27 October 2016
EMA/719647/2016
Human Medicines Evaluation Division

List of nationally authorised medicinal

Active substance(s): gliclazide

Procedure no.: PSUSA/00001532/201602

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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
Azildor 30 mg modified-release tablets	DK/H/2376/001	MA628/12701	ACTAVIS GROUP PTC EHF.	MT
Azildor 60 mg modified-release tablets	DK/H/2376/002	MA628/12702	ACTAVIS GROUP PTC EHF.	MT
Bilxona 30mg Modified-release Tablets	DK/H/2376/001-002	PL 30306/0577	ACTAVIS GROUP PTC EHF.	UK
Bilxona 60mg Modified-release Tablets	DK/H/2376/001-002	PL 30306/0578	ACTAVIS GROUP PTC EHF.	UK
Clazistada, 30 mg, tabletki o zmodyfikowanym uwalnianiu	DK/H/2270/001	21806	STADA ARZNEIMITTEL AG	PL
Dacadis MR30 mg Modified-release Tablets	DE/H/0893/001	PL 04569/1004	GENERICS [UK] LIMITED	UK
Diabrezide - Tabletten	UK/H/0246/001	1-23119	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	AT
DIABREZIDE 80 mg compresse	not available	031844018	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Diabrezide 80 mg tablets	UK/H/0246/001	PA 925/1/1	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IE
Diabrezide or Gliclazide 80 mg tablets	UK/H/0246/001	PL 16046/0001	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	UK
DIABREZIDE, 80 mg, tabletki	not available	R/3599	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
DIACLAZID 30 mg tablete s podaljšanim sproščanjem	not available	5363-I-754/14	BELUPO D.O.O.	SI
DIACLAZID 30 mg tablety s prodlouženým uvolňováním	not available	18/053/14-C	BELUPO, S.R.O.	CZ
Diaclide MR 30 mg Modified-release Tablets	DE/H/0893/001	PA0577/030/002	MCDERMOTT LABORATORIES LTD	IE
Diaclide MR 60 mg modified-released tablets	PT/H/1203/001	PA0577/030/003	MCDERMOTT LABORATORIES LTD	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
Diacronal MR 30 mg modified-release tablets	DE/H/0892/001	PA1347/024/002	KRKA, D.D., NOVO MESTO	IE
Diacronal MR 60 mg Modified-release Tablets	DE/H/0894/002	PA1347/024/001	KRKA, D.D., NOVO MESTO	IE
Diagen, 30 mg tabletki o zmodyfikowanym uwalnianiu	DE/H/0893/001	17050	GENERICS [UK] LIMITED	PL
Diagen, 60 mg, tabletki o zmodyfikowanym uwalnianiu	PT/H/1203/001	22365	GENERICS [UK] LIMITED	PL
Diaglyaran 60 mg modified-release tablets	UK/H/5466/01/DC	PL 14894/0726	RANBAXY (UK) LIMITED	UK
Diaglyc 30 mg modified-release tablets	DE/H/0895/001	PA0749/076/001	TEVA PHARMA B.V.	IE
DIAMICRON 30 mg	FR/H/171/01	11420	LES LABORATOIRES SERVIER (SURESNES)	PL
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404054/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404128/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404155/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404027/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/171/01	023404104/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/171/01	023404080/M	LES LABORATOIRES SERVIER (SURESNES)	IT
Diamicron 30 mg compresse a rilascio modificato	FR/H/0171/001	023404092/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404041/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404142/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404078/M	LES LABORATOIRES SERVIER (SURESNES)	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
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404066/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404130/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/171/01	023404116/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404167/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg compresse a rilascio modificato	FR/H/0171/001	023404039/M	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 30 mg comprimidos de liberación modificada	FR/H/171/01	63.644	LES LABORATOIRES SERVIER (SURESNES)	ES
DIAMICRON 30 mg MR Tablets	FR/H/171/01	PL 05815/0019	LES LABORATOIRES SERVIER (SURESNES)	UK
DIAMICRON 30MG comprimés à libération prolongée	FR/H/171/01	2006028385	LES LABORATOIRES SERVIER (SURESNES)	LU
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404318	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404306	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404270	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404243	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404294	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404181	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404229	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404282	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404256	LES LABORATOIRES SERVIER (SURESNES)	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
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404193	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/0171/002	023404231	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404205	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404268	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404320	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg compresse a rilascio modificato	FR/H/171/02/DC	023404217	LES LABORATOIRES SERVIER (SURESNES)	IT
Diamicron 60 mg compresse a rilascio modificato	FR/H/0171/002	023404179	LES LABORATOIRES SERVIER (SURESNES)	IT
DIAMICRON 60 mg, comprimé sécable à libération modifiée	FR/H/0171/002	338 107-2	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 60 mg, comprimé sécable à libération modifiée	FR/H/0171/002	338 234-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 60 mg, comprimé sécable à libération modifiée	FR/H/0171/002	338 233-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 60 mg, comprimé sécable à libération modifiée	FR/H/0171/002	338 146-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 60 mg, comprimé sécable à libération modifiée	FR/H/0171/002	338 232-1	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 60 mg, comprimidos de liberación modificada	FR/H/0171/002	71738	LES LABORATOIRES SERVIER (SURESNES)	ES
DIAMICRON 60 mg, modified release tablet	FR/H/171/02/DC	MA 066/00603	LES LABORATOIRES SERVIER (SURESNES)	MT
DIAMICRON 60MG comprimés à libération prolongée	FR/H/171/02/DC	2010080012	LES LABORATOIRES SERVIER (SURESNES)	LU
DIAMICRON 80 mg Tablets	not available	PL 00093/0024	SERVIER LABORATORIES LTD	UK
DIAMICRON 80 MG, comprimé sécable	not available	312 936-1	LES LABORATOIRES SERVIER (SURESNES)	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
DIAMICRON 80 MG, comprimé sécable	not available	372 289-2	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 80 MG, comprimé sécable	not available	312 937-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 80 mg, comprimé sécable	not available	320 060-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON 80 mg, scored tablet	not available	MA 066/00601	LES LABORATOIRES SERVIER (SURESNES)	MT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3597788	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3515780	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516481	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516788	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516184	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516689	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3515681	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516085	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516580	LES LABORATOIRES SERVIER (SURESNES)	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
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3515988	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516382	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3515889	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3516283	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 30 mg, comprimido de libertação modificada	FR/H/171/01	3515582	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 60 mg, comprimido de libertação modificada	FR/H/0171/002	5250261	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 60 mg, comprimido de libertação modificada	FR/H/0171/002	5250303	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON LM 60 mg, comprimido de libertação modificada	FR/H/0171/002	5250279	LES LABORATOIRES SERVIER (SURESNES)	PT
DIAMICRON MR	FR/H/171/01	MA 066/00602	LES LABORATOIRES SERVIER (SURESNES)	MT
Diamicron MR 30 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/171/01	1-24000	SERVIER AUSTRIA GMBH	AT
DIAMICRON MR 30 MG, MODIFIED RELEASE TABLET	FR/H/0171/001	PA 568/13/1	LES LABORATOIRES SERVIER (SURESNES)	IE
DIAMICRON MR 30 mg, tabletten met gereguleerde afgifte	FR/H/171/01	RVG 25617	LES LABORATOIRES SERVIER (SURESNES)	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
Diamicron MR 60 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/171/02/DC	1-28884	SERVIER AUSTRIA GMBH	AT
DIAMICRON MR 60 mg, modified release tablet	FR/H/0171/002	PA 568/013/002	LES LABORATOIRES SERVIER (SURESNES)	IE
DIAMICRON MR 60 mg, tabletten met gereguleerde afgifte	FR/H/171/02/DC	RVG 102828	LES LABORATOIRES SERVIER (SURESNES)	NL
DIAMICRON MR 60 mg, δισκίο ελεγχόμενης αποδέσμευσης	FR/H/171/02/DC	78542/08-12-2010	SERVIER HELLAS	GR
Diamicron Uno	FR/H/171/02/DC	43536	LES LABORATOIRES SERVIER (SURESNES)	DK
Diamicron Uno	FR/H/0171/001	31642	LES LABORATOIRES SERVIER (SURESNES)	DK
DIAMICRON UNO 30 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/171/01	50028.00.00	LES LABORATOIRES SERVIER (SURESNES)	DE
DIAMICRON UNO 30 mg, töflur með breyttan losunarhraða	FR/H/171/01	IS/1/00/021/01	LES LABORATOIRES SERVIER (SURESNES)	IS
DIAMICRON UNO 60 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/171/02/DC	74783.00.00	LES LABORATOIRES SERVIER (SURESNES)	DE
DIAMICRON UNO 60 mg, töflur með breyttan losunarhraða	FR/H/171/02/DC	IS/1/08/100/01	LES LABORATOIRES SERVIER (SURESNES)	IS
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 184-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 185-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 179-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 182-5	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	562 674-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 186-0	LES LABORATOIRES SERVIER (SURESNES)	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
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 183-1	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 191-4	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	354 188-3	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/171/01	357 019-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	562 672-1	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	354 190-8	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	354 187-7	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	354 180-2	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	354 181-9	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 30 mg, comprimé à libération modifiée	FR/H/0171/001	372 261-0	LES LABORATOIRES SERVIER (SURESNES)	FR
DIAMICRON® 60 mg MR Tablets	FR/H/0171/002	PL 05815/0073	LES LABORATOIRES SERVIER (SURESNES)	UK
DIAMICRON® MR 30 mg, δισκίο ελεγχόμενης αποδέσμευσης	FR/H/0171/001	19662	LES LABORATOIRES SERVIER (SURESNES)	CY
DIAMICRON® MR 30 mg, δισκίο ελεγχόμενης αποδέσμευσης	FR/H/0171/001	52743/19-7-2011	SERVIER HELLAS	GR
DIAMICRON® MR 60mg, δισκίο ελεγχόμενης αποδέσμευσης	FR/H/0171/002	20710	LES LABORATOIRES SERVIER (SURESNES)	CY
DIAPREL MR	FR/H/171/01	18/469/00-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
DIAPREL MR	not available	4430	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	PL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DIAPREL MR 30 mg comprimate cu eliberare modificată	not available	7648/2015/01	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 30 mg comprimate cu eliberare modificată	not available	7648/2015/02	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/005	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/011	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/015	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/007	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/003	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/012	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/002	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/008	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/001	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/013	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/006	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/010	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/014	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/009	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 30 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/001	LT/1/05/0200/004	LES LABORATOIRES SERVIER (SURESNES)	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
