EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

31 March 2016
EMA/333388/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: everolimus (indicated for rejection of transplanted organs)

Procedure no.: PSUSA/00010269/201507

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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
Certican	SE/H/0356/001	35644	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/002	35665	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/003	35666	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/004	35667	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/005	35668	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/006	35669	NOVARTIS HEALTHCARE A/S (ART57)	DK
Certican	SE/H/0356/004	088/04704	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
Certican	SE/H/0356/001	088/04701	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
Certican	SE/H/0356/002	088/04702	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
Certican	SE/H/0356/003	088/04703	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
Certican	SE/H/0356/005	088/04705	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
Certican	SE/H/0356/006	088/04706	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	MT
CERTICAN 0,75 mg comprimé	SE/H/0356/003	3400936410244	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,75 mg comprimé	SE/H/0356/003	3400936410305	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,75 mg comprimé	SE/H/0356/003	3400936410473	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,75 mg comprimé	SE/H/0356/003	3400956550425	NOVARTIS PHARMA S.A.S. (ART57)	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
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373177	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373189	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373191	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373203	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,1 mg comprimate pentru dispersie orala	SE/H/0356/005	7242/2014/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	SE/H/0356/005	7242/2014/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	SE/H/0356/005	7242/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	SE/H/0356/005	7242/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	not available	6072/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	not available	6072/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	not available	6072/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg comprimate pentru dispersie orala	not available	6072/2005/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,1 mg Comprimidos dispersíveis	SE/H/0356/005	5021688	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Certican 0,1 mg dispergeeruvad tabletid	SE/H/0356/005	464305	NOVARTIS FINLAND OY (ART57)	EE
Certican 0,1 mg disperģējamās tabletes	SE/H/0356/005	05-0030	NOVARTIS FINLAND OY (ART57)	LV
Certican 0,1 mg dispergerbara tabletter	SE/H/0356/005	18705	NOVARTIS FINLAND OY (ART57)	FI
Certican 0,1 mg dispergerbara tabletter	SE/H/0356/005	18694	NOVARTIS SVERIGE AB (ART57)	SE
Certican 0,1 mg dispergerbare tabletter	SE/H/0356/005	03-2079	NOVARTIS NORGE AS (ART57)	NO
Certican 0,1 mg dispergovatelne tablety	SE/H/0356/005	59/0296/04-S	NOVARTIS, S.R.O. (ART57)	SK
CERTICAN 0,1 mg dispergovatelné tablety	SE/H/0356/005	59/429/14-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/017	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/018	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/019	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/020	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,1 mg diszpergálódó tabletta	SE/H/0356/005	OGYI-T-9961/05	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 0,1 mg dreifitöflur	SE/H/0356/005	IS/1/03/040/05	NOVARTIS HEALTHCARE A/S (ART57)	IS
Certican 0,1 mg tabletki do sporządzania zawiesiny doustnej	SE/H/0356/005	R/22336	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 0,1 mg Tabletten zur Herstellung einer	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V. (ART57)	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Suspension zum Einnehmen				
Certican 0,1 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/005	58387.00.01	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 0,1 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0378/005	0010/04030109	NOVARTIS PHARMA N.V. (ART57)	LU
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411425	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411593	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411654	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400956550715	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,1 mg, comprimés dipsersibles	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,10 mg comprimés dispersibles	SE/H/0378/005	0010/04030109	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,10 mg dispergeerbare tabletten	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,1mg - Tabletten zur Herstellung einer Suspension	SE/H/0356/005	1-25275	NOVARTIS PHARMA GMBH (ART57)	AT
