES
Trimetazidina ratiopharm 35 mg Comprimidos de Libertação Prolongada		5297031 5297049	Ratiopharm-Comercio e Industria de Produtos Farmaceuticos Lda;	PT
Trimetazidina Rimafar 20 mg comprimidos recubiertos EFG		64394	Laboratorios Rimafar S.L.	ES
TRIMETAZIDINA TEVA 35 mg comprimate cu eliberare prelungita	DE/H/3522/001	5817/2013/01-03	Teva Pharmaceuticals S.R.L.	RO
TRIMETAZIDINA ZENTIVA		3577384	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TRIMETAZIDINA ZENTIVA		3925286	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TRIMETAZIDINA ZENTIVA		3577186	SANOFI - PRODUTOS FARMACEUTICOS 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
TRIMETAZIDINA ZENTIVA		3577285	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Trimetazidine Actavis	EE/H/0134/001	691010	Actavis Group PTC ehf.	ET
Trimetazidine Actavis	EE/H/0134/001	LT/01/10/2146/001	Actavis Group PTC ehf.	LT
Trimetazidine Actavis	EE/H/0134/001	10-0390	Actavis Group PTC ehf.	LV
TRIMETAZIDINE EG 35 mg, comprimé pelliculé à libération modifiée	not available	3400937997072	BIOGARAN	FR
TRIMETAZIDINE EG 35MG, comprimé pelliculé à libération modifiée		NL 33235	BIOGARAN	FR
Trimetazidine MR Servier 35 mg modifikuoto atpalaidavimo plėvele dengtos tabletės		LT/1/02/3178/001	LES LABORATOIRES SERVIER (SURESNES)	LT
Trimetazidine MR Servier 35mg ilgstošās darbības apvalkotās tabletes		03-0065	LES LABORATOIRES SERVIER (SURESNES)	LV
TRIMETAZIDINE MR SERVIER		412903	LES LABORATOIRES SERVIER (SURESNES)	ET
Trimetazidine Mylan 20 mg comprime pellicule		NL22978	Mylan S.A.S.	FR
Trimetazidine Mylan 20mg/ml solution buvable en gouttes		NL23040	Mylan S.A.S.	Fr
Trimetazidine Mylan 35mg retard tabletta	DE/H/2653/001	OGYI-T-21717/01 OGYI-T-21717/02 OGYI-T-21717/03 OGYI-T-21717/04 OGYI-T-21717/05 OGYI-T-21717/06 OGYI-T-21717/07	Generics [UK] Ltd	HU

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		OGYI-T-21717/08		
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/04	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/08	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/10	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/12	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/15	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/17	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/19	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/21	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/22	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/23	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/24	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/01	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/02	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/03	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/06	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/07	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/09	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35	HU/H/0347/001	OGYI-T-21756/11	SANDOZ HUNGÁRIA KFT	HU

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mg retard tabletta				
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/13	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/14	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/16	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/18	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/20	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Sandoz 35 mg retard tabletta	HU/H/0347/001	OGYI-T-21756/05	SANDOZ HUNGÁRIA KFT	HU
Trimetazidine Teva 35 mg pailginto atpalaidavimo tabletės	HU/H/0344/001-DC	LT/1/10/2345/001 LT/1/10/2345/002 LT/1/10/2345/003	Teva Pharma B.V.	LT
Trimetazidine Teva 35mg	HU/H/0344/001-DC	734911	Teva Pharma B.V.	ET
Trimetazidine Teva 35mg ilgstošās darbības tabletes	HU/H/0344/001-DC	10-0639	Teva Pharma B.V.	LV
TRIMETAZIDINE ZYDUS 20 mg comprimé pelliculé		34009 377 452 9 1	ZYDUS FRANCE	FR
TRIMETAZIDINE BIOGARAN 20MG, comprimé pelliculé		NL21034	BIOGARAN	FR
TRIMETAZIDINE BIOGARAN 20MG/ML, solution buvable, gouttes		NL21457	BIOGARAN	FR
TRIMETAZIDINE BIOGARAN 35MG, comprimé pelliculé à libération modifiée		33237	BIOGARAN	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TRIMETAZIDINE BIOGARAN 35MG, comprimé pelliculé à libération modifiée		NL33238	BIOGARAN	FR
