29 November 2018
EMA/846372/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: metformin

Procedure no.: PSUSA/00002001/201804

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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
Dianben 850 mg comprimidos recubiertos con película	FR/H/0181/002	55.211	MERCK SANTÉ S.A.S.	ES
Glucophage 1000 mg - Filmtabletten	FR/H/0181/001	1-24142	MERCK GESELLSCHAFT MBH	AT
GLUCOPHAGE 1000 mg apvalkotās tabletes	FR/H/0181/001	08-0335	MERCK SANTÉ S.A.S.	LV
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758044	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758057	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758069	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758071	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758083	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758095	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758107	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758119	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758032	BRUNO FARMACEUTICI	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
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758044	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758057	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758069	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758071	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758083	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758095	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758107	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758119	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg compresse rivestite con film	FR/H/0181/001	017758032	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/09	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/14	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/17	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/13	MERCK SANTÉ S.A.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
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/19	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/04	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/08	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/02	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/12	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/07	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/11	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/15	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/18	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/01	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/06	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/03	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/16	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/10	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg comprimate filmate	FR/H/0181/001	7538/2015/05	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 1000 mg film-coated tablets	FR/H/0181/001	PA 654/19/3	MERCK SERONO LTD.	IE
GLUCOPHAGE 1000 mg filmdragerad tablett	FR/H/0181/001	17143	MERCK SANTÉ S.A.S.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glucophage 1000 mg filmdrasjerte tabletter	FR/H/0181/001	01-1628	MERCK SANTÉ S.A.S.	NO
Glucophage 1000 mg filmom obalené tablety	FR/H/0181/001	18/0589/08-S	MERCK SANTÉ S.A.S.	SK
Glucophage 1000 mg filmom obložene tablete	not available	HR-H-841365882	MERCK D.O.O	HR
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/006	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/007	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/008	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/002	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/003	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/004	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/001	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/009	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/005	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/015	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/016	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/017	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/010	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/011	MERCK 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
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/012	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/013	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/018	MERCK D.O.O.	SI
GLUCOPHAGE 1000 mg filmsko obložene tablete	FR/H/0181/001	H/98/00703/014	MERCK D.O.O.	SI
Glucophage 1000 mg Filmtabletten	FR/H/0181/001	51643.00.00	MERCK SERONO GMBH	DE
GLUCOPHAGE 1000 mg filmuhúðaðar töflur	FR/H/0181/001	IS/1/01/019/01	MERCK SANTÉ S.A.S.	IS
GLUCOPHAGE 1000 mg potahované tablety	FR/H/0181/001	18/155/02-C	MERCK SANTÉ S.A.S.	CZ
GLUCOPHAGE 1000 mg tabletti, kalvopäällysteinen	FR/H/0181/001	16414	MERCK SANTÉ S.A.S.	FI
GLUCOPHAGE 1000 mg, 1000 mg, tabletki powlekane	FR/H/0181/001	15716	MERCK SANTÉ S.A.S.	PL