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373013	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373025	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373037	NOVARTIS FARMA S.P.A. (ART57)	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
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373049	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373215	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373227	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373239	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373241	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,25 mg comprimate	not available	6068/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate	not available	6068/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate	not available	6068/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate	not available	6068/2005/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate pentru dispersie orală	not available	6073/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate pentru dispersie orală	not available	6073/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate pentru dispersie orală	not available	6073/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate pentru dispersie orală	not available	6073/2005/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate/Certican 0,25 mg tablets	SE/H/0356/001	7238/2014/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate/Certican 0,25 mg tablets	SE/H/0356/001	7238/2014/02	NOVARTIS PHARMA GMBH (ART57)	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
Certican 0,25 mg comprimate/Certican 0,25 mg tablets	SE/H/0356/001	7238/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimate/Certican 0,25 mg tablets	SE/H/0356/001	7238/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25 mg comprimés	SE/H/0378/001	0010/04030105	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,25 mg comprimés dispersibles	SE/H/0356/006	0010/04030110	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,25 mg comprimidos dispersíveis	SE/H/0356/006	5021787	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Certican 0,25 mg dispergeerbare tabletten	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,25 mg dispergeeruvad tabletid	SE/H/0356/006	464205	NOVARTIS FINLAND OY (ART57)	EE
Certican 0,25 mg disperģējamās tabletes	SE/H/0356/006	05-0031	NOVARTIS FINLAND OY (ART57)	LV
Certican 0,25 mg dispergerbara tabletter	SE/H/0356/006	18704	NOVARTIS FINLAND OY (ART57)	FI
Certican 0,25 mg dispergerbara tabletter	SE/H/0356/006	18695	NOVARTIS SVERIGE AB (ART57)	SE
Certican 0,25 mg dispergerbare tabletter	SE/H/0356/006	03-2080	NOVARTIS NORGE AS (ART57)	NO
Certican 0,25 mg dispergovatelne tablety	SE/H/0356/006	59/0297/04-S	NOVARTIS, S.R.O. (ART57)	SK
CERTICAN 0,25 mg dispergovatelné tablety	SE/H/0356/006	59/430/14-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/021	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/022	NOVARTIS FINLAND OY (ART57)	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
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/023	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/024	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 mg diszpergálódó tabletta	SE/H/0356/006	OGYI-T-9961/06	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 0,25 mg dreifitöflur	SE/H/0356/006	IS/1/03/040/06	NOVARTIS HEALTHCARE A/S (ART57)	IS
Certican 0,25 mg tablete za oralnu suspenziju	SE/H/0356/006	HR-H-996683373	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 0,25 mg tabletes	SE/H/0356/001	05-0026	NOVARTIS FINLAND OY (ART57)	LV
Certican 0,25 mg tabletid	SE/H/0356/001	464705	NOVARTIS FINLAND OY (ART57)	EE
CERTICAN 0,25 MG TABLETKI	SE/H/0356/001	R/11215	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 0,25 mg tabletki do sporządzania zawiesiny doustnej	SE/H/0356/006	R/22337	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 0,25 mg tabletta	SE/H/0356/001	OGYI-T-9961/02	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 0,25 mg Tabletten	SE/H/0356/001	BE266445	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,25 mg tabletten	SE/H/0356/001	BE266445	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,25 mg Tabletten	SE/H/0356/001	58387.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 0,25 mg Tabletten	SE/H/0378/001	0010/04030105	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,25 mg Tabletten zur Herstellung einer Suspension	SE/H/0356/006	1-25276	NOVARTIS PHARMA GMBH (ART57)	AT
Certican 0,25 mg	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tabletten zur Herstellung einer Suspension zum Einnehmen			(ART57)	
Certican 0,25 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/006	58387.01.01	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 0,25 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/006	0010/04030110	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,25 mg tabletter	SE/H/0356/001	18698	NOVARTIS FINLAND OY (ART57)	FI