Diaprel MR 30 mg módosított hatóanyagleadású tabletta	FR/H/0171/001	OGYI-T-8448/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 30 mg módosított hatóanyagleadású tabletta	FR/H/0171/001	OGYI-T-8448/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 30 mg módosított hatóanyagleadású tabletta	FR/H/0171/001	OGYI-T-8448/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/004	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/011	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/005	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/014	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/012	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/013	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/007	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/015	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/006	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/008	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/002	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/010	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/009	SERVIER PHARMA D.O.O	SI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/003	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg tablete s prirejenim sproščanjem	FR/H/171/01	H/02/00464/001	SERVIER PHARMA D.O.O	SI
DIAPREL MR 30 mg, tablety s riadeným uvoľňovaním	FR/H/0171/001	18/0023/05-S	LES LABORATOIRES SERVIER (SURESNES)	SK
DIAPREL MR 60 MG	FR/H/171/02/DC	18/022/10-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/08	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/02	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/06	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/05	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/04	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/14	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/12	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/13	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/15	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/01	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/09	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/07	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/03	LES LABORATOIRES SERVIER (SURESNES)	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
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/10	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/11	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg comprimate cu eliberare modificată	FR/H/171/02/DC	8054/2015/16	LES LABORATOIRES SERVIER (SURESNES)	RO
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/025	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/022	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/017	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/030	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/028	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/029	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/019	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/024	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/018	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/031	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/023	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/016	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/026	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/027	LES LABORATOIRES SERVIER (SURESNES)	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
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/021	LES LABORATOIRES SERVIER (SURESNES)	LT
DIAPREL MR 60 mg modifikuoto atpalaidavimo tabletės	FR/H/0171/002	LT/1/05/0200/020	LES LABORATOIRES SERVIER (SURESNES)	LT
Diaprel MR 60 mg módosított hatóanyagleadású tabletta	FR/H/0171/002	OGYI-T-8448/08	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 60 mg módosított hatóanyagleadású tabletta	FR/H/0171/002	OGYI-T-8448/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 60 mg módosított hatóanyagleadású tabletta	FR/H/0171/002	OGYI-T-8448/07	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 60 mg módosított hatóanyagleadású tabletta	FR/H/0171/002	OGYI-T-8448/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
Diaprel MR 60 mg módosított hatóanyagleadású tabletta	FR/H/0171/002	OGYI-T-8448/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	HU
DIAPREL MR 60 mg tableta s riadeným uvoľňovaním	FR/H/0171/002	18/0763/09-S	LES LABORATOIRES SERVIER (SURESNES)	SK
DIAPREL MR 60 mg tablete s prilagođenim oslobađanjem	FR/H/171/02/DC	HR-H-276485617	SERVIER PHARMA D.O.O - CROATIA	HR
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/019	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/022	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/027	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/028	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/023	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/030	SERVIER PHARMA D.O.O	SI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/025	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/031	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/020	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/024	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/026	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/021	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/029	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/018	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/017	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg tablete s prirejenim sproščanjem	FR/H/171/02/DC	H/02/00464/016	SERVIER PHARMA D.O.O	SI
DIAPREL MR 60 mg, toimeainet modifitseeritult vabastavad tabletid	FR/H/0171/002	661109	LES LABORATOIRES SERVIER (SURESNES)	EE
Diaprel MR 60mg ilgstošās darbības tabletes	FR/H/171/02/DC	09-0536	LES LABORATOIRES SERVIER (SURESNES)	LV
DIAPREL MR, 30 mg toimeainet modifitseeritult vabastav tablett	FR/H/0171/001	350401	LES LABORATOIRES SERVIER (SURESNES)	EE
DIAPREL MR, 60 mg, tabletki o zmodyfikowanym uwalnianiu	FR/H/0171/002	16641	LES LABORATOIRES SERVIER (SURESNES)	PL
DIAPREL, 80 mg tabletid	not available	150596	LES LABORATOIRES SERVIER (SURESNES)	EE
DIAZID 80 mg compresse divisibili	not available	036328019	BENEDETTI & CO. S.P.A.	IT
Diazidan, 80 mg, tabletki	not available	9111	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
Diglical MR 30 mg tablete s prilagođenim oslobađanjem	not available	UP/I-530-09/11-01 /170	PLIVA HRVATSKA D.O.O.	HR
Diglical MR 60 mg tablete s prilagođenim oslobađanjem	DK/H/2460/001	HR-H-847714460	PLIVA HRVATSKA D.O.O.	HR
DIGLICAL tablet 80 mg	not available	20060387	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Dizirel	NL/H/3108/001	874715	SANDOZ PHARMACEUTICALS D.D.	EE
Dizirel	NL/H/3384/001	901116	SANDOZ PHARMACEUTICALS D.D.	EE
Dizirel 60 mg ilgstošās darbības	NL/H/3384	16-0017	SANDOZ PHARMACEUTICALS D.D.	LV
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/001	SANDOZ PHARMACEUTICALS D.D.	LT
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/002	SANDOZ PHARMACEUTICALS D.D.	LT
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/003	SANDOZ PHARMACEUTICALS D.D.	LT
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/004	SANDOZ PHARMACEUTICALS D.D.	LT
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/005	SANDOZ PHARMACEUTICALS D.D.	LT
Dizirel 60 mg modifikuoto atpalaidavimo tabletės	NL/H/3384/001	LT/1/16/3874/006	SANDOZ PHARMACEUTICALS D.D.	LT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564032/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564071/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564158/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564020/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564057/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564018/M	IST. FARMACO BIOLOGICO STRODER 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
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564083/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564069/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564044/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564133/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564145/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564107/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564095/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564119/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 30 mg compresse a rilascio modificato	FR/H/172/01	035564121/M	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564160	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564261	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564311	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564297	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564210	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564208	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564309	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564285	IST. FARMACO BIOLOGICO STRODER 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
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564259	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564273	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564172	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564234	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564222	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564196	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564184	IST. FARMACO BIOLOGICO STRODER SRL	IT
DRAMION 60 mg compresse a rilascio modificato	FR/H/172/02/DC	035564246	IST. FARMACO BIOLOGICO STRODER SRL	IT
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/01	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/02	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/04	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/03	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/06	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/05	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/07	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/08	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/10	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
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/12	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/11	SANDOZ HUNGÁRIA KFT	HU
Dyaclid 30 mg módosított hatóanyagleadású tabletta	NL/H/3108/001	OGYI-T-22847/09	SANDOZ HUNGÁRIA KFT	HU
EDICIL MR 30 mg modified-release tablets	DE/H/0896/001	PL 15773/0777	RATIOPHARM GMBH	UK
ESQUEL 80 mg comprimate	not available	6075/2014/01	GEDEON RICHTER ROMÂNIA S.A.	RO
GALTES 80 mg compresse	not available	035940016	CRINOS S.P.A.	IT
Glibemat 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0894/001	67672.00.00	KRKA, D.D., NOVO MESTO	DE
Glibemat 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0894/002	86771.00.00	KRKA, D.D., NOVO MESTO	DE
Glicarsten 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/3525/001	86773.00.00	KRKA, D.D., NOVO MESTO	DE
GLICAZIDA Teva 30 mg Comprimidos de liberacin modificada EFG	DE/H/0892/001/R/001	673074	TEVA	ES
Glicazida-Lupin 30 mg, comprimidos de libertação prolongada	PT/H/0534/001	5592266	LUPIN (EUROPE) LIMITED	PT
Glicazida-Lupin 30 mg, comprimidos de libertação prolongada	PT/H/0534/001	5592308	LUPIN (EUROPE) LIMITED	PT
Glicazida-Lupin 30 mg, comprimidos de libertação prolongada	PT/H/0534/001	5592258	LUPIN (EUROPE) LIMITED	PT
Glicazida-Lupin 30 mg, comprimidos de libertação prolongada	PT/H/0534/001	5592274	LUPIN (EUROPE) LIMITED	PT
Gliclabore 30 mg, tabletten met gereguleerde afgifte	NL/H/1711/001	RVG 104874	WÖRWAG PHARMA GMBH & CO. KG	NL
Gliclada 30 mg ilgstošās darbības tabletes	DE/H/0892/001	08-0047	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