GLUCOPHAGE 1000 mg, comprimé pelliculé sécable	FR/H/0181/001	356 020-2	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 1000 mg, comprimé pelliculé sécable	FR/H/0181/001	356 023-1	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 1000 mg, comprimé pelliculé sécable	FR/H/0181/001	563 259-0	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 1000 mg, comprimé pelliculé sécable	FR/H/0181/001	356 017-1	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 1000 mg, επικαλυμμένο με λεπτό υμένιο δισκίο	FR/H/0181/001	20469	MERCK A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GLUCOPHAGE 1000 mg, Επικαλυμμένο με λεπτό υμένιο δισκίο	FR/H/0181/001	27102/08/12-03-2009	MERCK A.E.	GR
Glucophage 500 mg - Filmtabletten	FR/H/0181/003	11.834	MERCK GESELLSCHAFT MBH	AT
GLUCOPHAGE 500 mg apvalkotās tabletes	FR/H/0181/003	09-0344	MERCK SANTÉ S.A.S.	LV
GLUCOPHAGE 500 mg compresse rivestite con film	FR/H/0181/003	017758018	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 500 mg compresse rivestite con film	FR/H/0181/003	017758018	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/10	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/14	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/15	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/03	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/11	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/12	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/02	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/05	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/28	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/24	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/20	MERCK SANTÉ S.A.S.	RO

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GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/01	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/26	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/23	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/07	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/29	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/21	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/27	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/25	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/08	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/18	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/06	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/16	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/22	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/17	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/04	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/19	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/09	MERCK SANTÉ S.A.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
GLUCOPHAGE 500 mg comprimate filmate	FR/H/0181/003	7536/2015/13	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 500 mg comprimés pelliculés	FR/H/0181/003	BE058414	MERCK N.V./S.A.	BE
GLUCOPHAGE 500 mg comprimés pelliculés	FR/H/0181/003	BE341451	MERCK N.V./S.A.	BE
GLUCOPHAGE 500 mg comprimés pelliculés	FR/H/0181/003	194484	MERCK N.V./S.A.	LU
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487591	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487690	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4485892	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487294	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487492	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487898	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4488698	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487096	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487799	MERCK, 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
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486791	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486296	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	8050815	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487195	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486593	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486197	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486395	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4485991	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	8050807	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487393	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4488391	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos	FR/H/0181/003	4488292	MERCK, 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
por película				
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486692	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4488599	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4488193	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4486890	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4488094	MERCK, S.A.	PT
GLUCOPHAGE 500 mg comprimidos revestidos por película	FR/H/0181/003	4487997	MERCK, S.A.	PT
GLUCOPHAGE 500 mg film-coated tablets	FR/H/0181/003	PA 654/19/1	MERCK SERONO LTD.	IE
GLUCOPHAGE 500 mg film-coated tablets	FR/H/0181/003	PL 11648/0085	MERCK SERONO LTD.	UK
GLUCOPHAGE 500 mg filmdragerad tablett	FR/H/0181/003	7690	MERCK SANTÉ S.A.S.	SE
Glucophage 500 mg filmdrasjerte tabletter	FR/H/0181/003	5204	MERCK SANTÉ S.A.S.	NO
Glucophage 500 mg filmom obalené tablety	FR/H/0181/003	18/0561/09-S	MERCK SANTÉ S.A.S.	SK
Glucophage 500 mg filmom obložene tablete	not available	HR-H-839391381	MERCK D.O.O	HR
GLUCOPHAGE 500 mg filmomhulde tabletten	FR/H/0181/003	BE341451	MERCK N.V./S.A.	BE
