23 May 2019
EMA/281903/20199
Human Medicines Evaluation Division

List of nationally authorised medicinal products

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

Procedure no.: PSUSA/00010269/201807

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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 0,1 mg dispergovatelné tablety	SE/H/0356/005	59/429/14-C	NOVARTIS S.R.O.,	CZ
Certican 0,1 mg – Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/005	1-25275	NOVARTIS PHARMA GMBH	AT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373177	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373189	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373191	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,1 mg compresse dispersibili	SE/H/0356/005	036373203	NOVARTIS FARMA S.P.A.	IT
Certican 0,1 mg comprimate pentru dispersie orală	SE/H/0356/005	9629/2017/02	NOVARTIS PHARMA GMBH	RO
Certican 0,1 mg comprimate pentru dispersie orală	SE/H/0356/005	9629/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 0,1 mg comprimate pentru dispersie orală	SE/H/0356/005	9629/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 0,1 mg comprimate pentru dispersie orală	SE/H/0356/005	9629/2017/01	NOVARTIS PHARMA GMBH	RO
Certican 0,1 mg comprimés dipsersibles	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V.	BE
Certican 0,1 mg comprimés dispersibles	SE/H/0356/005	0010/04030109	NOVARTIS PHARMA N.V.	LU
CERTICAN 0,1 mg comprimidos dispersables	SE/H/0356/005	66.009	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 0,1 mg Comprimidos dispersíveis	SE/H/0356/005	5021688	NOVARTIS FARMA - PRODUTOS	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
			FARMACÊUTICOS S.A.	
CERTICAN 0,1 MG DISPERGEERUVAD TABLETID	SE/H/0356/005	464305	SIA "NOVARTIS BALTICS"	EE
Certican 0,1 mg disperģējamās tabletes	SE/H/0356/005	05-0030	SIA "NOVARTIS BALTICS"	LV
Certican 0,1 mg dispergerbara tabletter	SE/H/0356/005	18705	NOVARTIS FINLAND OY	FI
Certican 0,1 mg dispergerbara tabletter	SE/H/0356/005	18694	NOVARTIS SVERIGE AB	SE
Certican 0,1 mg dispergoituva tabletti	SE/H/0356/005	18705	NOVARTIS FINLAND OY	FI
Certican 0,1 mg dispergovatelne tablety	SE/H/0356/005	59/0296/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/017	SIA "NOVARTIS BALTICS"	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/018	SIA "NOVARTIS BALTICS"	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/019	SIA "NOVARTIS BALTICS"	LT
Certican 0,1 mg disperguojamosios tabletės	SE/H/0356/005	LT/1/05/0321/020	SIA "NOVARTIS BALTICS"	LT
Certican 0,1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/001	NOVARTIS PHARMA GMBH	SI
Certican 0,1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/002	NOVARTIS PHARMA GMBH	SI
Certican 0,1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/003	NOVARTIS PHARMA GMBH	SI
Certican 0,1 mg disperzibilne tablete	SE/H/0356/005	H/05/00370/004	NOVARTIS PHARMA GMBH	SI
Certican 0,1 mg	SE/H/0356/005	OGYI-T-9961/05	NOVARTIS HUNGÁRIA KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
diszpergálódó tabletta			PHARMA	
Certican 0,1 mg dreifitöflur	SE/H/0356/005	IS/1/03/040/05	NOVARTIS HEALTHCARE A/S	IS
Certican 0,1 mg tablete za oralnu suspenziju	SE/H/0356/005	HR-H-542458362	NOVARTIS HRVATSKA D.O.O.	HR
Certican 0,1 mg tabletki do sporządzania zawiesiny doustnej	SE/H/0356/005	22336	NOVARTIS POLAND SP. Z O. O.	PL
Certican 0,1 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V.	BE
Certican 0,1 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/005	0010/04030109	NOVARTIS PHARMA N.V.	LU
