09 February 2023
EMA/PRAC/83881/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): glimepiride

Procedure No.: PSUSA/00001534/202206

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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-Zentiva 4 mg tabletta	not available	OGYI-T-10452/01	ZENTIVA, K.S.	HU
Glimepirid-Zentiva 2 mg tabletta	not available	OGYI-T-10452/02	ZENTIVA, K.S.	HU
Amaryl 1, 1 mg, tabletki	not available	7448	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 3, 3 mg, tabletki	not available	7450	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 3, 3 mg, tabletki	not available	7450	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 4, 4 mg, tabletki	not available	7451	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 1, 1 mg, tabletki	not available	7448	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 2, 2 mg, tabletki	not available	7449	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 2, 2 mg, tabletki	not available	7449	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Amaryl 4, 4 mg, tabletki	not available	7451	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
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
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

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 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

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	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
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

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 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
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 mg tabletta	not available	OGYI-T-5746/03	SANOFI-AVENTIS ZRT	HU
Glimepirid - 1 A Pharma® 4 mg Tabletten	DE/H/2142/004	63822.03.00	1 A PHARMA GMBH	DE
Glimepirid - 1 A Pharma® 6 mg Tabletten	DE/H/2142/005	63822.04.00	1 A PHARMA GMBH	DE
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 619 5 7	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 620 3 9	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 529 6 2	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 618 9 6	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 527 3 3	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 1 mg, comprimé	NL/H/0622/001	34009 370 526 7 2	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	34009 370 625 5 8	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	34009 370 630 9 8	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	34009 567 635 7 3	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	34009 370 622 6 8	ZENTIVA 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 2 mg, comprimé	NL/H/0622/002	34009 370 623 2 9	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 2 mg, comprimé	NL/H/0622/002	34009 370 624 9 7	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 370 628 4 8	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 370 627 8 7	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 370 629 0 9	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 567 638 6 3	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 371 187 1 2	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 3 mg, comprimé	NL/H/0622/003	34009 370 626 1 9	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 371 188 8 0	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 370 632 1 0	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 567 639 2 4	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 567 642 3 5	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 370 631 5 9	ZENTIVA FRANCE	FR
GLIMEPIRIDE ZENTIVA 4 mg, comprimé	NL/H/0622/004	34009 567 640 0 6	ZENTIVA FRANCE	FR
Glimepirid Winthrop 1 mg Tabletten	NL/H/0622/001	63045.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Glimepirid Winthrop 2 mg Tabletten	NL/H/0622/002	63045.01.00	WINTHROP ARZNEIMITTEL GMBH	DE
Glimepirid Winthrop 3 mg Tabletten	NL/H/0622/003	63045.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
Glimepirid Winthrop 4 mg Tabletten	NL/H/0622/004	63045.03.00	WINTHROP ARZNEIMITTEL GMBH	DE
Glimepirid Winthrop 6 mg Tabletten	NL/H/0622/005	63045.04.00	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
Glimepiride Zentiva 1 mg, tabletten	NL/H/0622/001	RVG 31974	ZENTIVA, K.S.	NL
Glimepiride Zentiva 2 mg, tabletten	NL/H/0622/002	RVG 31975	ZENTIVA, K.S.	NL
Glimepiride Zentiva 3 mg, tabletten	NL/H/0622/003	RVG 31976	ZENTIVA, K.S.	NL
Glimepiride Zentiva 4 mg, tabletten	NL/H/0622/004	RVG 31977	ZENTIVA, K.S.	NL
Glimepiride Zentiva 6 mg, tabletten	NL/H/0622/005	RVG 31978	ZENTIVA, K.S.	NL
Glimepirid HEXAL® 6 mg Tabletten	DE/H/2141/005	63817.04.00	HEXAL AG	DE
Glimepirid HEXAL® 4 mg Tabletten	DE/H/2141/004	63817.03.00	HEXAL AG	DE
Amaryl 2 mg tablete	not available	HR-H-836629436-02	SANOFI-AVENTIS GROUPE	HR
Amaryl 3 mg tablete	not available	HR-H-851734879-01	SANOFI-AVENTIS GROUPE	HR
Amaryl 2 mg tablete	not available	HR-H-836629436-01	SANOFI-AVENTIS GROUPE	HR
Amaryl 3 mg tablete	not available	HR-H-851734879-02	SANOFI-AVENTIS GROUPE	HR
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	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
Amaryl 3 mg, tablet	NL/H/0101/003	PA 540/28/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Amaryl 1 mg, tablet	NL/H/0101/001	PA 540/28/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Amaryl 1 mg, tablet	NL/H/0101/001	PA 540/28/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amarylle 2 mg, comprimés	NL/H/0101/002	0636751	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0244636	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0244653	SANOFI BELGIUM	LU
Amaryl 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND 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 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	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
AMARYL 3 MG TABLETTI	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
AMARYL 4 MG TABLETTI	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	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
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
AMARYL 1 MG TABLETTI	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/10	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	2419/2010/08	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 6 mg, tabletten	NL/H/0101/005	RVG 17847	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG TABLETTI	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845012	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845404	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845149	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845125	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845392	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845378	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845137	SANOFI S.R.L.	IT
AMARYL 2 MG, COMPRESSE	NL/H/0101/002	032845024	SANOFI S.R.L.	IT
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR