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/001	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/002	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/003	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/004	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/005	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/006	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/007	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/008	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/009	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/010	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/011	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/012	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/013	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/014	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/001	LT/1/08/1035/015	KRKA, D.D., NOVO MESTO	LT
Gliclada 30 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/01	KRKA, D.D., NOVO MESTO	HU
Gliclada 30 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/02	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
Gliclada 30 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/03	KRKA, D.D., NOVO MESTO	HU
Gliclada 30 mg tablete s podaljšanim sproščanjem	NP/H/0429/001	H/07/00698/001	KRKA, D.D., NOVO MESTO	SI
Gliclada 30 mg tablete s podaljšanim sproščanjem	NP/H/0429/001	H/07/00698/002	KRKA, D.D., NOVO MESTO	SI
Gliclada 30 mg tablete s podaljšanim sproščanjem	NP/H/0429/001	H/07/00698/003	KRKA, D.D., NOVO MESTO	SI
Gliclada 30 mg tablete s prilagođenim oslobađanjem	not available	HR-H-934668649	KRKA-FARMA D.O.O.	HR
Gliclada 30 mg tabletki o zmodyfikowanym uwalnianiu	DE/H/0892/001	14634	KRKA, D.D., NOVO MESTO	PL
Gliclada 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0892/001	1-27525	KRKA, D.D., NOVO MESTO	AT
Gliclada 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0892/001	67670.00.00	KRKA, D.D., NOVO MESTO	DE
Gliclada 30 mg tablety s riadeným uvoľňovaním	DE/H/0892/001	18/0065/08-S	KRKA, D.D., NOVO MESTO	SK
Gliclada 30 mg toimeainet modifitsseritult vabastavad tabletid	DE/H/0892/001	571408	KRKA, D.D., NOVO MESTO	EE
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/016	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/017	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/018	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/019	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/020	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/021	KRKA, D.D., NOVO MESTO	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
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/022	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/023	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/024	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg modifikuoto atpalaidavimo tabktės	DE/H/0892/002	LT/1/08/1035/025	KRKA, D.D., NOVO MESTO	LT
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/04	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/05	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/06	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/07	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/08	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/09	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/10	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/11	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/12	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/13	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/14	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/15	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/16	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
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/17	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/18	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/19	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/20	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/21	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/22	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/23	KRKA, D.D., NOVO MESTO	HU
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/004	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/005	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/006	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/007	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/008	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s podaljšanim sproščanjem	NP/H/0429/002	H/07/00698/009	KRKA, D.D., NOVO MESTO	SI
Gliclada 60 mg tablete s prilagođenim oslobađanjem	not available	UP/I-530-09/12-01/01	KRKA-FARMA D.O.O.	HR
Gliclada 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0892/002	1-31795	KRKA, D.D., NOVO MESTO	AT
Gliclada 60 mg tbl mod	DE/H/0892/002	18/0132/13-S	KRKA, D.D., NOVO MESTO	SK
Gliclada 60 mg, Ilgstošās darbības tabletes	DE/H/0892/002	13-0010	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
Gliclada 60 mg, toimeainet modifitseeritult vabastavad tabletid	DE/H/0892/002	807313	KRKA, D.D., NOVO MESTO	EE
Gliclada 60mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0892/002	86770.00.00	KRKA, D.D., NOVO MESTO	DE
Gliclada 90 mg tablety s riadeným uvoľňovaním	DE/H/0892/003	18/0094/16-S	KRKA, D.D., NOVO MESTO	SK
Gliclada 90 mg ilgstošās darbības tabletes	DE/H/0892/003	16-0027	KRKA, D.D., NOVO MESTO	LV
Gliclada 90 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/003	LT/1/08/1035/026	KRKA, D.D., NOVO MESTO	LT
Gliclada 90 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/003	LT/1/08/1035/027	KRKA, D.D., NOVO MESTO	LT
Gliclada 90 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/003	LT/1/08/1035/028	KRKA, D.D., NOVO MESTO	LT
Gliclada 90 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/003	LT/1/08/1035/029	KRKA, D.D., NOVO MESTO	LT
Gliclada 90 mg modifikuoto atpalaidavimo tabletės	DE/H/0892/003	LT/1/08/1035/030	KRKA, D.D., NOVO MESTO	LT
Gliclada 90 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/24	KRKA, D.D., NOVO MESTO	HU
Gliclada 90 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/25	KRKA, D.D., NOVO MESTO	HU
Gliclada 90 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/26	KRKA, D.D., NOVO MESTO	HU
Gliclada 90 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/27	KRKA, D.D., NOVO MESTO	HU
Gliclada 90 mg módosított hatóanyagleadású tabletta	not available	OGYI-T-20515/28	KRKA, D.D., NOVO MESTO	HU
Gliclada 90 mg tablete s prilagođenim oslobađanjem	DE/H/0892/003	HR-H-976762113	KRKA-FARMA D.O.O.	HR
Gliclada 90 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0892/003	136843	Krka, d. d., Novo mesto	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
Gliclada tabletki o zmodyfikowanym uwalnianiu, 60 mg	DE/H/0892/002	21068	KRKA, D.D., NOVO MESTO	PL
Gliclada, 90 mg toimeainet modifitseeritult vabastavad tabletid	DE/H/0892/003	903116	KRKA, D.D., NOVO MESTO	EE
Gliclada, 90 mg, tabletki o przedłużonym uwalnianiu	DE/H/0892/003	23056	KRKA, D.D., NOVO MESTO	PL
Glicladzida Krka 30 mg comprimidos de liberación modificada EFG	DE/H/0894/001	78226	KRKA, D.D., NOVO MESTO	ES
Gliclagamma MR	NL/H/1711/001/DC	18164	WÖRWAG PHARMA GMBH & CO. KG	PL
Gliclalane 30 mg, tabletten met gereguleerde afgifte	NL/H/1705/001	RVG 104870	DISPHAR INTERNATIONAL B.V.	NL
Gliclareze 30 mg, tabletten met gereguleerde afgifte	NL/H/1709/001	RVG 104872	DISPHAR INTERNATIONAL B.V.	NL
Gliclasan 30 mg tabletten met gereguleerde afgifte	NL/H/3108/001	BE474177	SANDOZ N.V.	BE
Gliclastad 30 mg, tabletki o przedłużonym uwalnianiu	PT/H/0534/001	20493	STADA ARZNEIMITTEL AG	PL
Gliclatim, tabletter med modificeret udløsning	DK/H/2270/001	51775	STADA ARZNEIMITTEL AG	DK
Gliclazid "Actavis", tabletter med modificeret udløsning	DK/H/2376/001	53938	ACTAVIS GROUP PTC EHF.	DK
Gliclazid "Actavis", tabletter med modificeret udløsning	DK/H/2376/002	53939	ACTAVIS GROUP PTC EHF.	DK
Gliclazid "Sigillata", tabletter med modificeret udløsning	DK/H/2377/001	53940	SIGILLATA LIMITED	DK
Gliclazid "Sigillata", tabletter med modificeret udløsning	DK/H/2377/002	53941	SIGILLATA LIMITED	DK
Gliclazid ”HCS”, tabletter med modificeret udløsning	DK/H/2195/001	50744	HCS BVBA	DK
Gliclazid ”HCS”, tabletter med modificeret udløsning	DK/H/2195/002	50745	HCS BVBA	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
Gliclazid ”Stada”, tabletter med modificeret udløsning	DK/H/2301/001	52501	STADA ARZNEIMITTEL AG	DK
Gliclazid ”Teva B.V.”, tabletter med modificeret udløsning	DK/H/2460/001	55042	TEVA B.V	DK
Gliclazid ”Teva”	DE/H/0895/001	40070	TEVA DENMARK A/S	DK
Gliclazid Arcana 30 mg - Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0893/001	1-28537	ARCANA ARZNEIMITTEL GMBH	AT
Gliclazid axcount 30 mg Tabletten mit veränderter Wirkstofffreisetzung	UK/H/5530/001/DC	90888.00.00	AXCOUNT GENERIKA GMBH	DE
Gliclazid Genericon 30 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/11-01/473	GENERICON PHARMA D.O.O.	HR
Gliclazid Genericon 30 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	1-30053	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Gliclazid Genericon 30 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	1-30054	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Gliclazid Krka 30 mg Tabletter med modificeret udløsning	DE/H/0892/001	MTNR. 40069	KRKA SVERIGE AB	DK
Gliclazid Krka 30 mg tbl mod	DE/H/0894/001	18/0498/13-S	KRKA, D.D., NOVO MESTO	SK
Gliclazid Krka 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/3524/001	86772.00.00	KRKA, D.D., NOVO MESTO	DE
Gliclazid Krka 60 mg Tabletter med modificeret udløsning	DE/H/0892/002	MTNR. 50103	KRKA SVERIGE AB	DK
Gliclazid Krka 60 mg tbl mod	DE/H/0894/002	18/0133/13-S	KRKA, D.D., NOVO MESTO	SK
Gliclazid Mylan 30 mg tablety s riadeným uvolnovaním	DE/H/0893/001	18/0461/09-S	GENERICS [UK] LIMITED	SK
Gliclazid Mylan 30 mg tablety s řízeným uvolňováním	DE/H/0893/001	18/035/10-C	GENERICS [UK] LIMITED	CZ
Gliclazid Mylan 60 mg tablety s predĺženým uvoľňovaním	PT/H/1203/001	18/0067/15-S	GENERICS [UK] LIMITED	SK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gliclazid ratiopharm 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DK/H/2460/001	136638	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Gliclazid Sandoz 30 mg - Tabletten mit veränderter Wirkstofffreisetzung	NL/H/1700/001	1-29677	SANDOZ GMBH	AT
Gliclazid STADA 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DK/H/2301/001	135744	STADA ARZNEIMITTEL GMBH	AT
Gliclazid Uno "Servier"	FR/H/172/02/DC	43535	LES LABORATOIRES SERVIER (SURESNES)	DK
Gliclazid Uno "Servier"	FR/H/172/01	31643	LES LABORATOIRES SERVIER (SURESNES)	DK
Gliclazidă Actavis 30 mg comprimate cu eliberare modificată.	DK/H/2376/001	7835/2015/01-24	ACTAVIS GROUP PTC EHF.	RO
Gliclazidă Actavis 60 mg comprimate cu eliberare modificată.	DK/H/2376/002	7836/2015/01-24	ACTAVIS GROUP PTC EHF.	RO
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052741	ALTER, S.A.	PT
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052733	ALTER, S.A.	PT
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052725	ALTER, S.A.	PT
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052717	ALTER, S.A.	PT
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052709	ALTER, 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
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052675	ALTER, S.A.	PT
Gliclazida Alter 30 mg Comprimidos de libertação prolongada	not available	5052758	ALTER, S.A.	PT
Gliclazida Alter 60 mg comprimidos de libertação modificada	not available	5643754	ALTER, S.A.	PT
Gliclazida Apotex 30 mg, comprimido de libertação prolongada	IT/H/0311/001/DC	5300462, 5300470, 5300504, 5300512, 5300520, 5300538, 5300546, 5300553, 5300561, 5300579	APOTEX EUROPE BV	PT
Gliclazida AZEVEDOS 80 mg comprimidos	not available	5938485	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Gliclazida Azevedos 80 mg comprimidos	not available	5938386	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Gliclazida Bluepharma 80 mg comprimidos	not available	4392080	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Gliclazida Bluepharma 80 mg comprimidos	not available	4392189	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Gliclazida Bluepharma 80 mg comprimidos	not available	4391983	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Gliclazida Brill Pharma 30 mg comprimidos de liberación modificada EFG	UK/H/5530/001	79808	BRILL PHARMA, S.L.	ES
Gliclazida Cinfa 30 mg comprimidos de liberación modificada EFG	not available	78.439	LABORATORIOS CINFA, 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
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067749	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067756	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067764	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067772	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067806	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067814	CINFA PORTUGAL, LDA.	PT