Certican 0,25 mg tabletter	SE/H/0356/001	03-2075	NOVARTIS NORGE AS (ART57)	NO
Certican 0,25 mg tabletter	SE/H/0356/001	18690	NOVARTIS SVERIGE AB (ART57)	SE
CERTICAN 0,25 mg TABLETY	SE/H/0356/001	59/014/05-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 0,25 mg tablety	SE/H/0356/001	59/0294/04-S	NOVARTIS, S.R.O. (ART57)	SK
Certican 0,25 mg töflur	SE/H/0356/001	IS/1/03/040/01	NOVARTIS HEALTHCARE A/S (ART57)	IS
Certican 0,25 mg, comprimé	SE/H/0356/001	3400936411074	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,25 mg, comprimé	SE/H/0356/001	3400936411135	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,25 mg, comprimé	SE/H/0356/001	3400936411364	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,25 mg, comprimé	SE/H/0356/001	3400956550654	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411715	NOVARTIS PHARMA S.A.S. (ART57)	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
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411883	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411944	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400956550883	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,25 mg, comprimés	SE/H/0356/001	BE266445	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,25 mg, comprimés dipsersibles	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,25 tabletės	SE/H/0356/001	LT/1/05/0321/001	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 tabletės	SE/H/0356/001	LT/1/05/0321/002	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 tabletės	SE/H/0356/001	LT/1/05/0321/003	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 tabletės	SE/H/0356/001	LT/1/05/0321/004	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,25 tabletten 0,25 mg	SE/H/0356/001	RVG 30041	NOVARTIS PHARMA B.V. (ART57)	NL
Certican 0,25mg comprimate pentru dispersie orala	SE/H/0356/006	7243/2014/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25mg comprimate pentru dispersie orala	SE/H/0356/006	7243/2014/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25mg comprimate pentru dispersie orala	SE/H/0356/006	7243/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25mg comprimate pentru dispersie orala	SE/H/0356/006	7243/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,25mg comprimidos	SE/H/0356/001	5021282	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Certican 0,25mg Tabletten	SE/H/0356/001	1-25271	NOVARTIS PHARMA GMBH (ART57)	AT
Certican 0,5 mg compresse	SE/H/0356/002	036373052	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373064	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373076	NOVARTIS FARMA S.P.A. (ART57)	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373088	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,5 mg comprimate	not available	6069/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate	not available	6069/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate	not available	6069/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate	not available	6069/2005/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate/certican 0,5 tablets	SE/H/0356/002	7239/2014/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate/certican 0,5 tablets	SE/H/0356/002	7239/2014/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate/certican 0,5 tablets	SE/H/0356/002	7239/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimate/certican 0,5 tablets	SE/H/0356/002	7239/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,5 mg comprimés	SE/H/0378/002	0010/04030106	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,5 mg Comprimidos	SE/H/0356/002	5021381	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Certican 0,5 mg tablete	not available	UP/I-530-09/11-02/196	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 0,5 mg tabletes	SE/H/0356/002	05-0027	NOVARTIS FINLAND OY (ART57)	LV
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/005	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/006	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/007	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/008	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,5 mg tabletid	SE/H/0356/002	464605	NOVARTIS FINLAND OY (ART57)	EE
CERTICAN 0,5 MG TABLETKI	SE/H/0356/002	R/11214	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 0,5 mg tabletta	SE/H/0356/002	OGYI-T-9961/01	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 0,5 mg Tabletten	SE/H/0356/002	BE266481	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,5 mg Tabletten	SE/H/0356/002	58387.01.00	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 0,5 mg Tabletten	SE/H/0378/002	0010/04030106	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,5 mg tabletter	SE/H/0356/002	18699	NOVARTIS FINLAND OY (ART57)	FI
Certican 0,5 mg tabletter	SE/H/0356/002	03-2076	NOVARTIS NORGE AS (ART57)	NO
Certican 0,5 mg tabletter	SE/H/0356/002	18691	NOVARTIS SVERIGE AB (ART57)	SE