GLUCOPHAGE 500 mg filmomhulde tabletten	FR/H/0181/003	BE058414	MERCK N.V./S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/019	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/034	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/030	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/021	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/031	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/022	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/023	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/024	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/032	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/025	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/026	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/027	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/033	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/028	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/020	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/029	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/040	MERCK 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
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/046	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/041	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/035	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/036	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/042	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/037	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/043	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/038	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/044	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/039	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg filmsko obložene tablete	FR/H/0181/003	H/98/00703/045	MERCK D.O.O.	SI
GLUCOPHAGE 500 mg Filmtabletten	FR/H/0181/003	BE058414	MERCK N.V./S.A.	BE
GLUCOPHAGE 500 mg Filmtabletten	FR/H/0181/003	BE341451	MERCK N.V./S.A.	BE
Glucophage 500 mg Filmtabletten	FR/H/0181/003	6207899.00.00	MERCK SERONO GMBH	DE
GLUCOPHAGE 500 mg filmuhúðaðar töflur	FR/H/0181/003	691229	MERCK SANTÉ S.A.S.	IS
GLUCOPHAGE 500 mg potahované tablety	FR/H/0181/003	18/824/96-C	MERCK SANTÉ S.A.S.	CZ
GLUCOPHAGE 500 mg tabletti,	FR/H/0181/003	4566	MERCK SANTÉ S.A.S.	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kalvopäällysteinen				
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	551 036-1	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	372 048-5	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	558 344-3	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	337 259-3	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	304 478-8	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, comprimé pelliculé	FR/H/0181/003	352 816-7	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 500 mg, tabletki powlekane	FR/H/0181/003	16876	MERCK SANTÉ S.A.S.	PL
GLUCOPHAGE 500 mg, επικαλυμμένα με λεπτό υμένιο δισκία	FR/H/0181/003	20697	MERCK A.E.	CY
GLUCOPHAGE 500 mg, επικαλυμμένα με λεπτό υμένιο δισκία	FR/H/0181/003	18517/18-03-2010	MERCK A.E.	GR
Glucophage 850 mg – Filmtabletten	FR/H/0181/002	14.190	MERCK GESELLSCHAFT MBH	AT
GLUCOPHAGE 850 mg compresse rivestite con film	FR/H/0181/002	017758020	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 850 mg compresse rivestite con film	FR/H/0181/002	017758020	BRUNO FARMACEUTICI	IT
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/24	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/17	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/13	MERCK SANTÉ S.A.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
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/14	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/08	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/07	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/09	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/21	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/16	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/05	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/23	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/15	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/19	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/18	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/03	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/22	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/01	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/11	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/20	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/02	MERCK SANTÉ S.A.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
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/04	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/12	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/06	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimate filmate	FR/H/0181/002	7537/2015/10	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE 850 mg comprimés pelliculés	FR/H/0181/002	BE058466	MERCK N.V./S.A.	BE
GLUCOPHAGE 850 mg comprimés pelliculés	FR/H/0181/002	BE341467	MERCK N.V./S.A.	BE
GLUCOPHAGE 850 mg comprimés pelliculés	FR/H/0181/002	286832	MERCK N.V./S.A.	LU
GLUCOPHAGE 850 mg comprimés pelliculés	FR/H/0181/002	194498	MERCK N.V./S.A.	LU
GLUCOPHAGE 850 mg film-coated tablets	FR/H/0181/002	PA 654/19/2	MERCK SERONO LTD.	IE
GLUCOPHAGE 850 mg film-coated tablets	FR/H/0181/002	PL 11648/0086	MERCK SERONO LTD.	UK