Gliclazida Cinfa 30 mg Comprimidos de libertação prolongada	not available	5067822	CINFA PORTUGAL, LDA.	PT
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/15	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/01	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/02	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/03	GENERICS [UK] LIMITED	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
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/04	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/05	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/06	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/07	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/08	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/09	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/10	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/11	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/13	GENERICS [UK] LIMITED	RO
Gliclazida Generics 30 mg comprimate cu eliberare modificata	DE/H/0893/001	6910/2014/14	GENERICS [UK] LIMITED	RO
Gliclazidă Generics 30 mg comprimate cu eliberare modificată	DE/H/0893/001	6910/2014/12	GENERICS [UK] LIMITED	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
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032305	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032313	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032321	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032339	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032347	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032354	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 30 mg Comprimidos de libertação prolongada	not available	5032362	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465711	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465679	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465703	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465729	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
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465737	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 60 mg Comprimidos de libertação modificada	not available	5465745	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 80 mg Comprimidos	not available	5400197	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Generis 80 mg Comprimidos	not available	5400296	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida GP 80 mg comprimidos	not available	4782082	GP – GENÉRICOS PORTUGUESES, LDA.	PT
Gliclazida GP 80 mg comprimidos	not available	4782181	GP – GENÉRICOS PORTUGUESES, LDA.	PT
Gliclazida Helm 80 mg comprimido	not available	5661400	HELM PORTUGAL, LDA.	PT
Gliclazida Helm 80 mg comprimido	not available	5661418	HELM PORTUGAL, LDA.	PT
Gliclazida Jaba 80 mg Comprimidos	not available	5400692	JABA RECORDATI, S.A.	PT
Gliclazida Jaba 80 mg Comprimidos	not available	5400593	JABA RECORDATI, S.A.	PT
Gliclazida KERN PHARMA 30 mg comprimidos de liberación modificada EFG	not available	74040	KERN PHARMA, S.L.	ES
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081401	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081419	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081427	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081435	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081443	KRKA FARMACÊUTICA, 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
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081450	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081468	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081476	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081500	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimido de libertação modificada	DE/H/0892/001	5081518	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimidos de libertação modificada	DE/H/0892/001	5081526	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimidos de libertação modificada	DE/H/0892/001	5081534	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimidos de libertação modificada	DE/H/0892/001	5081542	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimidos de libertação modificada	DE/H/0892/001	5081559	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 30 mg Comprimidos de libertação modificada	DE/H/0892/001	5081567	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/01	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/02	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/03	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/04	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/05	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
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/06	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/07	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/08	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/09	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/10	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/11	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/12	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/13	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/14	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/15	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/16	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/17	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/18	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/19	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg comprimate cu eliberare modificată	DE/H/0894/002	5468/2013/20	KRKA, D.D., NOVO MESTO	RO
Gliclazida Krka 60 mg Comprimidos de liberación modificada EFG	DE/H/0894/002	77493	KRKA, D.D., NOVO MESTO	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465612	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465620	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465638	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465646	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465653	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Krka 60 mg comprimidos de libertação modificada	not available	5465661	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Gliclazida Labesfal 80 mg comprimidos	not available	3983491	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal 80 mg comprimidos	not available	3983699	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal 80 mg comprimidos	not available	3983590	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝo modificada	not available	5603428	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝo modificada	not available	5603451	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝo modificada	not available	5603469	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
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝ o modificada	not available	5603436	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝ o modificada	not available	5603444	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LM 60 mg comprimidos de liberta��ァ��ゝ o modificada	not available	5603477	GENERIS FARMACÊUTICA, S.A.	PT
Gliclazida Labesfal LP	not available	5574959 5574975 5575006 5574967	Generis Farmacêutica, S.A.	PT
Gliclazida Lek 30 mg comprimidos de liberación modificada EFG	NL/H/3108/001	79929	SANDOZ FARMACÉUTICA, S.A.	ES
Gliclazida Lupin 30 mg comprimidos de liberación modificada	not available	76714	STADA Genéricos S.L.,	ES
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/01	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/02	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/03	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/04	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/05	LUPIN (EUROPE) LIMITED	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
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/06	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/07	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/08	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/09	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/10	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/11	LUPIN (EUROPE) LIMITED	RO
Gliclazida Lupin 60 mg comprimate cu eliberare prelungita	PT/H/1272/001/DC	8292/2015/12	LUPIN (EUROPE) LIMITED	RO
Gliclazida Mesiproc	NL/H/3108/001	5652938	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Mesiproc	NL/H/3108/001	5652946	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Mesiproc	NL/H/3108/001	5652953	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Mesiproc	NL/H/3108/001	5652961	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Mylan 30 mg comprimidos de liberación modificada EFG	DE/H/0897/001	69734	MYLAN PHARMACEUTICALS S.L.	ES
Gliclazida Mylan 30 mg, comprimidos de libertação modificada	DE/H/0893/001	5210208	MYLAN, LDA	PT
Gliclazida Mylan 30 mg, comprimidos de libertação modificada	DE/H/0893/001	5210216	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
Gliclazida Mylan 60 mg comprimido de libertação modificada	PT/H/1203/001	5642640	MYLAN, LDA	PT
Gliclazida Mylan 60 mg comprimido de libertação modificada	PT/H/1203/001	5642624	MYLAN, LDA	PT
Gliclazida Mylan 60 mg comprimido de libertação modificada	PT/H/1203/001	5642632	MYLAN, LDA	PT
Gliclazida NORMON 30 mg comprimidos de liberación modificada EFG	not available	75377	LABORATORIOS NORMON, S.A.	ES
Gliclazida Placasod 30 mg comprimidos de liberación modificada EFG	NL/H/1700/001	72991	SANDOZ FARMACÉUTICA, S.A.	ES
Gliclazida Ranbaxy 60 mg comprimidos de liberación modificada EFG	UK/H/5466/01/DC	79994	LABORATORIOS RANBAXY S.L.	ES
Gliclazida ratiopharm 60 mg comprimidos de libertação modificada	DK/H/2460/001	5666219	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm 60 mg comprimidos de libertação modificada	DK/H/2460/001	5666227	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm 60 mg comprimidos de libertação modificada	DK/H/2460/001	5666235	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm 60 mg comprimidos de libertação modificada	DK/H/2460/001	5666243	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081740	RATIOPHARM-COMERCIO E INDUSTRIA DE 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
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081757	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081773	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081765	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081807	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081864	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081823	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081906	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081732	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081856	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081872	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081724	RATIOPHARM-COMERCIO E INDUSTRIA DE 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
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081831	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081815	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida ratiopharm, 30 mg, comprimidos de libertação modificada	DE/H/0896/001	5081849	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Gliclazida Sandoz	NL/H/1700/001	5321708	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Sandoz	NL/H/1700/001	5321724	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Sandoz	NL/H/1700/001	5321674	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida Sandoz	NL/H/1700/001	5321716	SANDOZ FARMACÊUTICA LDA.	PT
Gliclazida STADA 30 mg comprimidos de liberación modificada	DK/H/2270/001	78.483	LABORATORIO STADA, S.L.	ES
Gliclazida STADA Genéricos 30 mg comprimidos de liberación modificada EFG	PT/H/0534/001	76.714	STADA GENÉRICOS, S.L.	ES
Gliclazida STADA Genéricos 60 mg comprimidos de liberación modificada	DK/H/2301/001	79.204	STADA GENÉRICOS, S.L.	ES
Gliclazida TAD 30 mg comprimidos de libertação modificada	DE/H/0894/001	5588819	TAD PHARMA GMBH	PT
Gliclazida TAD 30 mg comprimidos de libertação modificada	DE/H/0894/001	5588827	TAD PHARMA GMBH	PT
Gliclazida TAD 30 mg comprimidos de libertação modificada	DE/H/0894/001	5588835	TAD PHARMA GMBH	PT
Gliclazida TAD 60 mg comprimidos de libertação modificada	DE/H/0894/002	5559133	TAD PHARMA GMBH	PT
Gliclazida TAD 60 mg comprimidos de libertação modificada	DE/H/0894/002	5559141	TAD PHARMA GMBH	PT