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411425	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411593	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400936411654	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,1 mg, comprimé dispersible	SE/H/0356/005	3400956550715	NOVARTIS PHARMA S.A.S.	FR
Certican 0,10 mg dispergeerbare tabletten	SE/H/0356/005	BE266454	NOVARTIS PHARMA N.V.	BE
Certican 0,25 mg – Tabletten	SE/H/0356/001	1-25271	NOVARTIS PHARMA GMBH	AT
Certican 0,25 mg – Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/006	1-25276	NOVARTIS PHARMA GMBH	AT
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373013	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg	SE/H/0356/001	036373025	NOVARTIS FARMA S.P.A.	IT

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compresse				
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373037	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg compresse	SE/H/0356/001	036373049	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373215	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373227	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373239	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,25 mg compresse dispersibili	SE/H/0356/006	036373241	NOVARTIS FARMA S.P.A.	IT
Certican 0,25 mg comprimate	SE/H/0356/001	9625/2017/01	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate	SE/H/0356/001	9625/2017/02	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate	SE/H/0356/001	9625/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate	SE/H/0356/001	9625/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate pentru dispersie orală	SE/H/0356/006	9630/2017/01	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate pentru dispersie orală	SE/H/0356/006	9630/2017/02	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate pentru dispersie orală	SE/H/0356/006	9630/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimate pentru dispersie orală	SE/H/0356/006	9630/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 0,25 mg comprimés	SE/H/0356/001	BE266445	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
Certican 0,25 mg comprimés	SE/H/0356/001	0010/04030105	NOVARTIS PHARMA N.V.	LU
Certican 0,25 mg comprimés dispersibles	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V.	BE
Certican 0,25 mg comprimés dispersibles	SE/H/0356/006	0010/04030110	NOVARTIS PHARMA N.V.	LU
CERTICAN 0,25 mg comprimidos	SE/H/0356/001	66.001	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 0,25 mg comprimidos	SE/H/0356/001	5021282	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
CERTICAN 0,25 mg comprimidos dispersables	SE/H/0356/006	66.002	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 0,25 mg comprimidos dispersíveis	SE/H/0356/006	5021787	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Certican 0,25 mg dispergeerbare tabletten	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V.	BE
CERTICAN 0,25 MG DISPERGEERUVAD TABLETID	SE/H/0356/006	464205	SIA "NOVARTIS BALTICS"	EE
Certican 0,25 mg disperģējamās tabletes	SE/H/0356/006	05-0031	SIA "NOVARTIS BALTICS"	LV
Certican 0,25 mg dispergerbara tabletter	SE/H/0356/006	18704	NOVARTIS FINLAND OY	FI
Certican 0,25 mg dispergerbara tabletter	SE/H/0356/006	18695	NOVARTIS SVERIGE AB	SE
Certican 0,25 mg dispergoituva tabletti	SE/H/0356/006	18704	NOVARTIS FINLAND OY	FI
Certican 0,25 mg dispergovatelne tablety	SE/H/0356/006	59/0297/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 0,25 mg dispergovatelné tablety	SE/H/0356/006	59/430/14-C	NOVARTIS S.R.O.,	CZ
Certican 0,25 mg	SE/H/0356/006	LT/1/05/0321/021	SIA "NOVARTIS BALTICS"	LT

