11 January 2018
EMA/25531/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: glimepiride

Procedure no.: PSUSA/00001534/201706

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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
Amaryl 1 mg tablete	NL/H/0101/001	H/97/00159/001	SANOFI-AVENTIS D.O.O.	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/024	SANOFI-AVENTIS D.O.O.	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/013	SANOFI-AVENTIS D.O.O.	SI
GLIMEPIRID-ZENTIVA 4 MG TABLETTA	not available	OGYI-T-10452/01	ZENTIVA HU KFT	HU
AMARYL 2 MG TABLETE	not available	HR-H-836629436-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
AMARYL 3 MG TABLETE	not available	HR-H-851734879-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
AMARYL 2 MG TABLETE	not available	HR-H-836629436-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
AMARYL 3 MG TABLETE	not available	HR-H-851734879-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
AMARYL 1 MG-TABLETTEN	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Solosa 4 mg, δισκίο	NL/H/0101/004	228600411	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	228600211	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600112	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	228600311	SANOFI-AVENTIS AEBE	GR
AMARYL 4 mg tabletta	not available	OGYI-T-5746/04	SANOFI-AVENTIS ZRT	HU
AMARYL 1 mg tabletta	not available	OGYI-T-5746/01	SANOFI-AVENTIS ZRT	HU
AMARYL 2 mg tabletta	not available	OGYI-T-5746/02	SANOFI-AVENTIS ZRT	HU
AMARYL 3	not available	7450	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
AMARYL 4	not available	7451	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
AMARYL 1	not available	7448	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
AMARYL 2	not available	7449	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
AMARYL 6 MG TABLETTA	not available	OGYI-T-5746/05	SANOFI-AVENTIS ZRT	HU
AMARYL 3 MG TABLETTA	not available	OGYI-T-5746/03	SANOFI-AVENTIS ZRT	HU
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Glimepiride Winthrop 6 mg, tablet	NL/H/0622/005	RVG 31978	WINTHROP MEDICAMENTS	NL
Glimepirid Winthrop® 2 mg Tabletten	NL/H/0622/002	63045.01.00	WINTHROP ARZNEIMITTEL 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
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/06	SANOFI ROMANIA SRL	RO
Amaryl® 6 mg Tabletten	NL/H/0101/005	32638.04.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/07	SANOFI ROMANIA SRL	RO
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	567 635-7	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE WINTHROP 1 MG, TABLET	NL/H/0622/001	RVG 31974	WINTHROP MEDICAMENTS	NL
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	370 622-6	SANOFI-AVENTIS FRANCE	FR
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	370 624-9	SANOFI-AVENTIS FRANCE	FR
AMARYL 4 mg, tabletten	NL/H/0101/004	RVG 17846	SANOFI-AVENTIS NETHERLANDS B.V.	NL
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845455	SANOFI SPA	IT
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845238	SANOFI SPA	IT
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845226	SANOFI SPA	IT
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	370 623-2	SANOFI-AVENTIS FRANCE	FR
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845214	SANOFI SPA	IT
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845240	SANOFI SPA	IT
Amarylle 4 mg, comprimés	NL/H/0101/004	0351376	SANOFI BELGIUM	LU
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845479	SANOFI SPA	IT
Amarylle 4 mg, comprimés	NL/H/0101/004	0351345	SANOFI BELGIUM	LU
Amarylle 4 mg, comprimés	NL/H/0101/004	0351362	SANOFI BELGIUM	LU
GLIMEPIRIDE WINTHROP 3 MG, TABLET	NL/H/0622/003	RVG 31976	WINTHROP MEDICAMENTS	NL
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Amarylle 4 mg, comprimés	NL/H/0101/004	0351301	SANOFI BELGIUM	LU
Amarylle 4 mg, comprimés	NL/H/0101/004	0351328	SANOFI BELGIUM	LU
Amaryl 4 mg, tablet	NL/H/0101/004	PL 17780/0622	WINTHROP PHARMACEUTICALS UK LTD	UK
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845253	SANOFI SPA	IT
Amarylle 4 mg, comprimés	NL/H/0101/004	0351393	SANOFI BELGIUM	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
Amarylle 4 mg, comprimés	NL/H/0101/004	0351331	SANOFI BELGIUM	LU
Glimepirid Winthrop® 3 mg Tabletten	NL/H/0622/003	63045.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
Amaryl 4 mg, tablett	NL/H/0101/004	02-1339	SANOFI-AVENTIS NORGE AS	NO
Amaryl, 6 mg tabletid	NL/H/0101/005	642909	SANOFI-AVENTIS ESTONIA OÜ	EE
AMARYL 4 MG TABLETTEN	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amarylle 4 mg, comprimés	NL/H/0101/004	0636779	SANOFI BELGIUM	LU
Amarylle 4 mg, comprimés	NL/H/0101/004	0351359	SANOFI BELGIUM	LU
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	NL31672	SANOFI-AVENTIS FRANCE	FR
Glimepirid Winthrop® 4 mg Tabletten	NL/H/0622/004	63045.03.00	WINTHROP ARZNEIMITTEL GMBH	DE
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé.	NL/H/0622/002	370 630-9	SANOFI-AVENTIS FRANCE	FR