Gliclazida TAD 60 mg comprimidos de libertação modificada	DE/H/0894/002	5559158	TAD PHARMA GMBH	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
Gliclazida TAD 60 mg comprimidos de libertação modificada	DE/H/0894/002	5559166	TAD PHARMA GMBH	PT
Gliclazida TecniGen 30 mg comprimidos de liberación modificada EFG	not available	74.434	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/01	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/02	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/03	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/04	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/05	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/06	TERAPIA S.A.	RO
Gliclazidă Terapia 60 mg comprimate cu eliberare modificată	UK/H/5466/001	8149/2015/07	TERAPIA S.A.	RO
Gliclazida Teva 30 mg comprimidos de liberación modificada EFG	DE/H/0892/001	69740	TEVA GENERICOS ESPANOLA S.L.	ES
Gliclazida Teva 30 mg Comprimidos de libertação modificada	DE/H/0895/001	5081633	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazida Teva 30 mg Comprimidos de libertação modificada	DE/H/0895/001	5081658	TEVA PHARMA – PRODUTOS FARMACÊUTICOS 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
Gliclazida Teva 30mg comprimido de libertação modificada	DE/H/0895/001	5081641	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazida Teva 30mg comprimido de libertação modificada	DE/H/0895/001	5081625	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazidă Teva 60 mg comprimate cu eliberare modificată	DK/H/2460/001	8587/2016/01-08	TEVA PHARMACEUTICALS S.R.L	RO
Gliclazida Teva 60 mg comprimidos de libertação modificada	DK/H/2494/001	5666268	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazida Teva 60 mg comprimidos de libertação modificada	DK/H/2494/001	5666300	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazida Teva 60 mg comprimidos de libertação modificada	DK/H/2494/001	5666250	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Gliclazida Teva 60 mg comprimidos de libertação modificada	DK/H/2494/001	5666276	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
GLICLAZIDA ZENTIVA 30 MG COMPRIMIDOS DE LIBERACION MODIFICADA EFG	DK/H/2302/001	79125	ZENTIVA, K.S.	ES
GLICLAZIDA ZENTIVA 30 MG COMPRIMIDOS DE LIBERTAÇÃO MODIFICADA	DK/H/2302/001	5633268	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 30 MG COMPRIMIDOS DE LIBERTAÇÃO MODIFICADA	DK/H/2302/001	5625173	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 30 MG COMPRIMIDOS DE LIBERTAÇÃO MODIFICADA	DK/H/2302/001	5625207	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Gliclazidă Zentiva 60 mg comprimate cu eliberare modificată	DK/H/2302/002	6788/2014/03	ZENTIVA, K.S.	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
Gliclazidă Zentiva 60 mg comprimate cu eliberare modificată	DK/H/2302/002	6788/2014/04	ZENTIVA, K.S.	RO
Gliclazidă Zentiva 60 mg comprimate cu eliberare modificată	DK/H/2302/002	6788/2014/01	ZENTIVA, K.S.	RO
Gliclazidă Zentiva 60 mg comprimate cu eliberare modificată	DK/H/2302/002	6788/2014/02	ZENTIVA, K.S.	RO
GLICLAZIDA ZENTIVA 60 MG COMPRIMIDOS DE LIBERTAÇÃO MODIFICADA	DK/H/2302/002	5625223	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 60 MG COMPRIMIDOS DE LIBERTAÇÃO MODIFICADA	DK/H/2302/002	5625215	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 80 MG COMPRIMIDOS REVESTIDOS	not available	3576899	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 80 MG COMPRIMIDOS REVESTIDOS	not available	3576790	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
GLICLAZIDA ZENTIVA 80 MG COMPRIMIDOS REVESTIDOS	not available	3576998	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Gliclazide 30 mg PCH, tabletten met gereguleerde afgifte	DE/H/0895/001	RVG 34591	PHARMACHEMIE BV	NL
Gliclazide 40mg Tablets	not available	PL40496/0040	BRILL PHARMA LIMITED	UK
Gliclazide 40mg Tablets	not available	PL 30306/0631	ACTAVIS GROUP PTC EHF.	UK
Gliclazide 60 mg MR Tablets	FR/H/172/002	PL 05815/0074	LES LABORATOIRES SERVIER (SURESNES)	UK
Gliclazide 80 mg Tablets	not available	PL 20416/0279	CRESCENT PHARMA LIMITED	UK
Gliclazide 80 mg Tablets	not available	PL 06464/2062	WAYMADE PLC	UK
Gliclazide 80mg Tablets	not available	AA154/07401	WOCKHARDT UK LTD	MT
Gliclazide 80mg Tablets	not available	PL 17907/0068	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Gliclazide 80mg Tablets	not available	PL 44473/0001	ECOGEN EUROPE LIMITED	UK
Gliclazide 80mg Tablets	not available	PL 21880/0070	MEDREICH PLC	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gliclazide 80mg Tablets	not available	PL 16363/0006	MILPHARM LIMITED	UK
Gliclazide 80mg tablets	not available	PL 20395/0046	RELON CHEM LIMITED	UK
Gliclazide 80mg Tablets	not available	PL 00289/0480	TEVA UK LIMITED	UK
Gliclazide 80mg Tablets	not available	29831/0103	WOCKHARDT UK LTD	UK
Gliclazide 80mg Tablets BP	not available	AA054/04301	ACCORD HEALTHCARE LIMITED	MT
Gliclazide Actavis 30 mg ilgstošas darbibas tabletes	DK/H/2376/001	15-0087	ACTAVIS GROUP PTC EHF.	LV
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/001	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/002	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/003	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/004	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/005	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/006	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/007	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/008	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/009	ACTAVIS GROUP PTC EHF.	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
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/010	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/011	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/001	LT/1/15/3723/012	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/01	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/02	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/03	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/04	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/05	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/06	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 30 mg töflur með breyttan losunarhraða	DK/H/2376/001	IS/1/15/053/01	ACTAVIS GROUP PTC EHF.	IS
Gliclazide Actavis 60 mg ilgstošās darbības tabletes	DK/H/2376/002	15-0088	ACTAVIS GROUP PTC EHF.	LV
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/013	ACTAVIS GROUP PTC EHF.	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
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/014	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/015	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/016	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/017	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/018	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/019	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/020	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/021	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/022	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/023	ACTAVIS GROUP PTC EHF.	LT
Gliclazide Actavis 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2376/002	LT/1/15/3723/024	ACTAVIS GROUP PTC EHF.	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
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/07	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/08	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/09	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/10	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/11	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg módosított hatóanyagleadású tabletta	DK/H/2376/001-002	OGYI-T-22828/12	ACTAVIS GROUP PTC EHF.	HU
Gliclazide Actavis 60 mg töflur með breyttan losunarhraða	DK/H/2376/002	IS/1/15/053/02	ACTAVIS GROUP PTC EHF.	IS
Gliclazide Actavis, 30 mg toimeainet modifitseeritult vabastavad tabletid	DK/H/2376/001	871915	ACTAVIS GROUP PTC EHF.	EE
Gliclazide Actavis, 30 mg, tabletki o zmodyfikowanym uwalnianiu	DK/H/2376/001	22543	ACTAVIS GROUP PTC EHF.	PL
Gliclazide Actavis, 60 mg toimeainet modifitseeritult vabastavad tabletid	DK/H/2376/002	871815	ACTAVIS GROUP PTC EHF.	EE
Gliclazide Actavis, 60 mg, tabletki o zmodyfikowanym uwalnianiu	DK/H/2376/002	22544	ACTAVIS GROUP PTC EHF.	PL
GLICLAZIDE ALMUS 80 mg compresse	not available	036245013	ALMUS S.R.L	IT
Gliclazide Apotex 80 mg, tabletten met gereguleerde afgifte	not available	RVG 26546	APOTEX EUROPE 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
GLICLAZIDE ARROW 30 mg, comprimé à libération modifiée	not available	NL 37706	ARROW GENERIQUES	FR
GLICLAZIDE ARROW 60 mg, comprimé à libération modifiée	DE/H/3525/001	NL 41932	ARROW GENERIQUES	FR
GLICLAZIDE ARROW 80 mg, comprimé sécable	not available	NL 26115	ARROW GENERIQUES	FR
Gliclazide Aurobindo 80 mg, tabletten met gereguleerde afgifte	not available	RVG 16011=56872	AUROBINDO PHARMA B.V.	NL
GLICLAZIDE BGR® 30 mg, comprimé à libération modifiée	FR/H/0337/001	3400939049892	BIOGARAN	FR
GLICLAZIDE BGR® 30 mg, comprimé à libération modifiée	FR/H/0337/001	3400939050034	BIOGARAN	FR
GLICLAZIDE BGR® 30 mg, comprimé à libération modifiée	FR/H/0337/001	3400941899973	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 60 mg, comprimé sécable à libération modifiée	not available	3400927586231	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 60 mg, comprimé sécable à libération modifiée	not available	3400927586750	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 80 mg, comprimé sécable	not available	3400934903069	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 80 mg, comprimé sécable	not available	3400934902819	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 80 mg, comprimé sécable	not available	3400937842846	BIOGARAN	FR
GLICLAZIDE BIOGARAN® 80 mg, comprimé sécable	not available	3400934902987	BIOGARAN	FR
Gliclazide Bristol 30 mg, tabletten met verlengde afgif	UK/H/5530/001	RVG 114159	BRISTOL LABORATORIES LTD (BERKHAMSTED)	NL
Gliclazide CF 80 mg, tabletten met gereguleerde afgifte	not available	RVG 56872	CENTRAFARM B.V.	NL
GLICLAZIDE CRISTERS 80 mg, comprimé sécable	not available	34009 385 541 7 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
GLICLAZIDE CRISTERS 80 mg, comprimé sécable	not available	34009 385 542 3 6	CRISTERS	FR
GLICLAZIDE CRISTERS 80 mg, comprimé sécable	not available	34009 373 159 5 1	CRISTERS	FR
GLICLAZIDE CRISTERS 80 mg, comprimé sécable	not available	34009 562 369 7 5	CRISTERS	FR
GLICLAZIDE CRISTERS 80 mg, comprimé sécable	not available	34009 373 160 3 3	CRISTERS	FR
GLICLAZIDE DOC Generici 80 mg Compresse	not available	036528014	DOC GENERICI S.R.L.	IT
GLICLAZIDE EG 60 mg, comprimé sécable à libération modifiéeé	DK/H/2301/001	NL435336	EG LABO - LABORATOIRES EUROGENERICS	FR
GLICLAZIDE EG 80 mg compresse	not available	036282010	EG SPA	IT
GLICLAZIDE EG LABO Laboratoires EuroGenerics, 30 mg, comprimé à libération modifiée	DK/H/2270/001	NL43030	EG LABO - LABORATOIRES EUROGENERICS	FR
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020019	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020021	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020058	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020033	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020072	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020060	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
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020159	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020096	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020084	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020045	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020108	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020134	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020146	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020110	EG SPA	IT
GLICLAZIDE EUROGENERICI 30 mg compresse a rilascio modificato	DE/H/0892/001	039020122	EG SPA	IT
Gliclazide GAMMA 30 mg MR	NL/H/1711/001/DC	18/0680/10-S	WÖRWAG PHARMA GMBH & CO. KG	SK
GLICLAZIDE ISOMED 30 mg, comprimé à libération modifiée	not available	NL39120	TEVA SANTÉ	FR
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038017	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038029	KRKA, D.D., NOVO MESTO	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