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disperguojamosios tabletės				
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/022	SIA "NOVARTIS BALTICS"	LT
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/023	SIA "NOVARTIS BALTICS"	LT
Certican 0,25 mg disperguojamosios tabletės	SE/H/0356/006	LT/1/05/0321/024	SIA "NOVARTIS BALTICS"	LT
Certican 0,25 mg disperzibilne tablete	SE/H/0356/006	H/05/00370/005	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg disperzibilne tablete	SE/H/0356/006	H/05/00370/006	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg disperzibilne tablete	SE/H/0356/006	H/05/00370/007	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg disperzibilne tablete	SE/H/0356/006	H/05/00370/008	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg diszpergálódó tabletta	SE/H/0356/006	OGYI-T-9961/06	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Certican 0,25 mg dreifitöflur	SE/H/0356/006	IS/1/03/040/06	NOVARTIS HEALTHCARE A/S	IS
Certican 0,25 mg tablete	SE/H/0356/001	HR-H-809985014-02	NOVARTIS HRVATSKA D.O.O.	HR
Certican 0,25 mg tablete	SE/H/0356/001	H/05/00370/009	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg tablete	SE/H/0356/001	H/05/00370/010	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg tablete	SE/H/0356/001	H/05/00370/011	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg tablete	SE/H/0356/001	H/05/00370/012	NOVARTIS PHARMA GMBH	SI
Certican 0,25 mg tablete za oralnu suspenziju	SE/H/0356/006	HR-H-996683373	NOVARTIS HRVATSKA D.O.O.	HR
Certican 0,25 mg tabletes	SE/H/0356/001	05-0026	SIA "NOVARTIS BALTICS"	LV
CERTICAN 0,25 MG	SE/H/0356/001	LT/1/05/0321/001	SIA "NOVARTIS BALTICS"	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
TABLETĖS				
CERTICAN 0,25 MG TABLETĖS	SE/H/0356/001	LT/1/05/0321/002	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,25 MG TABLETĖS	SE/H/0356/001	LT/1/05/0321/003	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,25 MG TABLETĖS	SE/H/0356/001	LT/1/05/0321/004	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,25 MG TABLETID	SE/H/0356/001	464705	SIA "NOVARTIS BALTICS"	EE
Certican 0,25 mg tabletki	SE/H/0356/001	11215	NOVARTIS POLAND SP. Z O. O.	PL
Certican 0,25 mg tabletki do sporządzania zawiesiny doustnej	SE/H/0356/006	22337	NOVARTIS POLAND SP. Z O. O.	PL
Certican 0,25 mg tabletta	SE/H/0356/001	OGYI-T-9961/02	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Certican 0,25 mg Tabletten	SE/H/0356/001	BE266445	NOVARTIS PHARMA N.V.	BE
Certican 0,25 mg tabletten	SE/H/0356/001	BE266445	NOVARTIS PHARMA N.V.	BE
Certican 0,25 mg Tabletten	SE/H/0356/001	0010/04030105	NOVARTIS PHARMA N.V.	LU
Certican 0,25 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/006	BE266497	NOVARTIS PHARMA N.V.	BE
Certican 0,25 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	SE/H/0356/006	0010/04030110	NOVARTIS PHARMA N.V.	LU
Certican 0,25 mg tabletter	SE/H/0356/001	18698	NOVARTIS FINLAND OY	FI
Certican 0,25 mg tabletter	SE/H/0356/001	18690	NOVARTIS SVERIGE AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Certican 0,25 mg tabletti	SE/H/0356/001	18698	NOVARTIS FINLAND OY	FI
Certican 0,25 mg tablety	SE/H/0356/001	59/014/05-C	NOVARTIS S.R.O.,	CZ
Certican 0,25 mg tablety	SE/H/0356/001	59/0294/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 0,25 mg töflur	SE/H/0356/001	IS/1/03/040/01	NOVARTIS HEALTHCARE A/S	IS