GLUCOPHAGE 850 mg filmdragerad tablett	FR/H/0181/002	10799	MERCK SANTÉ S.A.S.	SE
Glucophage 850 mg filmdrasjerte tabletter	FR/H/0181/002	97-375	MERCK SANTÉ S.A.S.	NO
Glucophage 850 mg filmom obalené tablety	FR/H/0181/002	18/0562/09-S	MERCK SANTÉ S.A.S.	SK
Glucophage 850 mg filmom obložene tablete	not available	HR-H-213830516	MERCK D.O.O	HR
GLUCOPHAGE 850 mg filmomhulde tabletten	FR/H/0181/002	BE341467	MERCK N.V./S.A.	BE
GLUCOPHAGE 850 mg filmomhulde tabletten	FR/H/0181/002	BE058466	MERCK N.V./S.A.	BE
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/050	MERCK 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
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/064	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/047	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/061	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/051	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/052	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/053	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/062	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/054	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/055	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/056	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/057	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/058	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/063	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/048	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/059	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/049	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/060	MERCK 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
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/065	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/067	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/068	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/066	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg filmsko obložene tablete	FR/H/0181/002	H/98/00703/069	MERCK D.O.O.	SI
GLUCOPHAGE 850 mg Filmtabletten	FR/H/0181/002	BE058466	MERCK N.V./S.A.	BE
GLUCOPHAGE 850 mg Filmtabletten	FR/H/0181/002	BE341467	MERCK N.V./S.A.	BE
Glucophage 850 mg Filmtabletten	FR/H/0181/002	6373818.00.00	MERCK SERONO GMBH	DE
GLUCOPHAGE 850 mg filmuhúðaðar töflur	FR/H/0181/002	853604	MERCK SANTÉ S.A.S.	IS
GLUCOPHAGE 850 mg potahované tablety	FR/H/0181/002	18/825/96-C	MERCK SANTÉ S.A.S.	CZ
GLUCOPHAGE 850 mg επικαλυμμένα με λεπτό υμένιο δισκία	FR/H/0181/002	20698	MERCK A.E.	CY
GLUCOPHAGE 850 mg επικαλυμμένα με λεπτό υμένιο δισκία	FR/H/0181/002	34521/02-06-2008	MERCK A.E.	GR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 328 730 9 8	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 372 246 1 1	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 559 233 0 5	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 329 827 6 9	MERCK SANTÉ S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 337 260 1 0	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 558 346 6 3	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, comprimé pelliculé	FR/H/0181/002	34009 304 480 2 1	MERCK SANTÉ S.A.S.	FR
GLUCOPHAGE 850 mg, tabletki powlekane	FR/H/0181/002	16877	MERCK SANTÉ S.A.S.	PL
Glucophage SR 1000 mg prolonged release tablet	not available	PL 11648/0067	MERCK SERONO LTD.	UK
Glucophage SR 500mg prolonged release tablet	not available	PL 11648/0054	MERCK SERONO LTD.	UK
Glucophage SR 750 mg prolonged release tablet	not available	PL 11648/0066	MERCK SERONO LTD.	UK
GLUCOPHAGE XR 1000 mg comprimate cu eliberare prelungită	not available	10261/2017/01	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 1000 mg comprimate cu eliberare prelungită	not available	10261/2017/02	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 1000 mg comprimate cu eliberare prelungită	not available	10261/2017/03	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 1000 mg comprimate cu eliberare prelungită	not available	10261/2017/04	MERCK SANTÉ S.A.S.	RO
Glucophage XR 1000 mg ilgstošās darbības tabletes	not available	10-0372	MERCK SANTÉ S.A.S.	LV
Glucophage XR 1000 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-01/95	MERCK D.O.O	HR
Glucophage XR 1000 mg tablety s predlženým uvolnovaním	not available	18/0150/10-S	MERCK SANTÉ S.A.S.	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
GLUCOPHAGE XR 1000 mg tablety s prodlouženým uvolňováním	not available	18/222/11-C	MERCK SANTÉ S.A.S.	CZ
GLUCOPHAGE XR 500 mg comprimate cu eliberare prelungită	not available	7236/2014/02	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 500 mg comprimate cu eliberare prelungită	not available	7236/2014/01	MERCK SANTÉ S.A.S.	RO
Glucophage XR 500 mg ilgstošās darbības tabletes	not available	04-0289	MERCK SANTÉ S.A.S.	LV
Glucophage XR 500 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-01/93	MERCK D.O.O	HR
GLUCOPHAGE XR 500 mg tablety s prodlouženým uvolňováním	not available	18/166/04-C	MERCK SANTÉ S.A.S.	CZ
GLUCOPHAGE XR 750 mg comprimate cu eliberare prelungita	not available	10260/2017/01	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 750 mg comprimate cu eliberare prelungita	not available	10260/2017/02	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 750 mg comprimate cu eliberare prelungita	not available	10260/2017/03	MERCK SANTÉ S.A.S.	RO