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 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Amaryl 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 3 mg, compresse	NL/H/0101/003	032845442	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845467	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845481	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845265	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845238	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845226	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845240	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845214	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845479	SANOFI S.R.L.	IT
Amaryl 4 mg, compresse	NL/H/0101/004	032845253	SANOFI S.R.L.	IT
Amaryl 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS	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
			DEUTSCHLAND GMBH	
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR

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, δισκio	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκio	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκio	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκio	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 560 087-4 2	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 366 852-0 8	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 366 854-3 7	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 366 856-6 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 366 853-7 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 2 MG, COMPRIMÉ	NL/H/0101/002	34009 342 103-8 9	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 342 104-4 0	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 366 857-2 7	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 560 088-0 3	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 366 859-5 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 366 860-3 8	SANOFI-AVENTIS FRANCE	FR
AMAREL 3 MG, COMPRIMÉ	NL/H/0101/003	34009 366 858-9 5	SANOFI-AVENTIS FRANCE	FR
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl 2 mg comprimidos	NL/H/0101/002	61.406	SANOFI-AVENTIS, S.A.	ES
Amaryl 4 mg comprimidos	NL/H/0101/004	61.409	SANOFI-AVENTIS, S.A.	ES
Amaryl 4 mg comprimidos	NL/H/0101/004	61.409	SANOFI-AVENTIS, S.A.	ES
Amaryl 2 mg comprimidos	NL/H/0101/002	61.406	SANOFI-AVENTIS, S.A.	ES
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	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
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 3 mg tablety	NL/H/0101/003	18/0483/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/013	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/006	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/012	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/008	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/001	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/009	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/042	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/007	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/005	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/010	SANOFI-AVENTIS GROUPE	LT
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/041	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/015	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/021	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/018	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/002	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/017	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/016	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/022	SANOFI-AVENTIS GROUPE	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 tabletės	NL/H/0101/002	LT/1/99/1288/020	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/014	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/019	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/023	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/027	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/028	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/024	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/031	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/030	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/025	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/026	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/045	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/034	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/035	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/004	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/038	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/039	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/036	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/037	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/033	SANOFI-AVENTIS GROUPE	LT
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
AMARYL, 1 MG TABLETID	NL/H/0101/001	272799	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE

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 tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
AMARYL, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	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
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL® 1 MG, ΔΙΣΚΊΟ	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
AMARYL 2 MG, TABLET	NL/H/0101/002	H/97/00159/013	SANOFI-AVENTIS GROUPE	SI
AMARYL 3 MG, TABLET	NL/H/0101/003	H/97/00459/024	SANOFI-AVENTIS GROUPE	SI
AMARYL 6 MG TABLETE	NL/H/0101/005	H/97/00159/046	SANOFI-AVENTIS GROUPE	SI
AMARYL 4 MG TABLETE	NL/H/0101/004	H/97/00159/035	SANOFI-AVENTIS GROUPE	SI
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 2 mg таблетки	NL/H/0101/002	9800224	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG

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
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
AMARYL® 2 MG, ΔΙΣΚΊΟ	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
Amaryl® 1 mg, δισκίο	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
Amaryl® 1 mg, δισκίο	NL/H/0101/001	20550	SANOFI-AVENTIS GROUPE	CY
Amaryl® 2 mg, δισκίο	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg Tabletten	NL/H/0101/002	32638.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 6 mg Tabletten	NL/H/0101/005	1-21663	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE
Amaryl, 2 mg tabletid	NL/H/0101/002	272899	SANOFI-AVENTIS GROUPE	EE