GLIMÉPIRIDE ZENTIVA 2 mg, comprimé.	NL/H/0622/002	370 625-5	SANOFI-AVENTIS FRANCE	FR
Amarylle 3 mg, comprimés	NL/H/0101/003	0351278	SANOFI BELGIUM	LU
Amaryl 4 mg comprimidos	NL/H/0101/004	61.409	SANOFI-AVENTIS, S.A.	ES
Amarylle 4 mg, comprimés	NL/H/0101/004	0351314	SANOFI BELGIUM	LU
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 619-5	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 4 mg tabletti	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/046	SANOFI-AVENTIS D.O.O.	SI
AMAREL 3 mg, comprimé	NL/H/0101/003	366 859-5	SANOFI-AVENTIS FRANCE	FR
AMARYL 1 MG, TABLET	NL/H/0101/001	PL 17780/0619	WINTHROP PHARMACEUTICALS UK LTD	UK
Amarylle 3 mg, comprimés	NL/H/0101/003	0351216	SANOFI BELGIUM	LU
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 527-3	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 618-9	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 526-7	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/10	SANOFI ROMANIA SRL	RO
Amarylle 4 mg, tabletten	NL/H/0101/004	BE251282	SANOFI BELGIUM	BE
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/02	SANOFI ROMANIA SRL	RO
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 620-3	SANOFI-AVENTIS FRANCE	FR
GLIMÉPIRIDE ZENTIVA 1 mg, comprimé.	NL/H/0622/001	NL31671	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/08	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/09	SANOFI ROMANIA SRL	RO
AMAREL 3 mg, comprimé	NL/H/0101/003	366 858-9	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 MG, COMPRIME	NL/H/0622/001	370 529-6	SANOFI-AVENTIS FRANCE	FR
Amarylle 3 mg, comprimés	NL/H/0101/003	0351233	SANOFI BELGIUM	LU
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/03	SANOFI ROMANIA SRL	RO
AMAREL 3 mg, comprimé	NL/H/0101/003	342 104-4	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 mg, comprimé	NL/H/0101/003	366 860-3	SANOFI-AVENTIS FRANCE	FR
Amarylle 3 mg, comprimés	NL/H/0101/003	0351247	SANOFI BELGIUM	LU
Amarylle 3 mg, comprimés	NL/H/0101/003	0351251	SANOFI BELGIUM	LU
Amarylle 3 mg, comprimés	NL/H/0101/003	0351202	SANOFI BELGIUM	LU
Amarylle 3 mg, comprimés	NL/H/0101/003	0351264	SANOFI BELGIUM	LU
Amarylle 3 mg, comprimés	NL/H/0101/003	0351281	SANOFI BELGIUM	LU
Amaryl 3 mg, tablet	NL/H/0101/003	16884	SANOFI-AVENTIS DENMARK A/S	DK
Amaryl, tabletter	NL/H/0101/004	16885	SANOFI-AVENTIS DENMARK A/S	DK
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Amarylle 3 mg, comprimés	NL/H/0101/003	0636765	SANOFI BELGIUM	LU
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/08	SANOFI ROMANIA SRL	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
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/10	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/04	SANOFI ROMANIA SRL	RO
Amarylle 3 mg, comprimés	NL/H/0101/003	0351295	SANOFI BELGIUM	LU
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/11	SANOFI ROMANIA SRL	RO
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845428	SANOFI SPA	IT
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845176	SANOFI SPA	IT
Amaryl® 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845442	SANOFI SPA	IT
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845430	SANOFI SPA	IT
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845164	SANOFI SPA	IT
AMARYL	NL/H/0101/003	9800023	SANOFI BULGARIA EOOD	BG
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845152	SANOFI SPA	IT
AMARYL 3 MG TABLETY	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/01	SANOFI ROMANIA SRL	RO
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845416	SANOFI SPA	IT
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845202	SANOFI SPA	IT
AMAREL 1 mg, comprimé	NL/H/0101/001	NL 20477	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	NL 20477	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	560 086-8	SANOFI-AVENTIS FRANCE	FR
Amaryl 3 mg, tablet	NL/H/0101/003	PA 540/28/3	SANOFI-AVENTIS IRELAND LTD	IE
AMAREL 1 mg, comprimé	NL/H/0101/001	342 102-1	SANOFI-AVENTIS FRANCE	FR
Amaryl 3 mg tabletes	NL/H/0101/003	09-0323	SANOFI-AVENTIS LATVIA SIA	LV
AMAREL 1 mg, comprimé	NL/H/0101/001	366 850-8	SANOFI-AVENTIS FRANCE	FR
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845188	SANOFI SPA	IT
AMAREL 1 mg, comprimé	NL/H/0101/001	366 851-4	SANOFI-AVENTIS FRANCE	FR
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS ESTONIA OÜ	EE
AMARYL 3 MG, COMPRESSE	NL/H/0101/003	032845190	SANOFI 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
Amaryl 3 mg, tafla	NL/H/0101/003	IS/1/02/040/03	SANOFI-AVENTIS NORGE AS	IS
AMAREL 1 mg, comprimé	NL/H/0101/001	NL 20477	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	347 112-5	SANOFI-AVENTIS FRANCE	FR
AMARYL 3 MG, TABLET	NL/H/0101/003	PL 17780/0621	WINTHROP PHARMACEUTICALS UK LTD	UK
AMAREL 1 mg, comprimé	NL/H/0101/001	366 848-3	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	NL 20477	SANOFI-AVENTIS FRANCE	FR
AMARYL 1 MG TABLETT	NL/H/0101/001	02-1336	SANOFI-AVENTIS NORGE AS	NO