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038031	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038043	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038056	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038068	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038070	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038082	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038094	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038106	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038118	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038120	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038132	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038144	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg compresse a rilascio modificato	DE/H/0894/001	AIC N. 039038157	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 30 mg toimeainet modifitseeritult vabastavad tabletid	DE/H/0894/001	828113	KRKA, D.D., NOVO MESTO	EE
Gliclazide Krka 30 mg, comprimé à libération modifée	DK/H/2195/001	NL 42429, CIS: 6 548 524 5 (34009 273 688 6 8)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 30 mg, comprimé à libération modifée	DK/H/2195/001	NL 42429, CIS: 6 548 524 5 (34009 273 689 2 9)	KRKA, D.D., NOVO MESTO	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gliclazide Krka 30 mg, comprimé à libération modifée	DK/H/2195/001	NL 42429, CIS: 6 548 524 5 (34009 275 169 6 2)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038169	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038171	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038183	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038195	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038207	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038219	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038221	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038233	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038245	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038258	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038260	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038272	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038284	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038296	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038308	KRKA, D.D., NOVO MESTO	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
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038310	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038322	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038334	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038346	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg compresse a rilascio modificato	DE/H/0894/002	AIC N. 039038359	KRKA, D.D., NOVO MESTO	IT
Gliclazide Krka 60 mg toimeainet modifitseeritult vabastav tablett	DE/H/0894/002	806213	KRKA, D.D., NOVO MESTO	EE
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 703 5 9)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 704 1 0)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 705 8 8)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 706 4 9)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 707 0 0)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 60 mg, comprimé à libération modifée	DK/H/2195/002	NL 42430, CIS: 6 232 635 5 (34009 273 708 7 8)	KRKA, D.D., NOVO MESTO	FR
Gliclazide Krka 90 mg tabletten met gereguleerde afgifte	DE/H/0894/003	BE488053	KRKA, D.D., NOVO MESTO	BE
Gliclazide Krka tabletki o zmodyfikowanym uwalnianiu, 30 mg	DE/H/0894/001	21665	KRKA, D.D., NOVO MESTO	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
Gliclazide Krka tabletki o zmodyfikowanym uwalnianiu, 60 mg	DE/H/0894/002	21064	KRKA, D.D., NOVO MESTO	PL
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462248	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462299	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462236	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462325	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462263	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462275	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462287	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462251	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462313	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462301	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
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462059	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462097	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462010	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462085	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462061	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462046	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462073	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462109	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462022	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462111	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462034	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
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462135	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462123	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462174	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462198	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462147	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462150	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462186	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462224	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462212	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462200	EG SPA	IT
GLICLAZIDE LABORATORI EUROGENERICI 30 mg compresse a rilascio modificato	DK/H/2270/001	042462162	EG SPA	IT
Gliclazide Lupin 60 mg comprimés à libération prolongée	PT/H/1272/001	BE475742	LUPIN (EUROPE) LIMITED	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
Gliclazide Lupin 60 mg ilgstošās darbības tabletes	PT/H/1272/001	15-0186	LUPIN (EUROPE) LIMITED	LV
GLICLAZIDE MOLTENI 80 mg compresse	not available	033363019	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Gliclazide MR 30mg prolonged-release tablets	UK/H/5530/01/DC	PA1240/010/001	BRISTOL LABORATORIES LTD (BERKHAMSTED)	IE
Gliclazide MR Servier 60mg ilgstošās darbības tabletes	FR/H/172/02/DC	09-0537	LES LABORATOIRES SERVIER (SURESNES)	LV
Gliclazide MR Stada	DK/H/2301/001	18/0396/14-S	STADA ARZNEIMITTEL AG	SK
Gliclazide Mylan 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0893/001	67671.00.00	MYLAN S.A.S	DE
GLICLAZIDE MYLAN 30 mg, comprimé à libération modifiée	DE/H/0893/001	NL33442	MYLAN S.A.S	FR
Gliclazide Mylan 80 mg, comprimé sécable	not available	NL24598	MYLAN S.A.S	FR
GLICLAZIDE MYLAN GENERICS 80 mg compresse	not available	036244010	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469019	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469021	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469033	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469045	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469058	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
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469060	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469072	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469084	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469096	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469108	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469110	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469122	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469134	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469146	MYLAN S.P.A.	IT
GLICLAZIDE MYLAN GENERICS ITALIA 30 mg compresse a rilascio modificato	DE/H/0893/001	038469159	MYLAN S.P.A.	IT
Gliclazide Pensa 80 mg compresse	not available	036643017	PENSA PHARMA S.P.A.	IT
Gliclazide Qualimed 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0897/001	67675.00.00	MYLAN DURA 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
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644018	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644032	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644020	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644044	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644057	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644069	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg compresse a rilascio modificato	UK/H/5466/001	043644071	RANBAXY ITALIA S.P.A.	IT
Gliclazide Ranbaxy 60 mg, tabletten met gereguleerde afgifte	UK/H/5466/01/DC	RVG 115253	RANBAXY (UK) LIMITED	NL
GLICLAZIDE RANBAXY 60 mg, comprimé à libération modifiée	UK/H/5466/001/DC	NL 44462	RPG	FR
GLICLAZIDE RATIOPHARM 30 mg, comprimé à libération modifiée	DE/H/0896/001	NL33445	RATIOPHARM GMBH	FR
GLICLAZIDE RATIOPHARM 80 mg, comprimé sécable	not available	NL24815	TEVA SANTÉ	FR
Gliclazide retard CF 30 mg, tabletten met gereguleerde afgifte	DK/H/2270/001	RVG 112930	CENTRAFARM B.V.	NL
Gliclazide Retard Mylan 30 mg comprimés à libération modifiée	DE/H/0893/001	BE345651	MYLAN BVBA/SPRL	BE
Gliclazide Retard Mylan 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0893/001	BE345642	MYLAN BVBA/SPRL	BE
Gliclazide Retard Mylan 30 mg, tabletten met gereguleerde afgifte	DE/H/0893/001	RVG 34590	MYLAN B.V.	NL
Gliclazide Retard Mylan 60 mg, comprimé à libération modifiée	PT/H/1203/001	BE469564	MYLAN BVBA/SPRL	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
Gliclazide Retard Mylan 60 mg, comprimé à libération modifiée	PT/H/1203/001	BE469573	MYLAN BVBA/SPRL	BE
Gliclazide Retard Mylan 60 mg, comprimé à libération modifiée	PT/H/1203/001	2015120197	MYLAN BVBA/SPRL	LU
Gliclazide Retard Mylan 80 mg, tabletten met gereguleerde afgifte	not available	RVG 26307	MYLAN B.V.	NL
GLICLAZIDE RPG 30 mg, comprimé à libération modifiée	not available	NL 37705	RANBAXY PHARMACIE GENERIQUES	FR
GLICLAZIDE RPG 80 mg, comprimé sécable	not available	NL 25077	RANBAXY PHARMACIE GENERIQUES	FR
Gliclazide Sandoz 30 mg comprimés à libération modifiée	NL/H/3108/001	2015120198	SANDOZ N.V.	LU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/01	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/02	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/03	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/04	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/05	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/06	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/07	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 30 mg retard tabletta	PT/H/0534/001	OGYI-T-22298/08	SANDOZ HUNGÁRIA KFT	HU
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 628-5	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 629-1	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 631-6	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
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 632-2	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 633-9	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 634-5	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 635-1	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 636-8	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 637-4	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 638-0	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 639-7	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 640-5	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 641-1	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 626-2	SANDOZ	FR
GLICLAZIDE SANDOZ 30 mg, comprimé à libération modifiée	DE/H/0894/001	385 627-9	SANDOZ	FR
Gliclazide Sandoz 30 mg, tablet met gereguleerde afgifte	NL/H/1700/001	BE375873	SANDOZ N.V.	BE
Gliclazide Sandoz 30 mg, tablet met gereguleerde afgifte	NL/H/1700/001	BE375864	SANDOZ N.V.	BE
Gliclazide Sandoz 30 mg, tabletten met gereguleerde afgifte	NL/H/3108/001	RVG 114941	SANDOZ B.V.	NL
Gliclazide Sandoz 60 mg módosított hatóanyagleadású tabletta	NL/H/3384/001	OGYI-T-22298/09	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
Gliclazide Sandoz 60 mg módosított hatóanyagleadású tabletta	NL/H/3384/001	OGYI-T-22298/10	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 60 mg módosított hatóanyagleadású tabletta	NL/H/3384/001	OGYI-T-22298/11	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 60 mg módosított hatóanyagleadású tabletta	NL/H/3384/001	OGYI-T-22298/12	SANDOZ HUNGÁRIA KFT	HU