CERTICAN 0,25 mg, comprimé	SE/H/0356/001	3400936411074	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé	SE/H/0356/001	3400936411135	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé	SE/H/0356/001	3400936411364	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé	SE/H/0356/001	3400956550654	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411715	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411883	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400936411944	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,25 mg, comprimé dispersible	SE/H/0356/006	3400956550883	NOVARTIS PHARMA S.A.S.	FR
Certican 0,25, tabletten 0,25 mg	SE/H/0356/001	RVG 30041	NOVARTIS PHARMA B.V.	NL
Certican 0,5 mg – Tabletten	SE/H/0356/002	1-25272	NOVARTIS PHARMA GMBH	AT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373052	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373064	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373076	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,5 mg compresse	SE/H/0356/002	036373088	NOVARTIS FARMA S.P.A.	IT
Certican 0,5 mg comprimate	SE/H/0356/002	9626/2017/01	NOVARTIS PHARMA GMBH	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 0,5 mg comprimate	SE/H/0356/002	9626/2017/02	NOVARTIS PHARMA GMBH	RO
Certican 0,5 mg comprimate	SE/H/0356/002	9626/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 0,5 mg comprimate	SE/H/0356/002	9626/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 0,5 mg comprimés	SE/H/0356/002	BE266481	NOVARTIS PHARMA N.V.	BE
Certican 0,5 mg comprimés	SE/H/0356/002	0010/04030106	NOVARTIS PHARMA N.V.	LU
CERTICAN 0,5 mg comprimidos	SE/H/0356/002	66.006	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 0,5 mg Comprimidos	SE/H/0356/002	5021381	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Certican 0,5 mg tablete	SE/H/0356/002	HR-H-592546271-02	NOVARTIS HRVATSKA D.O.O.	HR
Certican 0,5 mg tablete	SE/H/0356/002	H/05/00370/013	NOVARTIS PHARMA GMBH	SI
Certican 0,5 mg tablete	SE/H/0356/002	H/05/00370/014	NOVARTIS PHARMA GMBH	SI
Certican 0,5 mg tablete	SE/H/0356/002	H/05/00370/015	NOVARTIS PHARMA GMBH	SI
Certican 0,5 mg tablete	SE/H/0356/002	H/05/00370/016	NOVARTIS PHARMA GMBH	SI
Certican 0,5 mg tabletes	SE/H/0356/002	05-0027	SIA "NOVARTIS BALTICS"	LV
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/005	SIA "NOVARTIS BALTICS"	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/006	SIA "NOVARTIS BALTICS"	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/007	SIA "NOVARTIS BALTICS"	LT
Certican 0,5 mg tabletės	SE/H/0356/002	LT/1/05/0321/008	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,5 MG TABLETID	SE/H/0356/002	464605	SIA "NOVARTIS BALTICS"	EE
Certican 0,5 mg tabletki	SE/H/0356/002	11214	NOVARTIS POLAND SP. Z O. O.	PL
Certican 0,5 mg tabletta	SE/H/0356/002	OGYI-T-9961/01	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Certican 0,5 mg Tabletten	SE/H/0356/002	BE266481	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
Certican 0,5 mg Tabletten	SE/H/0356/002	0010/04030106	NOVARTIS PHARMA N.V.	LU
Certican 0,5 mg tabletter	SE/H/0356/002	18699	NOVARTIS FINLAND OY	FI
Certican 0,5 mg tabletter	SE/H/0356/002	18691	NOVARTIS SVERIGE AB	SE
Certican 0,5 mg tabletti	SE/H/0356/002	18699	NOVARTIS FINLAND OY	FI
Certican 0,5 mg tablety	SE/H/0356/002	59/428/14-C	NOVARTIS S.R.O.,	CZ
Certican 0,5 mg tablety	SE/H/0356/002	59/0295/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 0,5 mg töflur	SE/H/0356/002	IS/1/03/040/02	NOVARTIS HEALTHCARE A/S	IS
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410763	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410824	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400936410992	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,5 mg, comprimé	SE/H/0356/002	3400956550593	NOVARTIS PHARMA S.A.S.	FR