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845380	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845378	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845024	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845125	SANOFI SPA	IT
AMAREL 1 mg, comprimé	NL/H/0101/001	366 847-7	SANOFI-AVENTIS FRANCE	FR
Amaryl 2 mg comprimidos	NL/H/0101/002	61.406	SANOFI-AVENTIS, S.A.	ES
Amarylle 3 mg, tabletten	NL/H/0101/003	BE251273	SANOFI BELGIUM	BE
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI BULGARIA EOOD	BG
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845113	SANOFI SPA	IT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg, tablet	NL/H/0101/001	16882	SANOFI-AVENTIS DENMARK A/S	DK
Amarylle 2 mg, comprimés	NL/H/0101/002	0636751	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	244636	SANOFI BELGIUM	LU
AMARYL 2 MG TABLETT	NL/H/0101/002	02-1337	SANOFI-AVENTIS NORGE AS	NO
Amaryl 4 mg tablett	NL/H/0101/004	18711	SANOFI AB	SE
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845137	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845392	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845012	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845404	SANOFI SPA	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845149	SANOFI SPA	IT
Amarylle 2 mg, comprimés	NL/H/0101/002	244653	SANOFI BELGIUM	LU
AMARYL 3 MG TABLETT	NL/H/0101/003	02-1338	SANOFI-AVENTIS NORGE AS	NO
Solosa 1 mg, δισκίο	NL/H/0101/001	228600110	SANOFI-AVENTIS AEBE	GR
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS ESTONIA OÜ	EE
AMAREL 3 mg, comprimé	NL/H/0101/003	560 088-0	SANOFI-AVENTIS FRANCE	FR

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AMAREL 3 mg, comprimé	NL/H/0101/003	NL 21528	SANOFI-AVENTIS FRANCE	FR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600108	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600101	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600103	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600107	SANOFI-AVENTIS AEBE	GR
AMAREL 3 mg, comprimé	NL/H/0101/003	NL 21528	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 mg, comprimé	NL/H/0101/002	366 854-3	SANOFI-AVENTIS FRANCE	FR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600111	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600105	SANOFI-AVENTIS AEBE	GR
AMAREL 3 mg, comprimé	NL/H/0101/003	366 857-2	SANOFI-AVENTIS FRANCE	FR
AMARYL® 3 MG, ΔΙΣΚΊΟ	NL/H/0101/003	20552	SANOFI-AVENTIS CYPRUS LTD	CY
Solosa 1 mg, δισκίο	NL/H/0101/001	228600102	SANOFI-AVENTIS AEBE	GR
AMAREL 2 mg, comprimé	NL/H/0101/002	366 853-7	SANOFI-AVENTIS FRANCE	FR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600106	SANOFI-AVENTIS AEBE	GR
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
AMAREL 3 mg, comprimé	NL/H/0101/003	NL 21528	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 mg, comprimé	NL/H/0101/002	366 856-6	SANOFI-AVENTIS FRANCE	FR
Amarylle 2 mg, tabletten	NL/H/0101/002	BE185534	SANOFI BELGIUM	BE
Solosa 1 mg, δισκίο	NL/H/0101/001	228600109	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	228600104	SANOFI-AVENTIS AEBE	GR
Amaryl 3 mg, tabletten	NL/H/0101/003	17845	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Solosa 2 mg, δισκίο	NL/H/0101/002	228600201	SANOFI-AVENTIS AEBE	GR
Amaryl 1 mg tabletes	NL/H/0101/001	09-0321	SANOFI-AVENTIS LATVIA SIA	LV
Amaryl 2 mg, tablet	NL/H/0101/002	16883	SANOFI-AVENTIS DENMARK A/S	DK
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/10	SANOFI ROMANIA SRL	RO
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI BULGARIA EOOD	BG
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
Solosa 2 mg, δισκίο	NL/H/0101/002	228600206	SANOFI-AVENTIS AEBE	GR
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/08	SANOFI ROMANIA SRL	RO
Solosa 2 mg, δισκίο	NL/H/0101/002	228600205	SANOFI-AVENTIS AEBE	GR
AMAREL 3 mg, comprimé	NL/H/0101/003	NL 21528	SANOFI-AVENTIS FRANCE	FR
Solosa 2 mg, δισκίο	NL/H/0101/002	228600202	SANOFI-AVENTIS AEBE	GR

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Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/023	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/021	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/017	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/016	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/031	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/018	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/002	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/028	UAB SANOFI-AVENTIS LIETUVA	LT
AMARYL® 4 MG, ΔΙΣΚΊΟ	NL/H/0101/004	20553	SANOFI-AVENTIS CYPRUS LTD	CY
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/038	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/022	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg, tafla	NL/H/0101/004	IS/1/02/040/04	SANOFI-AVENTIS NORGE AS	IS
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/004	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600309	SANOFI-AVENTIS AEBE	GR
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/024	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/030	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg, tafla	NL/H/0101/004	IS/1/02/040/04	SANOFI-AVENTIS NORGE AS	IS