Gliclazide Sandoz 60 mg tabletten met gereguleerde afgifte	NL/H/3384/001	BE488035	SANDOZ N.V.	BE
Gliclazide Sandoz 60 mg tabletten met gereguleerde afgifte	NL/H/3384/001	BE488044	SANDOZ N.V.	BE
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	368 184-5	SANDOZ	FR
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	368 185-1	SANDOZ	FR
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	372 922-7	SANDOZ	FR
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	372 923-3	SANDOZ	FR
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	372 925-6	SANDOZ	FR
GLICLAZIDE SANDOZ 80 mg, comprimé sécable	NL 24744	561 184-0	SANDOZ	FR
GLICLAZIDE SANDOZ 80, tabletten met gereguleerde afgifte 80 mg	RVG 16221=56872	RVG 16221=56872	SANDOZ B.V.	NL
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578015	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578027	SANDOZ GMBH	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
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578039	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578041	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578054	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578066	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578078	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578080	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578104	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578092	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578116	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578128	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578130	SANDOZ GMBH	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
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578142	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578155	SANDOZ GMBH	IT
GLICLAZIDE SANDOZ GMBH 30MG COMPRESSE A RILASCIO MODIFICATO	PT/H/0534/001	040578167	SANDOZ GMBH	IT
Gliclazide Sandoz retard 30 mg, tabletten met gereguleerde afgifte	NL/H/1700/001	RVG 105042	SANDOZ B.V.	NL
Gliclazide Sandoz retard 60 mg, tabletten met gereguleerde afgifte	NL/H/3384/001	RVG 116611	SANDOZ B.V.	NL
Gliclazide Servier 30 mg MR Tablets	FR/H/172/01	PL 05815/0020	LES LABORATOIRES SERVIER (SURESNES)	UK
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 203-2	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 196-6	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 193-7	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 198-9	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 194-3	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 205-5	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	357 018-1	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 200-3	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	562 673-8	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	562 675-0	LES LABORATOIRES SERVIER (SURESNES)	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
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 202-6	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 197-2	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	372 262-7	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 192-0	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 204-9	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 30MG, comprimé à libération modifiée	FR/H/0172/001	354 199-5	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 60 MG, comprimé sécable à libération modifiée	FR/H/0172/002	339 467-2	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 60 MG, comprimé sécable à libération modifiée	FR/H/0172/002	339 469-5	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 60 MG, comprimé sécable à libération modifiée	FR/H/0172/002	338 973-1	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 60 MG, comprimé sécable à libération modifiée	FR/H/0172/002	339 477-8	LES LABORATOIRES SERVIER (SURESNES)	FR
GLICLAZIDE SERVIER 60 mg, comprimé sécable à libération modifiée	FR/H/0172/002	338 978-3	LES LABORATOIRES SERVIER (SURESNES)	FR
Gliclazide Servier 60 mg, tabletten met gereguleerde afgifte	FR/H/172/02/DC	RVG102836	LES LABORATOIRES SERVIER (SURESNES)	NL
Gliclazide Tablets 80mg	PL 04416/0321	PL 04416/0321	SANDOZ LTD	UK
Gliclazide Tablets 80mg BP	not available	PL 20075/0114	ACCORD HEALTHCARE LIMITED	UK
GLICLAZIDE TABLETS BP 80mg	not available	PL 0142/0400	ACTAVIS UK LTD.	UK
Gliclazide Tablets BP 80mg	not available	PL/4569/0274	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
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/01	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/02	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/03	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/04	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/05	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/06	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/07	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/08	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/09	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/10	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/11	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/12	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/13	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/14	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/15	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/16	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/17	TAD PHARMA GMBH	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
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/18	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/19	TAD PHARMA GMBH	HU
Gliclazide TAD 60 mg, Módosított hatóanyagleadású tabletta	DE/H/0894/002	OGYI-T-22410/20	TAD PHARMA GMBH	HU
GLICLAZIDE TEVA 30 mg, comprimé à libération modifiée	DE/H/0895/001	NL33444	TEVA SANTÉ	FR
Gliclazide Teva 60 mg ilgstošās darbības tabletes	DK/H/2460/001	15-0296	TEVA B.V	LV
Gliclazide Teva 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2460/001	LT/1/15/3835/001	TEVA B.V	LT
Gliclazide Teva 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2460/001	LT/1/15/3835/002	TEVA B.V	LT
Gliclazide Teva 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2460/001	LT/1/15/3835/004	TEVA B.V	LT
Gliclazide Teva 60 mg modifikuoto atpalaidavimo tabletės	DK/H/2460/001	LT/1/15/3835/003	TEVA B.V	LT
GLICLAZIDE TEVA 60 mg, comprimé sécable à libération modifiée	DE/H/3526/002	NL41933	TEVA SANTÉ	FR
Gliclazide Teva 80 mg, comprimé sécable	not available	NL24262	TEVA SANTÉ	FR
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372037	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372064	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372076	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372126	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372013	TEVA 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
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372153	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372090	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372052/M	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372140	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372102	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372025/M	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372138	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372114	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372049	TEVA ITALIA S.R.L.	IT
Gliclazide Teva Italia 30 mg compresse a rilascio modificato	DE/H/0895/001	038372088	TEVA ITALIA S.R.L.	IT
Gliclazide Teva, 60 mg toimeainet modifitseeritult vabastavad tabletid	DK/H/2460/001	892715	TEVA B.V	EE
GLICLAZIDE ZENTIVA 30 MG ILGSTOŠĀS DARBĪBAS TABLETES	DK/H/2302/001	14-0166	ZENTIVA, K.S.	LV
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 226-6	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 233-2	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 235-5	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
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 234-9	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 222-0	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 227-2	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	578 739-3	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 236-1	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 232-6	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 229-5	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 MG, COMPRIME A LIBERATION MODIFIEE	not available	497 224-3	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 mg, comprimé à libération modifiée	not available	497 223 7	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 mg, comprimé à libération modifiée	not available	497 237-8	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 mg, comprimé à libération modifiée	not available	497 230-3	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 30 mg, comprimé à libération modifiée	not available	497 228-9	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 60 MG ILGSTOŠĀS DARBĪBAS TABLETES	DK/H/2302/002/DC	14-0167	ZENTIVA, K.S.	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
Gliclazide Zentiva 60 mg tabletter med modificeret udløsning	DK/H/2302/002	52504	ZENTIVA, K.S.	DK
GLICLAZIDE ZENTIVA 60 MG, COMPRIMÉ SÉCABLE À LIBÉRATION MODIFIÉE	DK/H/2302/002	34009 300 110 2 7	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 60 MG, COMPRIMÉ SÉCABLE À LIBÉRATION MODIFIÉE	DK/H/2302/002	34009 300 109 9 0	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 60 MG, COMPRIMÉ SÉCABLE À LIBÉRATION MODIFIÉE	DK/H/2302/002	34009 300 110 1 0	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 60 MG, COMPRIMÉ SÉCABLE À LIBÉRATION MODIFIÉE	DK/H/2302/002	34009 300 110 03	SANOFI-AVENTIS FRANCE	FR
GLICLAZIDE ZENTIVA 80 mg compresse	not available	036376010	ZENTIVA ITALIA SRL	IT
Gliclazide Zentiva 80 mg, comprimé sécable	not available	367 457-8	SANOFI-AVENTIS FRANCE	FR
Gliclazide Zentiva 80 mg, comprimé sécable	not available	561 265-3	SANOFI-AVENTIS FRANCE	FR
Gliclazide Zentiva 80 mg, comprimé sécable	not available	367 458-4	SANOFI-AVENTIS FRANCE	FR
Gliclazide Zentiva Lab 30 mg compresse a rilascio modificato	DK/H/2302/001	042893065	ZENTIVA ITALIA SRL	IT
Gliclazide Zentiva Lab 30 mg compresse a rilascio modificato	DK/H/2302/001	042893038	ZENTIVA ITALIA SRL	IT
Gliclazide Zentiva Lab 30 mg compresse a rilascio modificato	DK/H/2302/001	042893014	ZENTIVA ITALIA SRL	IT
Gliclazide Zentiva Lab 30 mg compresse a rilascio modificato	DK/H/2302/001	042893053	ZENTIVA ITALIA SRL	IT
GLICLAZIDE ZENTIVA, 30 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	DK/H/2302/001	22307	ZENTIVA, K.S.	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
GLICLAZIDE ZENTIVA, 60 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	DK/H/2302/002	22308	ZENTIVA, K.S.	PL
Gliclazide Zentiva30 mg tabletter med modificeret udløsning	DK/H/2302/001	52502	ZENTIVA, K.S.	DK
Gliclazide/Mylan 30 mg δισκία ελεγχόμενης αποδέσμευσης	DE/H/0897/001	16579/15-02-2016	GENERICS PHARMA HELLAS LTD	GR
Gliclazid-ratiopharm 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0896/001	1-27539	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Gliclazid-ratiopharm 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0896/001	67674.00.00	RATIOPHARM GMBH	DE
Gliclazid-TEVA 30 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/0895/001	67673.00.00	TEVA GMBH	DE
Gliglical, tabletter med modificeret udløsning	DK/H/2460/001	55841	TEVA B.V	DK
GLIKA 30 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/11-01/116	BELUPO D.D.	HR
Gliklazid Actavis 30 mg tablety s řízeným uvolňováním	DK/H/2376/001	18/250/15-C	ACTAVIS GROUP PTC EHF.	CZ
Gliklazid Actavis 60 mg tablety s řízeným uvolňováním	DK/H/2376/002	18/251/15-C	ACTAVIS GROUP PTC EHF.	CZ
Gliklazid Krka 60 mg tablete s prilagođenim oslobađanjem	not available	UP/I-530-09/12-01/02	KRKA-FARMA D.O.O.	HR
GLIKLAZID LUPIN 60 mg retard tabletta	PT/H/1272/001/DC	OGYI-T-22888/01-02	Lupin (Europe) Ltd.	HU
Gliklazid Lupin 60 mg tablety s predlženým uvoľňovaním	PT/H/1272/001	18/0316/15-S	LUPIN (EUROPE) LIMITED	SK
Gliklazid Lupin, 60 mg, tabletki o zmodyfikowanym uwalnianiu	PT/H/1272/001/DC	22928	LUPIN (EUROPE) LIMITED	PL
Gliklazid Sandoz 30 mg tablete s prilagođenim oslobađanjem	NL/H/3108/001	HR-H-404023017	SANDOZ D.O.O.	HR
Gliklazid Sandoz 60 mg	NL/H/3384/001	18/093/16-C	SANDOZ S.R.O.	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