Certican 0,5, tabletten 0,5 mg	SE/H/0356/002	RVG 30042	NOVARTIS PHARMA B.V.	NL
Certican 0,50 mg tabletten	SE/H/0356/002	BE266481	NOVARTIS PHARMA N.V.	BE
Certican 0,75 mg – Tabletten	SE/H/0356/003	1-25273	NOVARTIS PHARMA GMBH	AT
CERTICAN 0,75 mg compresse	SE/H/0356/003	036373090	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,75 mg compresse	SE/H/0356/003	036373102	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,75 mg compresse	SE/H/0356/003	036373114	NOVARTIS FARMA S.P.A.	IT
CERTICAN 0,75 mg compresse	SE/H/0356/003	036373126	NOVARTIS FARMA S.P.A.	IT
Certican 0,75 mg comprimate	SE/H/0356/003	9627/2017/01	NOVARTIS PHARMA GMBH	RO
Certican 0,75 mg comprimate	SE/H/0356/003	9627/2017/02	NOVARTIS PHARMA GMBH	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 0,75 mg comprimate	SE/H/0356/003	9627/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 0,75 mg comprimate	SE/H/0356/003	9627/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 0,75 mg comprimés	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V.	BE
Certican 0,75 mg comprimés	SE/H/0356/003	0010/04030107	NOVARTIS PHARMA N.V.	LU
CERTICAN 0,75 mg comprimidos	SE/H/0356/003	66.007	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 0,75 mg Comprimidos	SE/H/0356/003	5021480	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Certican 0,75 mg tablete	SE/H/0356/003	HR-H-941540378	NOVARTIS HRVATSKA D.O.O.	HR
Certican 0,75 mg tablete	SE/H/0356/003	H/05/00370/017	NOVARTIS PHARMA GMBH	SI
Certican 0,75 mg tablete	SE/H/0356/003	H/05/00370/018	NOVARTIS PHARMA GMBH	SI
Certican 0,75 mg tablete	SE/H/0356/003	H/05/00370/019	NOVARTIS PHARMA GMBH	SI
Certican 0,75 mg tablete	SE/H/0356/003	H/05/00370/020	NOVARTIS PHARMA GMBH	SI
Certican 0,75 mg tabletes	SE/H/0356/003	05-0028	SIA "NOVARTIS BALTICS"	LV
CERTICAN 0,75 MG TABLETĖS	SE/H/0356/003	LT/1/05/0321/009	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,75 MG TABLETĖS	SE/H/0356/003	LT/1/05/0321/010	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,75 MG TABLETĖS	SE/H/0356/003	LT/1/05/0321/011	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,75 MG TABLETĖS	SE/H/0356/003	LT/1/05/0321/012	SIA "NOVARTIS BALTICS"	LT
CERTICAN 0,75 MG TABLETID	SE/H/0356/003	464505	SIA "NOVARTIS BALTICS"	EE
Certican 0,75 mg tabletki	SE/H/0356/003	11226	NOVARTIS POLAND SP. Z O. O.	PL
Certican 0,75 mg	SE/H/0356/003	OGYI-T-9961/03	NOVARTIS HUNGÁRIA KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletta			PHARMA	
Certican 0,75 mg Tabletten	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V.	BE
Certican 0,75 mg tabletten	SE/H/0356/003	BE266472	NOVARTIS PHARMA N.V.	BE
Certican 0,75 mg Tabletten	SE/H/0356/003	0010/04030107	NOVARTIS PHARMA N.V.	LU
Certican 0,75 mg tabletter	SE/H/0356/003	18702	NOVARTIS FINLAND OY	FI
Certican 0,75 mg tabletter	SE/H/0356/003	18692	NOVARTIS SVERIGE AB	SE
Certican 0,75 mg tabletti	SE/H/0356/003	18702	NOVARTIS FINLAND OY	FI
Certican 0,75 mg tablety	SE/H/0356/003	59/016/05-C	NOVARTIS S.R.O.,	CZ
Certican 0,75 mg tablety	SE/H/0356/003	59/0298/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 0,75 mg töflur	SE/H/0356/003	IS/1/03/040/03	NOVARTIS HEALTHCARE A/S	IS
CERTICAN 0,75 mg, comprimé	SE/H/0356/003	3400936410244	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,75 mg, comprimé	SE/H/0356/003	3400936410305	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,75 mg, comprimé	SE/H/0356/003	3400936410473	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 0,75 mg, comprimé	SE/H/0356/003	3400956550425	NOVARTIS PHARMA S.A.S.	FR
Certican 0,75, tabletten 0,75 mg	SE/H/0356/003	RVG 30043	NOVARTIS PHARMA B.V.	NL