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 tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/032	SANOFI-AVENTIS GROUPE	LT
Amaryl 6 mg comprimate	NL/H/0101/005	2419/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amarylle 2 mg, comprimés	NL/H/0101/002	0883007	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882979	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882948	SANOFI BELGIUM	LU
Amaryl® 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg tabletter	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 3 mg tabletter	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletti	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 3 mg tabletti	NL/H/0101/003	12716	SANOFI OY	FI
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg, compresse	NL/H/0101/002	032845380	SANOFI S.R.L.	IT
Amaryl 2 mg tabletės	NL/H/0101/002	LT/1/99/1288/043	SANOFI-AVENTIS GROUPE	LT
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 2 mg, tabletten	NL/H/0101/002	RVG 17844	GENZYME EUROPE B.V.	NL
Amaryl 3 mg, tabletten	NL/H/0101/003	RVG 17845	GENZYME EUROPE B.V.	NL
Amaryl® 2 mg, δισκίο	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	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
Amaryl 4 mg, tabletten	NL/H/0101/004	RVG 17846	GENZYME EUROPE B.V.	NL
Amaryl® 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 2 mg, δισκίο	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
Solosa 1 mg, δισκίο	NL/H/0101/001	33821/95/18-10-96	SANOFI-AVENTIS AEBE	GR
Solosa 4 mg, δισκίο	NL/H/0101/004	35690/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Амарил 1 mg таблетки	NL/H/0101/001	9800222	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 3 mg таблетки	NL/H/0101/003	9800223	SANOFI-AVENTIS GROUPE	BG
Амарил 4 mg таблетки	NL/H/0101/004	20020600	SANOFI-AVENTIS GROUPE	BG
Solosa 2 mg, δισκίο	NL/H/0101/002	35688/17-05-2011	SANOFI-AVENTIS AEBE	GR
Amaryl® 2 mg, δισκίο	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/018	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/022	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/015	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/017	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/020	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/019	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/014	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/023	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/016	SANOFI-AVENTIS GROUPE	SI
Amaryl 2 mg tablete	NL/H/0101/002	H/97/00159/021	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/032	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/033	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/027	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/026	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/034	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/029	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/028	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/031	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/025	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg tablete	NL/H/0101/003	H/97/00159/030	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/040	SANOFI-AVENTIS GROUPE	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
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/043	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/042	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/036	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/039	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/038	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/037	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/044	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/041	SANOFI-AVENTIS GROUPE	SI
Amaryl 4 mg tablete	NL/H/0101/004	H/97/00159/045	SANOFI-AVENTIS GROUPE	SI
AMAREL 1 mg, comprimé	NL/H/0101/001	NL 20477	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	34009 366 847-7 5	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	34009 342 102-1 1	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	34009 560 086-8 1	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	34009 366 850-8 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	34009 366 851-4 7	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	34009 347 112 5 1	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	34009 366 848-3 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 366 865-5 7	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 560 089-7 1	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 366 864-9 6	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 366 862-6 7	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 342 105-0 1	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	34009 366 863-2 8	SANOFI-AVENTIS FRANCE	FR
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
AMAREL 4 mg, comprimé	NL/H/0101/004	NL 20479	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	NL20477	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	NL20479	SANOFI-AVENTIS FRANCE	FR
AMAREL 4 mg, comprimé	NL/H/0101/004	NL20479	SANOFI-AVENTIS FRANCE	FR
AMAREL 1 mg, comprimé	NL/H/0101/001	NL20477	SANOFI-AVENTIS FRANCE	FR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR