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/020	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600308	SANOFI-AVENTIS AEBE	GR
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/027	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/039	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600304	SANOFI-AVENTIS AEBE	GR
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/025	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/036	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tablett	NL/H/0101/002	12231	SANOFI AB	SE
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/044	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600303	SANOFI-AVENTIS AEBE	GR
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/014	UAB SANOFI-AVENTIS LIETUVA	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
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/09	SANOFI ROMANIA SRL	RO
Solosa 3 mg, δισκίο	NL/H/0101/003	228600301	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	228600302	SANOFI-AVENTIS AEBE	GR
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/019	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/003	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/026	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600310	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/043	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600306	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	228600307	SANOFI-AVENTIS AEBE	GR
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/032	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/037	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/033	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/045	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/029	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/015	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg, tablet	NL/H/0101/002	PL 17780/0620	WINTHROP PHARMACEUTICALS UK LTD	UK
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/034	UAB SANOFI-AVENTIS LIETUVA	LT
Solosa 3 mg, δισκίο	NL/H/0101/003	228600305	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845354	SANOFI SPA	IT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/035	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/040	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 2 mg tabletes	NL/H/0101/002	09-0322	SANOFI-AVENTIS LATVIA SIA	LV
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS CYPRUS LTD	CY
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Solosa 2 mg, δισκίο	NL/H/0101/002	228600209	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845366	SANOFI 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
Amaryl 1 mg, tablet	NL/H/0101/001	PA 540/28/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Solosa 2 mg, δισκίο	NL/H/0101/002	228600204	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	228600207	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845099	SANOFI SPA	IT
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845339	SANOFI SPA	IT
Solosa 2 mg, δισκίο	NL/H/0101/002	228600203	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845063	SANOFI SPA	IT
Solosa 2 mg, δισκίο	NL/H/0101/002	228600210	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845101	SANOFI SPA	IT
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845087	SANOFI SPA	IT
Solosa 2 mg, δισκίο	NL/H/0101/002	228600208	SANOFI-AVENTIS AEBE	GR
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845341	SANOFI SPA	IT
Amaryl 1 mg, tablet	NL/H/0101/001	PA 540/28/1	SANOFI-AVENTIS IRELAND LTD	IE
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845075	SANOFI SPA	IT
AMARYL 1 MG TABLETT	NL/H/0101/001	12230	SANOFI AB	SE
AMARYL 1 MG, COMPRESSE	NL/H/0101/001	032845051	SANOFI SPA	IT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/013	UAB SANOFI-AVENTIS LIETUVA	LT
AMAREL 2 mg, comprimé	NL/H/0101/002	366 852-0	SANOFI-AVENTIS FRANCE	FR
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/006	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/012	UAB SANOFI-AVENTIS LIETUVA	LT
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS CYPRUS LTD	CY
AMAREL 2 mg, comprimé	NL/H/0101/002	NL 20478	SANOFI-AVENTIS FRANCE	FR
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/008	UAB SANOFI-AVENTIS LIETUVA	LT
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/001	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/009	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/007	UAB SANOFI-AVENTIS LIETUVA	LT
AMAREL 2 mg, comprimé	NL/H/0101/002	NL 20478	SANOFI-AVENTIS FRANCE	FR
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/041	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl, 1 mg tabletid	NL/H/0101/001	272799	SANOFI-AVENTIS ESTONIA OÜ	EE
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/042	UAB SANOFI-AVENTIS LIETUVA	LT
AMAREL 2 mg, comprimé	NL/H/0101/002	NL 20478	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