CERTICAN 0.1 MG DISPERSIBLE TABLETS	SE/H/0356/005	MA1249/00405	NOVARTIS IRELAND LIMITED	MT
CERTICAN 0.25 MG DISPERSIBLE TABLETS	SE/H/0356/006	MA1249/00406	NOVARTIS IRELAND LIMITED	MT
CERTICAN 0.25 MG TABLETS	SE/H/0356/001	MA1249/00401	NOVARTIS IRELAND LIMITED	MT
CERTICAN 0.5 MG TABLETS	SE/H/0356/002	MA1249/00402	NOVARTIS IRELAND LIMITED	MT
CERTICAN 0.75 MG	SE/H/0356/003	MA1249/00403	NOVARTIS IRELAND	MT

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
TABLETS			LIMITED	
Certican 1 mg – Tabletten	SE/H/0356/004	1-25274	NOVARTIS PHARMA GMBH	AT
Certican 1 mg comprimate	SE/H/0356/004	9628/2017/01	NOVARTIS PHARMA GMBH	RO
Certican 1 mg comprimate	SE/H/0356/004	9628/2017/02	NOVARTIS PHARMA GMBH	RO
Certican 1 mg comprimate	SE/H/0356/004	9628/2017/03	NOVARTIS PHARMA GMBH	RO
Certican 1 mg comprimate	SE/H/0356/004	9628/2017/04	NOVARTIS PHARMA GMBH	RO
Certican 1 mg tablety	SE/H/0356/004	59/017/05-C	NOVARTIS S.R.O.,	CZ
Certican 1 mg tablety	SE/H/0356/004	59/0299/04-S	NOVARTIS SLOVAKIA S.R.O.	SK
Certican 1,0 mg tabletti	SE/H/0356/004	18703	NOVARTIS FINLAND OY	FI
CERTICAN 1,0 mg compresse	SE/H/0356/004	036373138	NOVARTIS FARMA S.P.A.	IT
CERTICAN 1,0 mg compresse	SE/H/0356/004	036373140	NOVARTIS FARMA S.P.A.	IT
CERTICAN 1,0 mg compresse	SE/H/0356/004	036373153	NOVARTIS FARMA S.P.A.	IT
CERTICAN 1,0 mg compresse	SE/H/0356/004	036373165	NOVARTIS FARMA S.P.A.	IT
Certican 1,0 mg comprimés	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V.	BE
Certican 1,0 mg comprimés	SE/H/0356/004	0010/04030108	NOVARTIS PHARMA N.V.	LU
CERTICAN 1,0 mg comprimidos	SE/H/0356/004	66.008	NOVARTIS FARMACÉUTICA S.A.	ES
Certican 1,0 mg Comprimidos	SE/H/0356/004	5021589	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Certican 1,0 mg tablete	SE/H/0356/004	HR-H-694937531	NOVARTIS HRVATSKA D.O.O.	HR
Certican 1,0 mg tablete	SE/H/0356/004	H/05/00370/021	NOVARTIS PHARMA GMBH	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
Certican 1,0 mg tablete	SE/H/0356/004	H/05/00370/022	NOVARTIS PHARMA GMBH	SI
Certican 1,0 mg tablete	SE/H/0356/004	H/05/00370/023	NOVARTIS PHARMA GMBH	SI
Certican 1,0 mg tablete	SE/H/0356/004	H/05/00370/024	NOVARTIS PHARMA GMBH	SI
Certican 1,0 mg tabletes	SE/H/0356/004	05-0029	SIA "NOVARTIS BALTICS"	LV
Certican 1,0 mg tabletės	SE/H/0356/004	LT/1/05/0321/013	SIA "NOVARTIS BALTICS"	LT
Certican 1,0 mg tabletės	SE/H/0356/004	LT/1/05/0321/014	SIA "NOVARTIS BALTICS"	LT
Certican 1,0 mg tabletės	SE/H/0356/004	LT/1/05/0321/015	SIA "NOVARTIS BALTICS"	LT
Certican 1,0 mg tabletės	SE/H/0356/004	LT/1/05/0321/016	SIA "NOVARTIS BALTICS"	LT
CERTICAN 1,0 MG TABLETID	SE/H/0356/004	464405	SIA "NOVARTIS BALTICS"	EE
Certican 1,0 mg tabletki	SE/H/0356/004	22335	NOVARTIS POLAND SP. Z O. O.	PL
Certican 1,0 mg tabletta	SE/H/0356/004	OGYI-T-9961/04	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Certican 1,0 mg Tabletten	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V.	BE
Certican 1,0 mg tabletten	SE/H/0356/004	BE266463	NOVARTIS PHARMA N.V.	BE
Certican 1,0 mg Tabletten	SE/H/0356/004	0010/04030108	NOVARTIS PHARMA N.V.	LU
Certican 1,0 mg tabletter	SE/H/0356/004	18703	NOVARTIS FINLAND OY	FI
Certican 1,0 mg tabletter	SE/H/0356/004	18693	NOVARTIS SVERIGE AB	SE
Certican 1,0 mg töflur	SE/H/0356/004	IS/1/03/040/04	NOVARTIS HEALTHCARE A/S	IS
CERTICAN 1,0 mg, comprimé	SE/H/0356/004	3400930136669	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 1,0 mg, comprimé	SE/H/0356/004	3400930136676	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 1,0 mg, comprimé	SE/H/0356/004	3400955043621	NOVARTIS PHARMA S.A.S.	FR