Certican 0,5 mg tablety	SE/H/0356/002	59/428/14-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 0,5 mg tablety	SE/H/0356/002	59/0295/04-S	NOVARTIS, S.R.O. (ART57)	SK
Certican 0,5 mg töflur	SE/H/0356/002	IS/1/03/040/02	NOVARTIS HEALTHCARE A/S (ART57)	IS

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
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410763	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410824	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410992	NOVARTIS PHARMA S.A.S. (ART57)	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400956550593	NOVARTIS PHARMA S.A.S. (ART57)	FR
Certican 0,5 mg, comprimés	SE/H/0356/002	BE266481	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,5 tabletten 0,5 mg	SE/H/0356/002	RVG 30042	NOVARTIS PHARMA B.V. (ART57)	NL
Certican 0,50 mg - Tabletten	SE/H/0356/002	1-25272	NOVARTIS PHARMA GMBH (ART57)	AT
Certican 0,50 mg tabletten	SE/H/0356/002	BE266481	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,75 mg - Tabletten	SE/H/0356/003	1-25273	NOVARTIS PHARMA GMBH (ART57)	AT
Certican 0,75 mg compresse	SE/H/0356/003	036373090	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,75 mg compresse	SE/H/0356/003	036373102	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,75 mg compresse	SE/H/0356/003	036373114	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,75 mg compresse	SE/H/0356/003	036373126	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 0,75 mg comprimate	not available	6070/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate	not available	6070/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate	not available	6070/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate	not available	6070/2005/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate/Certican	SE/H/0356/003	7240/2014/01	NOVARTIS PHARMA GMBH (ART57)	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
0,75 mg tablets				
Certican 0,75 mg comprimate/Certican 0,75 mg tablets	SE/H/0356/003	7240/2014/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate/Certican 0,75 mg tablets	SE/H/0356/003	7240/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg comprimate/Certican 0,75 mg tablets	SE/H/0356/003	7240/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 0,75 mg Comprimidos	SE/H/0356/003	5021480	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Certican 0,75 mg tabletes	SE/H/0356/003	05-0028	NOVARTIS FINLAND OY (ART57)	LV
Certican 0,75 mg tabletid	SE/H/0356/003	464505	NOVARTIS FINLAND OY (ART57)	EE
CERTICAN 0,75 MG TABLETKI	SE/H/0356/003	R/11226	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 0,75 mg tabletta	SE/H/0356/003	OGYI-T-9961/03	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 0,75 mg Tabletten	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,75 mg tabletten	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,75 mg Tabletten	SE/H/0356/003	58387.02.00	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 0,75 mg Tabletten	SE/H/0378/003	0010/04030107	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 0,75 mg tabletten 0,75 mg	SE/H/0356/003	RVG 30043	NOVARTIS PHARMA B.V. (ART57)	NL
Certican 0,75 mg tabletter	SE/H/0356/003	18702	NOVARTIS FINLAND OY (ART57)	FI
Certican 0,75 mg tabletter	SE/H/0356/003	03-2077	NOVARTIS NORGE AS (ART57)	NO

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
Certican 0,75 mg tabletter	SE/H/0356/003	18692	NOVARTIS SVERIGE AB (ART57)	SE
CERTICAN 0,75 mg TABLETY	SE/H/0356/003	59/016/05-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 0,75 mg töflur	SE/H/0356/003	IS/1/03/040/03	NOVARTIS HEALTHCARE A/S (ART57)	IS
Certican 0,75 mg, comprimés	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 0,75 tabletės	SE/H/0356/003	LT/1/05/0321/009	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,75 tabletės	SE/H/0356/003	LT/1/05/0321/010	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,75 tabletės	SE/H/0356/003	LT/1/05/0321/011	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,75 tabletės	SE/H/0356/003	LT/1/05/0321/012	NOVARTIS FINLAND OY (ART57)	LT
Certican 0,75mg comprimés	SE/H/0378/003	0010/04030107	NOVARTIS PHARMA N.V. (ART57)	LU
CERTICAN 0.1 mg comprimidos dispersables	SE/H/0356/005	66009	NOVARTIS FARMACÉUTICA S.A. (ART57)	ES
Certican 0.1 mg dispergoituva tabletti	SE/H/0356/005	18705	NOVARTIS FINLAND OY (ART57)	FI
Certican 0.1 mg dispersible tablets	SE/H/0378/005	PA 13/124/005	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 0.1 mg dispersible tablets	SE/H/0356/005	PL 00101/0965	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 0.1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/001	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/002	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.1 mg tablete za oralnu suspenziju	SE/H/0356/005	HR-H-542458362	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
CERTICAN 0.25 mg	SE/H/0356/001	66001	NOVARTIS	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos			FARMACÉUTICA S.A. (ART57)	