Amaryl 2 mg, tafla	NL/H/0101/002	IS/1/02/040/02	SANOFI-AVENTIS NORGE AS	IS
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/005	UAB SANOFI-AVENTIS LIETUVA	LT
AMAREL 2 mg, comprimé	NL/H/0101/002	560 087-4	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 mg, comprimé	NL/H/0101/002	NL 20478	SANOFI-AVENTIS FRANCE	FR
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/010	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/011	UAB SANOFI-AVENTIS LIETUVA	LT
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/01	SANOFI ROMANIA SRL	RO
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI BULGARIA EOOD	BG
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/12	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/10	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/06	SANOFI ROMANIA SRL	RO
AMAREL 2 mg, comprimé	NL/H/0101/002	342 103-8	SANOFI-AVENTIS FRANCE	FR
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/08	SANOFI ROMANIA SRL	RO
Amaryl® 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	567 639-2	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	371 187-1	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	370 631-5	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	370 632-1	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	370 626-1	SANOFI-AVENTIS FRANCE	FR
Glimepiride Winthrop 2 mg, tablet	NL/H/0622/002	RVG 31975	WINTHROP MEDICAMENTS	NL
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	371 188-8	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	370 629-0	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	370 628-4	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	567 642-3	SANOFI-AVENTIS FRANCE	FR
GLIMÉPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	NL31673	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 MG, COMPRIME	NL/H/0622/004	567 640-0	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	370 627-8	SANOFI-AVENTIS FRANCE	FR
GLIMÉPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	NL31674	SANOFI-AVENTIS FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 MG, COMPRIME	NL/H/0622/003	567 638-6	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg, compresse	NL/H/0101/005	032845529	SANOFI SPA	IT
Amaryl 6 mg, compresse	NL/H/0101/005	032845315	SANOFI SPA	IT
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845467	SANOFI SPA	IT
Amaryl 6 mg, compresse	NL/H/0101/005	032845277	SANOFI SPA	IT
Amaryl 6 mg, compresse	NL/H/0101/005	032845327	SANOFI SPA	IT
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845481	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	366 865-5	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg, compresse	NL/H/0101/005	032845289	SANOFI SPA	IT
Amaryl 6 mg tabletes	NL/H/0101/005	09-0333	SANOFI-AVENTIS LATVIA SIA	LV
AMAREL 4 mg, comprimé	NL/H/0101/004	560 089-7	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	366 864-9	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg, compresse	NL/H/0101/005	032845291	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	366 863-2	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	NL 20479	SANOFI-AVENTIS FRANCE	FR
AMARYL 4 MG, COMPRESSE	NL/H/0101/004	032845265	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	366 862-6	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg, compresse	NL/H/0101/005	032845505	SANOFI SPA	IT
Amaryl 6 mg, compresse	NL/H/0101/005	032845493	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	342 105-0	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amaryl 6 mg, compresse	NL/H/0101/005	032845517	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	NL 20479	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	NL 20479	SANOFI-AVENTIS FRANCE	FR
Amaryl 6 mg, compresse	NL/H/0101/005	032845303	SANOFI SPA	IT
AMAREL 4 mg, comprimé	NL/H/0101/004	NL 20479	SANOFI-AVENTIS FRANCE	FR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600410	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600409	SANOFI-AVENTIS AEBE	GR
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/08	SANOFI ROMANIA SRL	RO
Glimepiride Winthrop 4 mg, tablet	NL/H/0622/004	RVG 31977	WINTHROP MEDICAMENTS	NL
Glimepirid Winthrop® 1 mg Tabletten	NL/H/0622/001	63045.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/10	SANOFI ROMANIA SRL	RO
Glimepirid Winthrop® 6 mg Tabletten	NL/H/0622/005	63045.04.00	WINTHROP ARZNEIMITTEL GMBH	DE
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Solosa 4 mg, δισκίο	NL/H/0101/004	228600405	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600403	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600401	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600402	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600408	SANOFI-AVENTIS AEBE	GR
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS ESTONIA OÜ	EE
Solosa 4 mg, δισκίο	NL/H/0101/004	228600407	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	228600404	SANOFI-AVENTIS AEBE	GR
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS LATVIA SIA	LV