CERTICAN 1,0 mg, comprimé	SE/H/0356/004	3400955043614	NOVARTIS PHARMA S.A.S.	FR
Certican 1,0, tabletten 1,0 mg	SE/H/0356/004	RVG 30044	NOVARTIS PHARMA B.V.	NL
CERTICAN 1.0 MG	SE/H/0356/004	MA1249/00404	NOVARTIS IRELAND	MT

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
TABLETS			LIMITED	
Certican Dispers 0,1, dispergeerbare tabletten 0,1 mg	SE/H/0356/005	RVG 30045	NOVARTIS PHARMA B.V.	NL
Certican Dispers 0,25, dispergeerbare tabletten 0,25 mg	SE/H/0356/006	RVG 30046	NOVARTIS PHARMA B.V.	NL
Certican, dispergible tabletter	SE/H/0356/005	35668	NOVARTIS HEALTHCARE A/S	DK
Certican, dispergible tabletter	SE/H/0356/006	35669	NOVARTIS HEALTHCARE A/S	DK
Certican, tabletter	SE/H/0356/001	35644	NOVARTIS HEALTHCARE A/S	DK
Certican, tabletter	SE/H/0356/002	35665	NOVARTIS HEALTHCARE A/S	DK
Certican, tabletter	SE/H/0356/003	35666	NOVARTIS HEALTHCARE A/S	DK
Certican, tabletter	SE/H/0356/004	35667	NOVARTIS HEALTHCARE A/S	DK
Certican® 0,1 mg dispergerbare tabletter	SE/H/0356/005	03-2079	NOVARTIS NORGE AS	NO
CERTICAN® 0,1 MG TABLETTEN ZUR HERSTELLUNG EINER SUSPENSION ZUM EINNEHMEN	SE/H/0356/005	58387.00.01	NOVARTIS PHARMA GMBH	DE
Certican® 0,1mg διασπειρόμενα δισκία	SE/H/0356/005	261570501	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,1mg διασπειρόμενα δισκία	SE/H/0356/005	261570502	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,1mg διασπειρόμενα δισκία	SE/H/0356/005	261570503	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,1mg διασπειρόμενα δισκία	SE/H/0356/005	261570504	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg dispergerbare tabletter	SE/H/0356/006	03-2080	NOVARTIS NORGE AS	NO
CERTICAN® 0,25 MG TABLETTEN	SE/H/0356/001	58387.00.00	NOVARTIS PHARMA 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
CERTICAN® 0,25 MG TABLETTEN ZUR HERSTELLUNG EINER SUSPENSION ZUM EINNEHMEN	SE/H/0356/006	58387.01.01	NOVARTIS PHARMA GMBH	DE
Certican® 0,25 mg tabletter	SE/H/0356/001	03-2075	NOVARTIS NORGE AS	NO
Certican® 0,25 mg διασπειρόμενα δισκία	SE/H/0356/006	261570601	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg διασπειρόμενα δισκία	SE/H/0356/006	261570602	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg διασπειρόμενα δισκία	SE/H/0356/006	261570603	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg διασπειρόμενα δισκία	SE/H/0356/006	261570604	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg δισκία	SE/H/0356/001	261570101	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg δισκία	SE/H/0356/001	261570102	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg δισκία	SE/H/0356/001	261570103	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,25 mg δισκία	SE/H/0356/001	261570104	NOVARTIS (HELLAS) S.A.C.I.	GR
CERTICAN® 0,5 MG TABLETTEN	SE/H/0356/002	58387.01.00	NOVARTIS PHARMA GMBH	DE
Certican® 0,5 mg tabletter	SE/H/0356/002	03-2076	NOVARTIS NORGE AS	NO
Certican® 0,5 mg δισκία	SE/H/0356/002	261570201	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,5 mg δισκία	SE/H/0356/002	261570202	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,5 mg δισκία	SE/H/0356/002	261570203	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,5 mg δισκία	SE/H/0356/002	261570204	NOVARTIS (HELLAS) S.A.C.I.	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® 0,75 MG TABLETTEN	SE/H/0356/003	58387.02.00	NOVARTIS PHARMA GMBH	DE
Certican® 0,75 mg tabletter	SE/H/0356/003	03-2077	NOVARTIS NORGE AS	NO