CERTICAN 0.25 mg comprimidos dispersables	SE/H/0356/006	66002	NOVARTIS FARMACÉUTICA S.A. (ART57)	ES
Certican 0.25 mg dispergoituva tabletti	SE/H/0356/006	18704	NOVARTIS FINLAND OY (ART57)	FI
Certican 0.25 mg dispersible tablets	SE/H/0356/006	PA 13/124/006	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 0.25 mg dispersible tablets	SE/H/0356/006	PL 00101/0966	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 0.25 mg disperzibilne tablete	SE/H/0356/006	H/05/00370/005	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.25 mg disperzibilne tablete	SE/H/0356/006	~~5363-I-1011/09~~ H/05/00370/006	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.25 mg tablete	SE/H/0356/001	HR-H-809985014	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 0.25 mg tablete	SE/H/0356/001	H/05/00370/009	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.25 mg tablete	SE/H/0356/001	H/05/00370/010	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.25 mg tablets	SE/H/0378/001	PA 13/124/001	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 0.25 mg tablets	SE/H/0356/001	PL 00101/0961	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 0.25 mg tabletti	SE/H/0356/001	18698	NOVARTIS FINLAND OY (ART57)	FI
Certican 0.5 mg tablete	SE/H/0356/002	HR-H-592546271	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 0.5 mg tablete	SE/H/0356/002	H/05/00370/013	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.5 mg tablete	SE/H/0356/002	H/05/00370/014	NOVARTIS PHARMA	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
			GMBH (ART57)	
Certican 0.5 mg tablets	SE/H/0378/002	PA 13/124/002	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 0.5 mg tablets	SE/H/0356/002	PL 00101/0962	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 0.5 mg tabletti	SE/H/0356/002	18699	NOVARTIS FINLAND OY (ART57)	FI
CERTICAN 0.50 mg comprimidos	SE/H/0356/002	66006	NOVARTIS FARMACÉUTICA S.A. (ART57)	ES
CERTICAN 0.75 mg comprimidos	SE/H/0356/003	66007	NOVARTIS FARMACÉUTICA S.A. (ART57)	ES
Certican 0.75 mg tablete	SE/H/0356/003	HR-H-941540378	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 0.75 mg tablete	SE/H/0356/003	H/05/00370/017	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.75 mg tablete	SE/H/0356/003	H/05/00370/018	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 0.75 mg tablets	SE/H/0378/003	PA 13/124/003	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 0.75 mg tablets	SE/H/0356/003	PL 00101/0963	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 0.75 mg tabletti	SE/H/0356/003	18702	NOVARTIS FINLAND OY (ART57)	FI
Certican 1 mg comprimate	not available	6071/2005/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimate	not available	6071/2005/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimate	not available	6071/2005/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg	not available	6071/2005/04	NOVARTIS PHARMA	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
comprimate			GMBH (ART57)	
Certican 1 mg comprimate/certican 1 mg tablets	SE/H/0356/004	7241/2014/01	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimate/certican 1 mg tablets	SE/H/0356/004	7241/2014/02	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimate/certican 1 mg tablets	SE/H/0356/004	7241/2014/03	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimate/certican 1 mg tablets	SE/H/0356/004	7241/2014/04	NOVARTIS PHARMA GMBH (ART57)	RO
Certican 1 mg comprimés	SE/H/0378/004	0010/04030108	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 1 mg Comprimidos	SE/H/0356/004	5021589	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Certican 1 mg tabletes	SE/H/0356/004	05-0029	NOVARTIS FINLAND OY (ART57)	LV
Certican 1 mg tabletės	SE/H/0356/004	LT/1/05/0321/013	NOVARTIS FINLAND OY (ART57)	LT
Certican 1 mg tabletės	SE/H/0356/004	LT/1/05/0321/014	NOVARTIS FINLAND OY (ART57)	LT
Certican 1 mg tabletės	SE/H/0356/004	LT/1/05/0321/015	NOVARTIS FINLAND OY (ART57)	LT
Certican 1 mg tabletės	SE/H/0356/004	LT/1/05/0321/016	NOVARTIS FINLAND OY (ART57)	LT
Certican 1 mg tabletten	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 1 mg tabletter	SE/H/0356/004	18693	NOVARTIS SVERIGE AB (ART57)	SE
CERTICAN 1 mg TABLETY	SE/H/0356/004	59/017/05-C	NOVARTIS, S.R.O. (ART57)	CZ
Certican 1,0 mg	SE/H/0356/004	036373138	NOVARTIS FARMA 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
compresse			(ART57)	
Certican 1,0 mg compresse	SE/H/0356/004	036373140	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 1,0 mg compresse	SE/H/0356/004	036373153	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 1,0 mg compresse	SE/H/0356/004	036373165	NOVARTIS FARMA S.P.A. (ART57)	IT
Certican 1,0 mg tabletid	SE/H/0356/004	464405	NOVARTIS FINLAND OY (ART57)	EE
Certican 1,0 mg tabletta	SE/H/0356/004	OGYI-T-9961/04	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Certican 1,0 mg Tabletten	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 1,0 mg Tabletten	SE/H/0356/004	58387.03.00	NOVARTIS PHARMA GMBH (ART57)	DE
Certican 1,0 mg Tabletten	SE/H/0378/004	0010/04030108	NOVARTIS PHARMA N.V. (ART57)	LU