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 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
Solosa 3 mg, δισκίο	NL/H/0101/003	35689/17-05-2011	SANOFI-AVENTIS AEBE	GR
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
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/10	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	2417/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 3 mg comprimate	NL/H/0101/003	2417/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/03	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/09	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/07	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/10	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/08	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
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/04	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/01	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/05	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/02	SANOFI ROMANIA SRL	RO
Amaryl 1 mg comprimate	NL/H/0101/001	2415/2010/12	SANOFI ROMANIA SRL	RO
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/05	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 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	2415/2010/08	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
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/08	SANOFI ROMANIA SRL	RO
Amaryl 4 mg comprimate	NL/H/0101/004	2418/2010/10	SANOFI ROMANIA SRL	RO
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	2418/2010/11	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	2418/2010/06	SANOFI ROMANIA SRL	RO
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	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
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amarylle 2 mg, comprimés	NL/H/0101/002	0882934	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882982	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882951	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882965	SANOFI BELGIUM	LU
Amarylle 2 mg, comprimés	NL/H/0101/002	0882996	SANOFI BELGIUM	LU
Amaryl 3 mg, compresse	NL/H/0101/003	032845428	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845416	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845176	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845202	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845152	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845164	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845430	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845190	SANOFI S.R.L.	IT
Amaryl 3 mg, compresse	NL/H/0101/003	032845188	SANOFI S.R.L.	IT
Amaryl® 1 mg Tabletten	NL/H/0101/001	32638.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl® 3 mg Tabletten	NL/H/0101/003	32638.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH	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
			OSTERREICH	
Amaryl 1 mg Tabletten	NL/H/0101/001	1-21662	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg comprimate	NL/H/0101/002	2416/2010/11	SANOFI ROMANIA SRL	RO
Amaryl 3 mg tablety	NL/H/0101/003	18/234/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/053	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/048	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/047	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/050	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/055	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/056	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/052	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/051	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/054	SANOFI-AVENTIS GROUPE	SI
Amaryl 6 mg tablete	NL/H/0101/005	H/97/00159/049	SANOFI-AVENTIS GROUPE	SI
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	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
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg Tabletten	NL/H/0101/004	1-21665	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 3 mg Tabletten	NL/H/0101/003	1-21666	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl, 4 mg tabletid	NL/H/0101/004	413703	SANOFI-AVENTIS GROUPE	EE
Amaryl, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
Amaryl, 3 mg tabletid	NL/H/0101/003	272999	SANOFI-AVENTIS GROUPE	EE
Amaryl® 4 mg Tabletten	NL/H/0101/004	32638.03.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Amaryl 2 mg Tabletten	NL/H/0101/002	1-21664	SANOFI-AVENTIS GMBH OSTERREICH	AT
Amaryl 4 mg, compresse	NL/H/0101/004	032845455	SANOFI S.R.L.	IT
Amaryl 2 mg, compresse	NL/H/0101/002	032845113	SANOFI 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
Amaryl® 2 mg, δισκίο	NL/H/0101/002	20551	SANOFI-AVENTIS GROUPE	CY
Amaryl® 4 mg, δισκίο	NL/H/0101/004	20553	SANOFI-AVENTIS GROUPE	CY
Amaryl® 3 mg, δισκίο	NL/H/0101/003	20552	SANOFI-AVENTIS GROUPE	CY
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 2 mg tablety	NL/H/0101/002	18/0482/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 4 mg tablety	NL/H/0101/004	18/0484/09-S	SANOFI-AVENTIS GROUPE	SK
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg, tabletten	NL/H/0101/001	RVG 17843	GENZYME EUROPE B.V.	NL
Amaryl 1 mg tabletės	NL/H/0101/001	LT/1/99/1288/011	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/003	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/044	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletės	NL/H/0101/004	LT/1/99/1288/040	SANOFI-AVENTIS GROUPE	LT
Amaryl 3 mg tabletės	NL/H/0101/003	LT/1/99/1288/029	SANOFI-AVENTIS GROUPE	LT
Amaryl 4 mg tabletes	NL/H/0101/004	09-0324	SANOFI-AVENTIS GROUPE	LV
Amaryl 2 mg tablety	NL/H/0101/002	18/233/97-C	SANOFI-AVENTIS SRO	CZ
Amaryl 2 mg tabletter	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletti	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 1 mg tabletti	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 2 mg tabletti	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 2 mg tabletti	NL/H/0101/002	12715	SANOFI OY	FI
Amaryl 1 mg tabletti	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 1 mg tabletti	NL/H/0101/001	12714	SANOFI OY	FI
Amaryl 4 mg tabletti	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletti	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletti	NL/H/0101/004	17571	SANOFI OY	FI
Amaryl 4 mg tabletter	NL/H/0101/004	17571	SANOFI OY	FI