Solosa 4 mg, δισκίο	NL/H/0101/004	228600406	SANOFI-AVENTIS AEBE	GR
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/09	SANOFI ROMANIA SRL	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
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/035	SANOFI-AVENTIS D.O.O.	SI
Glimepirid-Zentiva 1 mg tabletta	not available	OGYI-T-10452/03	ZENTIVA HU KFT	HU
Glimepirid-Zentiva 2 mg tabletta	not available	OGYI-T-10452/02	ZENTIVA HU KFT	HU
Solosa 6 mg, compresse	not available	032117499	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117273	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117323	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117513	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117309	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117525	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117311	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117297	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117285	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 6 mg, compresse	not available	032117501	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117020	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117057	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117386	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117400	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117398	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117374	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117083	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117071	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117069	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 2 mg, compresse	not available	032117018	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117335	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117145	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117347	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117095	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117350	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117487	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117210	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117261	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117463	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117475	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117222	LABORATORI GUIDOTTI 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
Solosa 4 mg, compresse	not available	032117234	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117259	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117246	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117362	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117119	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117121	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117133	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117436	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117208	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117412	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117424	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117158	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117160	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117184	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117448	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117172	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 3 mg, compresse	not available	032117196	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 1 mg, compresse	not available	032117107	LABORATORI GUIDOTTI S.P.A.	IT
Solosa 4 mg, compresse	not available	032117451	LABORATORI GUIDOTTI S.P.A.	IT
RONAME 2 mg comprimidos	not available	62.808	LACER S.A.	ES
RONAME 4 mg comprimidos	not available	62.807	LACER S.A.	ES
Glimepirid Sandoz 4 mg tablety	DK/H/0802/004	18/397/05-C	SANDOZ GMBH	CZ
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0417111	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0417075	HEXAL AG	LU
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0417139	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0417092	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0715/05110014	HEXAL AG	LU
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0417108	HEXAL AG	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
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0417125	HEXAL AG	LU
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0417142	HEXAL AG	LU
GLIMEPIRID HEXAL 6 MG TABLETTEN	DE/H/2141/005	0715/05110015	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0417058	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0417089	HEXAL AG	LU
GLIMEPIRID HEXAL 4 MG TABLETTEN	DE/H/2141/004	0417061	HEXAL AG	LU
Glimepirid 1A Pharma 4 mg Tabletten	DE/H/2142/004	63822.03.00	1 A PHARMA GMBH	DE
Glimepirid HEXAL 6 mg Tabletten	DE/H/2141/005	63817.04.00	HEXAL AG	DE
Glimepirid 1A Pharma 6 mg Tabletten	DE/H/2142/005	63822.04.00	1 A PHARMA GMBH	DE
Glimepirid HEXAL 4 mg Tabletten	DE/H/2141/004	63817.03.00	HEXAL AG	DE