GLUCOPHAGE XR 750 mg comprimate cu eliberare prelungita	not available	10260/2017/04	MERCK SANTÉ S.A.S.	RO
Glucophage XR 750 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-01/94	MERCK D.O.O	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glucophage XR 750 mg tablety s predĺženým uvoľňovaním	not available	18/0515/06-S	MERCK SANTÉ S.A.S.	SK
GLUCOPHAGE XR 750 mg tablety s prodlouženým uvolňováním	not available	18/221/11-C	MERCK SANTÉ S.A.S.	CZ
Glucophage XR tablety s predĺženým uvoľňovaním	not available	18/0290/04-S	MERCK SANTÉ S.A.S.	SK
Glucophage XR, 1000 mg, tabletki o przedłużonym uwalnianiu	not available	18106	MERCK SANTÉ S.A.S.	PL
Glucophage XR, 500 mg, tabletki o przedłużonym uwalnianiu	not available	12244	MERCK SANTÉ S.A.S.	PL
Glucophage XR, 750 mg, tabletki o przedłużonym uwalnianiu	not available	15192	MERCK SANTÉ S.A.S.	PL
GLUCOSTOP®, αναβράζον δισκίο 1000 mg/tab	not available	71732/13/04.02.2014	IOULIA AND IRENE TSETI PHARMACEUTICAL LABORATORIES S.A.	GR
LYOMET, depottabletter	DK/H/2272/001/DC	51937	GENERIC PARTNERS UK LTD	DK
LYOMET, depottabletter	DK/H/2272/002/DC	51938	GENERIC PARTNERS UK LTD	DK
LYOMET, depottabletter	DK/H/2272/003/DC	51939	GENERIC PARTNERS UK LTD	DK
Merckformin 1000 mg filmtabletta	FR/H/0181/001	OGYI-T-5157/10	MERCK KFT.	HU
Merckformin 1000 mg filmtabletta	FR/H/0181/001	OGYI-T-5157/11	MERCK KFT.	HU
Merckformin 1000 mg filmtabletta	FR/H/0181/001	OGYI-T-5157/12	MERCK KFT.	HU
Merckformin 500 mg filmtabletta	FR/H/0181/003	OGYI-T-5157/07	MERCK KFT.	HU
Merckformin 850 mg filmtabletta	FR/H/0181/002	OGYI-T-5157/02	MERCK KFT.	HU
Merckformin 850 mg	FR/H/0181/002	OGYI-T-5157/17	MERCK 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
filmtabletta				
Merckformin 850 mg filmtabletta	FR/H/0181/002	OGYI-T-5157/03	MERCK KFT.	HU
Merckformin 850 mg filmtabletta	FR/H/0181/002	OGYI-T-5157/01	MERCK KFT.	HU
Merckformin 850 mg filmtabletta	FR/H/0181/002	OGYI-T-5157/18	MERCK KFT.	HU
Merckformin XR 1000 mg retard tabletta	not available	OGYI-T-5157/16	MERCK KFT.	HU
Merckformin XR 1000 mg retard tabletta	not available	OGYI-T-5157/15	MERCK KFT.	HU
Merckformin XR 500 mg retard tabletta	not available	OGYI-T-5157/13	MERCK KFT.	HU
Merckformin XR 500 mg retard tabletta	not available	OGYI-T-5157/14	MERCK KFT.	HU
Merckformin XR 750 mg retard tabletta	not available	OGYI-T-5157/04	MERCK KFT.	HU
Merckformin XR 750 mg retard tabletta	not available	OGYI-T-5157/05	MERCK KFT.	HU
Merckformin XR 750 mg retard tabletta	not available	OGYI-T-5157/06	MERCK KFT.	HU
MetfoLiquid GeriaSan 1000 mg/5 ml Lösung zum Einnehmen	DE/H/5203/001	95448.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
METFORAL 500 mg compresse rivestite con film	not available	019449014	LABORATORI GUIDOTTI S.P.A.	IT
METFORAL 850 mg compresse rivestite con film	not available	019449038	LABORATORI GUIDOTTI S.P.A.	IT
Metformax 850 mg comprimés enrobés	not available	BE195264	MENARINI BENELUX N.V./S.A.	BE
Metformax 850 mg comprimés enrobés	not available	2004068285	MENARINI BENELUX N.V./S.A.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Metformax 850 mg omhulde tabletten	not available	BE195264	MENARINI BENELUX N.V./S.A.	BE
Metformax 850 mg überzogene Tabletten	not available	BE195264	MENARINI BENELUX N.V./S.A.	BE
Metformin 100mg/ml Oral Solution	not available	PL 20046/0255	FOCUS PHARMACEUTICALS LIMITED	UK
Metformin 500 mg Film-Coated Tablets	not available	PL 29831/0133	WOCKHARDT UK LTD	UK
Metformin 500 mg film-coated tablets	not available	PL 21880/0199	MEDREICH PLC	UK
Metformin 500 mg film-coated tablets BP	not available	PL 46447/0011	JCSH PHARMA LIMITED	UK
Metformin 500 mg Tablets BP	not available	PL 39484/0032	FOURRTS (UK) PHARMACARE LIMITED	UK
Metformin 500mg tablets	not available	PL 36722/0030	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Metformin 500mg Tablets	not available	PL 20395/0027	RELON CHEM LIMITED	UK
METFORMIN 500mg TABLETS BP	not available	PL 0142/0505	ACTAVIS UK LTD.	UK
Metformin 500mg Tablets BP	not available	PL 17907/0178	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Metformin 850 mg film-coated tablets	not available	PL 21880/0200	MEDREICH PLC	UK
Metformin 850 mg film-coated tablets BP	not available	PL 46447/0012	JCSH PHARMA LIMITED	UK
Metformin 850 mg Tablets	not available	PL 20395/0028	RELON CHEM LIMITED	UK
Metformin 850 mg Tablets BP	not available	PL 39484/0033	FOURRTS (UK) PHARMACARE LIMITED	UK
Metformin 850mg Film-Coated Tablets	not available	PL 29831/0134	WOCKHARDT UK LTD	UK
Metformin 850mg tablets	not available	PL 36722/0031	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
METFORMIN 850mg TABLETS BP	not available	PL 0142/0506	ACTAVIS UK LTD.	UK
Metformin 850mg Tablets BP	not available	PL 17907/0179	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Metformin Actavis 850 mg filmuhúðaðar töflur	FR/H/0387/002	IS/1/09/055/02	ACTAVIS GROUP PTC EHF.	IS
Metformin Arcana 850 mg Filmtabletten	not available	1-19303	ARCANA ARZNEIMITTEL GMBH	AT
Metformin Aurobindo 850 mg apvalkotās tabletes	NL/H/1459/002	09-0031	AUROBINDO PHARMA (MALTA) LIMITED	LV
Metformin Hydrochloride 500 mg film-coated tablets	not available	PL 42289/0003	WAVE PHARMA LIMITED	UK
Metformin Hydrochloride 850 mg film-coated tablets	not available	PL 42289/0004	WAVE PHARMA LIMITED	UK
Metformin Sandoz 1000 mg filmtabletta	NL/H/3301/001	OGYI-T-7107/05	SANDOZ HUNGÁRIA KFT	HU