Certican® 0,75 mg δισκία	SE/H/0356/003	261570301	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,75 mg δισκία	SE/H/0356/003	261570302	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,75 mg δισκία	SE/H/0356/003	261570303	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0,75 mg δισκία	SE/H/0356/003	261570304	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 0.1 mg dispersible tablets	SE/H/0356/005	PL 00101/0965	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Certican® 0.1mg διασπειρόμενα δισκία	SE/H/0356/005	19646	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Certican® 0.1mg διασπειρόμενα δισκία	SE/H/0356/006	19647	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Certican® 0.25 mg dispersible tablets	SE/H/0356/006	PL 00101/0966	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Certican® 0.25 mg tablets	SE/H/0356/001	PL 00101/0961	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Certican® 0.25 mg δισκία	SE/H/0356/001	19642	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Certican® 0.5 mg tablets	SE/H/0356/002	PL 00101/0962	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Certican® 0.5 mg δισκία	SE/H/0356/002	19643	NOVARTIS PHARMACEUTICALS UK	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
			LIMITED	
Certican® 0.75 mg tablets	SE/H/0356/003	PL 00101/0963	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Certican® 0.75 mg δισκία	SE/H/0356/003	19644	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Certican® 1 mg tablets	SE/H/0356/004	PL 00101/0964	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
CERTICAN® 1,0 MG TABLETTEN	SE/H/0356/004	58387.03.00	NOVARTIS PHARMA GMBH	DE
Certican® 1,0 mg tabletter	SE/H/0356/004	03-2078	NOVARTIS NORGE AS	NO
Certican® 1,0 mg δισκία	SE/H/0356/004	261570401	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 1,0 mg δισκία	SE/H/0356/004	261570402	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 1,0 mg δισκία	SE/H/0356/004	261570403	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 1,0 mg δισκία	SE/H/0356/004	261570404	NOVARTIS (HELLAS) S.A.C.I.	GR
Certican® 1.0 mg δισκία	SE/H/0356/004	19645	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Everolimus Novartis 0,1 mg dispergerbara tabletter	SE/H/0378/005	19846	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,1 mg dispergerbara tabletter	SE/H/0378/005	19846	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,25 mg dispergerbara tabletter	SE/H/0378/006	19847	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,25	SE/H/0378/006	19847	NOVARTIS SVERIGE AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg dispergerbara tabletter				
Everolimus Novartis 0,25 mg tabletter	SE/H/0378/001	19842	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,25 mg tabletter	SE/H/0378/001	19842	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,5 mg tabletter	SE/H/0378/002	19843	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,5 mg tabletter	SE/H/0378/002	19843	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,75 mg tableter	SE/H/0378/003	19844	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 0,75 mg tableter	SE/H/0378/003	19844	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 1 mg tabletter	SE/H/0378/004	19845	NOVARTIS SVERIGE AB	SE
Everolimus Novartis 1 mg tabletter	SE/H/0378/004	19845	NOVARTIS SVERIGE AB	SE
Сертикан 0,1 mg диспергиращи се таблетки	SE/H/0356/005	20050109	NOVARTIS PHARMA GMBH	BG
Сертикан 0,25 mg диспергиращи се таблетки	SE/H/0356/006	20050108	NOVARTIS PHARMA GMBH	BG
Сертикан 0,25 mg таблетки	SE/H/0356/001	20050110	NOVARTIS PHARMA GMBH	BG
Сертикан 0,5 mg таблетки	SE/H/0356/002	20050111	NOVARTIS PHARMA GMBH	BG
Сертикан 0,75 mg таблетки	SE/H/0356/003	20050112	NOVARTIS PHARMA GMBH	BG
Сертикан 1,0 mg таблетки	SE/H/0356/004	20050113	NOVARTIS PHARMA GMBH	BG