TRIMETAZIDINE CRISTERS 20 mg, comprimé pelliculé		NL 19389 Presentation : 34009 360 392 8 5	Cristers	FR
TRIMETAZIDINE CRISTERS 20 mg, comprimé pelliculé		NL 19389 Presentation : 34009 341 679 3 5	Cristers	FR
TRIMETAZIDINE MYLAN 35 mg, comprimé pelliculé à libération modifiée		NL 38502- CIS 66870692	MYLAN S.A.S	FR
Trimetazidin-ratiopharm 20 mg	not available	41/0196/04-S	RATIOPHARM GMBH	SK
Trimetazidin-ratiopharm 20 mg	not available	41/0196/04-S	RATIOPHARM GMBH	SK
Trimetazidin-ratiopharm 20 mg	not available	41/0196/04-S	RATIOPHARM GMBH	SK
Trimetazidin-ratiopharm 20 mg	not available	41/0196/04-S	RATIOPHARM GMBH	SK
Trimetazidin-ratiopharm 20 mg	not available	41/0196/04-S	RATIOPHARM GMBH	SK
Trimetazidinratiopharm 35 mg retard tabletta	HU/H/0344/001-DC	OGYI-T-21552/01	ratiopharm GmbH	HU
Trimetazidinratiopharm 35 mg Retardtabletten	DE/H/3522/001	86595.00.00	ratiopharm GmbH;	DE
Trimetazidinum 123ratio	HU/H/0344/001-DC	18077	123ratio Sp. z o.o.	PL
Trimetazigen MR 35mg Таблетки с удължено освобождаване	DE/H/2653/001	201110150/09.03.2011	Generics [UK] Ltd	BG
Trioxigen 35mg Tabletki	DE/H/2653/001	18762	Generics [UK] Ltd	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
o przezdłużonym Uwalnianiu				
Vascotasin	EE/H/0134/001	20100413	Actavis Group PTC ehf.	BG
Vascotasin	EE/H/0134/001	OGYI-T-21590/01	Actavis Group PTC ehf.	HU
Vascotasin	EE/H/0134/001	17685	Actavis Group PTC ehf.	PL
Vascotasin	EE/H/0134/001	41/0587/10-S	Actavis Group PTC ehf.	SK
Vastarel		11398	LES LABORATOIRES SERVIER (SURESNES)	DA
VASTAREL		9553024	SERVIER PORTUGAL, Especialidades Farmacêuticas, Lda.	PT
VASTAREL		9553032	SERVIER PORTUGAL, Especialidades Farmacêuticas, Lda.	PT
VASTAREL		9598615	SERVIER PORTUGAL, Especialidades Farmacêuticas, Lda.	PT
VASTAREL 20 MG FILM-COATED TABLETS		PA 68/10/1	SERVIER LABORATORIES (IRELAND) LTD	IE
VASTAREL 20 mg compresse rivestite		27511029	IST. FARMACO BIOLOGICO SRL STRODER	IT
VASTAREL 20 MG, comprimé pelliculé	not available	322 0452	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 MG, comprimé pelliculé	not available	322 0475	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 MG, comprimé pelliculé	not available	322 0469	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 MG, comprimé pelliculé	not available	322 0481	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 MG, comprimé pelliculé	not available	322 0498	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 mg, filmcoated tablets		MA066/01001	LES LABORATOIRES SERVIER (SURESNES)	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VASTAREL 20 mg, επικαλυμμένο με λεπτό υμένιο δισκίο		43585/07/12-6-2008	SERVIER HELLAS	GR
VASTAREL 20 mg/ml, solution buvable, gouttes	not available	322.748 3	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20 mg/ml, πόσιμες σταγόνες, διάλυμα		43586/12.06.2008	SERVIER HELLAS	GR
VASTAREL 20MG		9990	LES LABORATOIRES SERVIER (SURESNES)	CY
VASTAREL 20MG, comprimé pelliculé		NL 11321	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 20MG, comprimé pelliculé		2007029194	LES LABORATOIRES SERVIER (SURESNES)	LU
VASTAREL 20MG/ML		9989	LES LABORATOIRES SERVIER (SURESNES)	CY
VASTAREL 20MG/ML, solution buvable, gouttes		NL 11706	LES LABORATOIRES SERVIER (SURESNES)	FR
Vastarel 35 mg - Filmtabletten mit veränderter Wirkstofffreisetzung		1-27263	SERVIER AUSTRIA GMBH	AT
VASTAREL 35MG		20069	LES LABORATOIRES SERVIER (SURESNES)	CY
VASTAREL 35MG PROLONGED-RELEASE TABLETS		PA 68/10/2	SERVIER LABORATORIES (IRELAND) LTD	IE
VASTAREL 35MG, comprimé pelliculé à libération modifiée		NL 25653	LES LABORATOIRES SERVIER (SURESNES)	FR
VASTAREL 35MG, comprimé pelliculé à libération modifiée		2007049258	LES LABORATOIRES SERVIER (SURESNES)	LU
VASTAREL LM		5012992	SERVIER PORTUGAL,	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
			Especialidades Farmacêuticas, Lda.	
VASTAREL MR 35 mg, modified release filmcoated tablets		MA066/01002	LES LABORATOIRES SERVIER (SURESNES)	MT
VASTAREL® 35 mg, δισκίο ελεγχόμενης αποδέσμευσης	not available	43588/12-6-2008	SERVIER HELLAS	GR
ZIDIN		6678/2-9-2004	FINIXFARM LTD	GR
ЕНЕРГОТРИМ 35 mg таблетки с удължено освобождаване	HU/H/0347/001	20110185	SANDOZ PHARMACEUTICALS D.D.	BG
ПРЕДУКТАЛ MR 35 mg таблетки с изменено освобождаване		20020977	LES LABORATOIRES SERVIER (SURESNES)	BG
Пректазидин MR		20110175	Medica AD	BG
Триметазидин		20080137	Sopharma AD	BG