Metformin Sandoz 1000 mg filmtabletta	NL/H/3301/001	OGYI-T-7107/06	SANDOZ HUNGÁRIA KFT	HU
Metformin Tablets 500 mg	not available	PL 20416/0323	CRESCENT PHARMA LIMITED	UK
Metformin Tablets 500mg	not available	PL 04416/0300	SANDOZ LTD	UK
Metformin Tablets 850 mg	not available	PL 20416/0324	CRESCENT PHARMA LIMITED	UK
Metformin Tablets 850mg	not available	PL 04416/0301	SANDOZ LTD	UK
Metformin Zentiva 1 000 mg potahované tablety	not available	18/177/06-C	ZENTIVA, K.S.	CZ
Metformin Zentiva 1 000 mg potahované tablety	not available	18/177/06-C	ZENTIVA, K.S.	CZ
Metformin Zentiva 1 000 mg potahované tablety	not available	18/177/06-C	ZENTIVA, K.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Metformina Aurobindo 1000 mg compresse rivestite con film	IT/H/0464/003	040592040	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Metformina Aurobindo 1000 mg compresse rivestite con film	IT/H/0464/003	040592089	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066372	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066384	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066396	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066408	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066410	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066422	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066434	DOC GENERICI S.R.L.	IT
METFORMINA DOC	NL/H/1459/003	039066446	DOC GENERICI 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
Generici 1000 mg compresse rivestite con film				
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066459	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066461	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066473	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066840	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066853	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066865	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066877	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066889	DOC GENERICI 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
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066891	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066903	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066915	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066927	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066939	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	039066941	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	044531046	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	044531059	DOC GENERICI S.R.L.	IT
METFORMINA DOC Generici 1000 mg compresse rivestite con	NL/H/1459/003	044531061	DOC GENERICI 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
film				
METFORMINA DOC Generici 1000 mg compresse rivestite con film	NL/H/1459/003	044531097	DOC GENERICI S.R.L.	IT
Metformine HCl Disper Mylan 850 mg, dispergeerbare tabletten	FR/H/0362/002	RVG 103134	MYLAN B.V.	NL
METFORMINE MYLAN 850 mg, comprimé dispersible	FR/H/0362/002	NL 33877	MYLAN S.A.S	FR
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3638889	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639085	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4022984	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639184	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4022786	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3638988	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639689	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639382	MERCK, 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
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639481	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4022885	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023586	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023081	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023388	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023289	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023180	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	4023487	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639283	MERCK, S.A.	PT
RISIDON 1000 mg comprimidos revestidos por película	FR/H/0181/001	3639580	MERCK, S.A.	PT
RISIDON 850 mg comprimidos revestidos por película	FR/H/0181/002	8248930	MERCK, S.A.	PT
RISIDON 850 mg comprimidos revestidos	FR/H/0181/002	8248922	MERCK, 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
por película				
Siofor 500 mg filmsko obložene tablete	not available	H/02/01427/002	BERLIN-CHEMIE AG	SI
Siofor 500 mg filmsko obložene tablete	not available	H/02/01427/001	BERLIN-CHEMIE AG	SI
Siofor 850 mg filmsko obložene tablete	not available	H/02/01427/004	BERLIN-CHEMIE AG	SI
Siofor 850 mg filmsko obložene tablete	not available	H/02/01427/003	BERLIN-CHEMIE AG	SI
Stagid 700 mg comprimidos	not available	9546820	MERCK, S.A.	PT
Stagid 700 mg comprimidos	not available	9546838	MERCK, S.A.	PT
STAGID 700 mg, comprimé sécable	not available	3400931927679	MERCK SANTÉ S.A.S.	FR
STAGID 700 mg, comprimé sécable	not available	3400931927501	MERCK SANTÉ S.A.S.	FR
Глюкофаж 1000 mg филмирани таблетки	FR/H/0181/001	II-4493	MERCK SANTÉ S.A.S.	BG
Глюкофаж 500 mg филмирани таблетки	FR/H/0181/003	II-9301	MERCK SANTÉ S.A.S.	BG
Глюкофаж 850 mg филмирани таблетки	FR/H/0181/002	II-9302	MERCK SANTÉ S.A.S.	BG