Gliklazid Sandoz 60 mg tablete s prilagođenim oslobađanjem	NL/H/3384/001	HR-H-898920477	SANDOZ D.O.O.	HR
Gliklazid Sandoz 60 mg tablety s riadeným uvoľňovaním	NL/H/3384/001	18/0057/16-S	SANDOZ PHARMACEUTICALS D.D.	SK
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/007	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/006	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/010	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/004	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/018	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/003	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/015	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/013	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/001	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/008	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/005	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/002	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/016	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/017	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/014	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
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/009	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/012	STADA ARZNEIMITTEL AG	SI
Gliklazid STADA 60 mg tablete s prirejenim sproščanjem	DK/H/2301/001	H/14/1904/011	STADA ARZNEIMITTEL AG	SI
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/001	LUPIN (EUROPE) LIMITED	LT
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/002	LUPIN (EUROPE) LIMITED	LT
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/003	LUPIN (EUROPE) LIMITED	LT
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/004	LUPIN (EUROPE) LIMITED	LT
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/005	LUPIN (EUROPE) LIMITED	LT
Gliklazidas Lupin 60 mg pailginto atpalaidavimo tabletės	PT/H/1272/001	LT/1/15/3790/006	LUPIN (EUROPE) LIMITED	LT
Glirike 60 mg Tabletten mit veränderter Wirkstofffreisetzung	DE/H/3526/001	81782.00.00	KRKA, D.D., NOVO MESTO	DE
Glizorem		AA084/08901	REMEDICA LTD	MT
Glizorem 80 mg Δισκία	not available	20347	REMEDICA LTD	CY
Gluctam 60 mg módosított hatóanyagleadású tabletta	FR/H/0172/002/DC	OGYI-T-10108/03-04	EGIS PHARMACEUTICALS PLC	HU
Gluctam 80 mg tabletta	not available	OGYI-T-6191/01	EGIS PHARMACEUTICALS PLC	HU
Gluctam MR 60 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/172/02/DC	1-30194	SERVIER AUSTRIA GMBH	AT
GLUCTAM MR 60MG	FR/H/172/02/DC	18/0764/09-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Glyakt 30 mg tablete s prilagođenim oslobađanjem	DK/H/2376/001	HR-H-398480928	ACTAVIS GROUP PTC EHF.	HR
Glyakt 60 mg tablete s prilagođenim oslobađanjem	DK/H/2376/002	HR-H-959639573	ACTAVIS GROUP PTC EHF.	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
Glyclada 30 mg Tablety s řízeným uvolňováním	DE/H/0892/001	18/105/08-C	KRKA, D.D., NOVO MESTO	CZ
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/01	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/02	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/03	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/04	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/05	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/06	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/07	KRKA, D.D., NOVO MESTO	RO
Glyclada 30 mg, comprimate cu eliberare prelungită	not available	1139/2008/08	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/01	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/02	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/03	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/04	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/05	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/06	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/07	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/08	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
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/09	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/10	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/11	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/12	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/13	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/14	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/15	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/16	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/17	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/18	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/19	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg comprimate cu eliberare modificată	DE/H/0892/002	5467/2013/20	KRKA, D.D., NOVO MESTO	RO
Glyclada 60 mg por.tbl.ret.	DE/H/0892/002	18/321/13-C	KRKA, D.D., NOVO MESTO	CZ
Glyclada 90 mg tablety s řízeným uvolňováním	DE/H/0892/003	18/020/16-C	KRKA, D.D., NOVO MESTO	CZ
Glydium 30 mg Tabletten mit veränderter Wirkstofffreisetzung	FR/H/336/01/DC	1-27729	LES LABORATOIRES SERVIER (SURESNES)	AT
GLYDIUM 30 mg, comprimé à libération modifiée	FR/H/0336/001	390 517-3	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 511-5	LES LABORATOIRES SERVIER (SURESNES)	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
GLYDIUM 30 mg, comprimé à libération modifiée	FR/H/336/01/DC	390 514-4	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 520-4	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	574 169-8	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 516-7	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	574 170-6	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 510-9	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 507-8	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 509-0	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 519-6	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 513-8	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30 MG, comprimé à libération modifiée	FR/H/0336/001	390 512-1	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30MG, comprimé à libération modifiée	FR/H/336/01/DC	390 508-4	LES LABORATOIRES SERVIER (SURESNES)	FR
GLYDIUM 30MG, comprimé à libération modifiée	FR/H/336/01/DC	390 515-0	LES LABORATOIRES SERVIER (SURESNES)	FR
Gulrike 60 mg Comprimidos de liberación modificada EFG	DE/H/3526/001	77494	KRKA, D.D., NOVO MESTO	ES
Laaglyda MR 60 mg modified-release tablets	DE/H/0894/002	PL 01656/0182	KRKA, D.D., NOVO MESTO	UK
Medoclazide	not available	MA032/01901	MEDOCHEMIE LTD.	MT
Medoclazide 80 mg tablets	not available	9819	MEDOCHEMIE LTD.	CY
Nazdol MR 30 mg modified-release tablets	DE/H/0892/001	PL 01656/0033	KRKA, D.D., NOVO MESTO	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nazdol MR 60 mg modified-release tablets	DE/H/0892/002	PL 01656/0181	KRKA, D.D., NOVO MESTO	UK
Normodiab 80 mg tablets	IS/H/0145/001	20030689	ACTAVIS GROUP PTC EHF.	BG
Normodiab 80 mg tablets	IS/H/0145/001	IS/1/10/068/01	ACTAVIS GROUP PTC EHF.	IS
Oziclide MR, 60 mg, tabletki o zmodyfikowanym uwalnianiu	UK/H/5466/001	22511	RANBAXY POLAND SP. ZO.O	PL
Salson, 60 mg, tabletki o zmodyfikowanym uwalnianiu	NL/H/3384	23060	SANDOZ GMBH	PL
Symazide MR 60, 60 mg, tabletki o zmodyfikowanym uwalnianiu	PL/H/0436/001	22936	SYMPHAR SP. Z O.O.	PL
Symazide MR, 30 mg, tabletka o zmodyfikowanym uwalnianiu	PL/H/0266/001	17622	SYMPHAR SP. Z O.O.	PL
UNI DIAMICRON 30 MG comprimés à libération modifiée	FR/H/0171/001	BE220315	SERVIER BENELUX S.A./N.V.	BE
UNI DIAMICRON 60 mg Tablette mit veränderter Wirkstofffreisetzung	FR/H/171/02/DC	BE354137	SERVIER BENELUX S.A./N.V.	BE
Uni Gliclazide EG 30 mg tabletten met gereguleerde afgifte	DK/H/2270/001	BE446462	EUROGENERICS SA	BE
Uni Gliclazide EG 30 mg tabletten met gereguleerde afgifte	DK/H/2270/001	BE446453	EUROGENERICS SA	BE
Uni Gliclazide EG 60 mg tabletten met gereguleerde afgifte	DK/H/2301/001	BE460240	EUROGENERICS SA	BE
Uni Gliclazide EG 60 mg tabletten met gereguleerde afgifte	DK/H/2301/001	BE460257	EUROGENERICS SA	BE
Uni Gliclazide EG 60mg comprimés à libération modifiée	DK/H/2301/001	2015120261	EUROGENERICS SA	LU
Vamju 30 mg modified-release tablets	DK/H/2381/001	PL 12762/0506	MERCURY PHARMACEUTICALS LTD.	UK
Vamju 60 mg modified-release tablets	DK/H/2381/002	PL 12762/0507	MERCURY PHARMACEUTICALS LTD.	UK
Vitile MR 30 mg modified-release tablets	DK/H/2376/001	PA1380/166/1	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
Ziclaseg 30mg prolonged-release tablets	PT/H/0534/001	PL 35507/0084	LUPIN (EUROPE) LIMITED	UK
Zicron 40mg Tablets	not available	PL 17907/0067	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Zicron PR 30 mg prolonged-release tablets	UK/H/5530/01/DC	MA737/01501	BRISTOL LABORATORIES LTD (BERKHAMSTED)	MT
Zicron PR 30 mg prolonged-release tablets	UK/H/5530/01/DC	PL 17907/0398	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
ZODEDIAB 30 mg comprimate cu eliberare modificată	DK/H/2300/001	6784/2014/01-08	ALVOGEN IPCO S.AR.L	RO
ZODEDIAB 60 mg comprimate cu eliberare modificată	DK/H/2300/002	6785/2014/01-08	ALVOGEN IPCO S.AR.L	RO
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/01	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/02	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/03	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/04	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/05	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/06	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/07	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg módosított hatóanyagleadású tabletta	DK/H/2300/001	OGYI-T-22698/08	ALVOGEN IPCO S.AR.L	HU
Zodediab 30 mg töflur með breyttan losunarhraða	DK/H/2300/001	IS/1/14/063/01	ALVOGEN IPCO S.AR.L	IS
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/09	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/10	ALVOGEN IPCO S.AR.L	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
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/11	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/12	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/16	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/15	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/14	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg módosított hatóanyagleadású tabletta	DK/H/2300/002	OGYI-T-22698/13	ALVOGEN IPCO S.AR.L	HU
Zodediab 60 mg töflur með breyttan losunarhraða	DK/H/2300/002	IS/1/14/063/02	ALVOGEN IPCO S.AR.L	IS
Zodediab MR	DK/H/2300/002	52498	ALVOGEN IPCO S.AR.L	DK
Zodediab MR	DK/H/2300/001	52497	ALVOGEN IPCO S.AR.L	DK
Zycron MR 30 mg modified-release tablets	DK/H/2270/001	PA 126/248/1	CLONMEL HEALTHCARE LTD.	IE
ГЛИКЛАДА 30 mg таблетки с удължено освобождаване	not available	20100262	KRKA, D.D., NOVO MESTO	BG
ГЛИКЛАДА 60 mg таблетки с изменено освобождаване	DE/H/0892/002	20130083	KRKA, D.D., NOVO MESTO	BG
Гликлада 90 mg таблетки с изменено освобождаване	DE/H/0892/003	20160071	KRKA, D.D., NOVO MESTO	BG
Гликлазид ГАММА 30 MR 30 mg таблетки с изменено освобождаване	NL/H/1711/001	20100697	WÖRWAG PHARMA GMBH & CO. KG	BG
Гликлазид Дженерикс 30 mg таблетки с изменено освобождаване	DE/H/0893/001	20090308	GENERICS [UK] LIMITED	BG
Гликлазид Зентива 60 mg таблетки с изменено освобождаване	DK/H/2302/002	20140311	ZENTIVA, K.S.	BG

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Гликлазид Лупин 60 mg таблетки с удължено освобождаване	PT/H/1272/001	20150831	LUPIN (EUROPE) LIMITED	BG
Диаб 80 mg таблетки	not available	II-15101	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Диаб MP 30mg таблетки с изменено освобождаване	not available	II-20677	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
ДИАПРЕЛ MR 60 mg таблетки с изменено освобождаване	FR/H/171/02/DC	20100003	LES LABORATOIRES SERVIER (SURESNES)	BG
Дигликал ER 60 mg таблетки с изменено освобождаване	DK/H/2460/001	II-31635 РЕГ. №: 20150399	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Екозид SR 30 mg таблетки с удължено освобождаване	not available	20120076	ECOPHARM GROUP	BG
Зодедиаб MR 30 mg таблетки с изменено освобождаване	DK/H/2300/001	II-26942	ALVOGEN IPCO S.AR.L	BG
Зодедиаб MR 60 mg таблетки с изменено освобождаване	DK/H/2300/002	II-26943	ALVOGEN IPCO S.AR.L	BG
МАДРАС MR 30 mg таблетки с изменено освобождаване	DK/H/2270/001	20140046	STADA ARZNEIMITTEL AG	BG
Мадрас MR 60 mg таблетки с изменено освобождаване	DK/H/2301/001	20140256	STADA ARZNEIMITTEL AG	BG
Нормодиаб MR 30 mg таблетки с изменено освобождаване	DK/H/2376/001	20150142	ACTAVIS GROUP PTC EHF.	BG
Нормодиаб MR 60 mg таблетки с изменено освобождаване	DK/H/2376/002	20150143	ACTAVIS GROUP PTC EHF.	BG