Certican 1,0 mg tabletter	SE/H/0356/004	18703	NOVARTIS FINLAND OY (ART57)	FI
Certican 1,0 mg tabletter	SE/H/0356/004	03-2078	NOVARTIS NORGE AS (ART57)	NO
Certican 1,0 mg tablety	SE/H/0356/004	59/0299/04-S	NOVARTIS, S.R.O. (ART57)	SK
Certican 1,0 mg töflur.	SE/H/0356/004	IS/1/03/040/04	NOVARTIS HEALTHCARE A/S (ART57)	IS
Certican 1,0 mg, comprimés	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V. (ART57)	BE
Certican 1,0 mg, tabletki	SE/H/0356/004	R/22335	NOVARTIS POLAND SP. Z O. O. (ART57)	PL
Certican 1,0 tabletten 1,0 mg	SE/H/0356/004	RVG 30044	NOVARTIS PHARMA B.V. (ART57)	NL
Certican 1.0 mg tabletti	SE/H/0356/004	18703	NOVARTIS FINLAND OY (ART57)	FI
CERTICAN 1.0 mg comprimidos	SE/H/0356/004	66008	NOVARTIS FARMACÉUTICA S.A. (ART57)	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Certican 1.0 mg tablete	SE/H/0356/004	HR-H-694937531	NOVARTIS HRVATSKA D.O.O. (ART57)	HR
Certican 1.0 mg tablete	SE/H/0356/004	H/05/00370/021	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 1.0 mg tablete	SE/H/0356/004	H/05/00370/022	NOVARTIS PHARMA GMBH (ART57)	SI
Certican 1.0 mg tablets	SE/H/0356/004	PA 13/124/004	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	IE
Certican 1.0mg tablets	SE/H/0356/004	PL 00101/0964	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	UK
Certican 1mg - Tabletten	SE/H/0356/004	1-25274	NOVARTIS PHARMA GMBH (ART57)	AT
Certican Dispers 0,1, dispergeerbare tabletten 0,1 mg	SE/H/0356/005	RVG 30045	NOVARTIS PHARMA B.V. (ART57)	NL
Certican Dispers 0,25, dispergeerbare tabletten 0,25 mg	SE/H/0356/006	RVG 30046	NOVARTIS PHARMA B.V. (ART57)	NL
Certican διασπειρόμενα δισκία 0.1mg	SE/H/0356/005	19646	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	CY
Certican διασπειρόμενα δισκία 0.25mg	SE/H/0356/006	19647	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	CY
Certican δισκία 0.75mg	SE/H/0356/003	19644	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	CY
Certican δισκία 0,25mg	SE/H/0356/001	19642	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	CY
Certican δισκία 0.5mg	SE/H/0356/002	19643	NOVARTIS PHARMACEUTICALS UK LIMITED (ART57)	CY
Certican δισκία 1mg	SE/H/0356/004	19645	NOVARTIS	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
			PHARMACEUTICALS UK LIMITED (ART57)	
Certican, διασπειρόμενο δισκίο 0.1mg	SE/H/0378/005	261570501	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.1mg	SE/H/0378/005	261570502	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.1mg	SE/H/0378/005	261570503	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.1mg	SE/H/0378/005	261570504	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.25mg	SE/H/0356/006	261570601	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.25mg	SE/H/0356/006	261570602	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.25mg	SE/H/0356/006	261570603	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, διασπειρόμενο δισκίο 0.25mg	SE/H/0356/006	261570604	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.25mg	SE/H/0378/001	261570101	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.25mg	SE/H/0378/001	261570102	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.25mg	SE/H/0378/001	261570103	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.25mg	SE/H/0378/001	261570104	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.5mg	SE/H/0378/002	261570201	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.5mg	SE/H/0378/002	261570202	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.5mg	SE/H/0378/002	261570203	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.5mg	SE/H/0378/002	261570204	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.75mg	SE/H/0378/003	261570301	NOVARTIS (HELLAS) S.A.C.I. (ART57)	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
Certican, δισκίο 0.75mg	SE/H/0378/003	261570302	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.75mg	SE/H/0378/003	261570303	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 0.75mg	SE/H/0378/003	261570304	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 1mg	SE/H/0378/004	261570401	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 1mg	SE/H/0378/004	261570402	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 1mg	SE/H/0378/004	261570403	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican, δισκίο 1mg	SE/H/0378/004	261570404	NOVARTIS (HELLAS) S.A.C.I. (ART57)	GR
Certican0,75 mg tbl	SE/H/0356/003	59/0298/04-S	NOVARTIS, S.R.O. (ART57)	SK
Everolimus Novartis 0,1 mg dispergerbara tabletter	SE/H/378/005	19846	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,25 mg dispergerbara tabletter	SE/H/378/006	19847	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,25 mg tabletter	SE/H/378/001	19842	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,5 mg tabletter	SE/H/378/002	19843	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,75 mg tabletter	SE/H/378/003	19844	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 1 mg tabletter	SE/H/378/004	19845	NOVARTIS SVERIGE AB	SE