9 April 2021
EMA/PRAC/222202/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: oxcarbazepine

Procedure no.: PSUSA/00002235/202008

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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
Apydan extent 150 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65225.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Apydan extent 300 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65226.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Apydan extent 600 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65227.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Timox 60 mg/ml Suspension zum Einnehmen	DE/H/4778/004	52125.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Timox extent 150 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65230.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Timox extent 300 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65231.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Timox extent 600 mg Tabletten mit veränderter Wirkstofffreisetzung	not available	65232.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Tolep 300 mg compresse	not available	028304018	NOVARTIS FARMA S.P.A.	IT
Tolep 300 mg compresse	not available	028304018	NOVARTIS FARMA S.P.A.	IT
Tolep 600 mg compresse	not available	028304020	NOVARTIS FARMA S.P.A.	IT
Tolep 600 mg compresse	not available	028304020	NOVARTIS FARMA S.P.A.	IT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127081	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127180	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127289	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127388	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127487	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127081	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127180	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127289	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127388	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg comprimidos revestidos por película	DK/H/0168/001	3127487	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 150 mg Film-coated tablets	not available	19109	NOVARTIS IRELAND LIMITED	CY
Trileptal 150 mg Film-coated tablets	not available	19109	NOVARTIS IRELAND LIMITED	CY
Trileptal 150 mg Film-coated tablets	DK/H/0168/001	PA0896/033/001	NOVARTIS IRELAND LIMITED	IE
Trileptal 150 mg Film-	DK/H/0168/001	PA0896/033/001	NOVARTIS IRELAND	IE

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
coated tablets			LIMITED	
Trileptal 150 mg filmdragerade tabletter	DK/H/0168/001	15781	NOVARTIS SVERIGE AB	SE
Trileptal 150 mg filmdragerade tabletter	DK/H/0168/001	15781	NOVARTIS SVERIGE AB	SE
Trileptal 150 mg filmdrasjerte tabletter	not available	99-315	NOVARTIS NORGE AS	NO
Trileptal 150 mg filmdrasjerte tabletter	not available	99-315	NOVARTIS NORGE AS	NO
Trileptal 150 mg filmomhulde tabletten	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg filmomhulde tabletten	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg Filmtabletten	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg Filmtabletten	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg filmuhúðaðar töflur	DK/H/0168/001	IS/1/00/004/01	NOVARTIS HEALTHCARE A/S	IS
Trileptal 150 mg filmuhúðaðar töflur	DK/H/0168/001	IS/1/00/004/01	NOVARTIS HEALTHCARE A/S	IS
Trileptal 150 mg plėvele dengtos tabletės	not available	LT/1/97/1473/001	SIA NOVARTIS BALTICS	LT
Trileptal 150 mg plėvele dengtos tabletės	not available	LT/1/97/1473/001	SIA NOVARTIS BALTICS	LT
Trileptal 150 mg tablett, filmdragerad	DK/H/0168/001	14797	NOVARTIS FINLAND OY	FI
Trileptal 150 mg tablett, filmdragerad	DK/H/0168/001	14797	NOVARTIS FINLAND OY	FI
Trileptal 150 mg tabletti, kalvopäällysteinen	DK/H/0168/001	14797	NOVARTIS FINLAND OY	FI
Trileptal 150 mg tabletti, kalvopäällysteinen	DK/H/0168/001	14797	NOVARTIS FINLAND OY	FI
Trileptal 150 mg	DK/H/0168/001	198880301	NOVARTIS (HELLAS)	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
Επικαλυμένα με λεπτό υμένιο δισκία			S.A.C.I.	
Trileptal 150 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/001	198880302	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 150 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/001	198880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 150 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/001	198880302	NOVARTIS (HELLAS) S.A.C.I.	GR
TRILEPTAL 150 mg, comprimé pelliculé	DK/H/0168/001	34009 353 570 1 4	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 150 mg, comprimé pelliculé	DK/H/0168/001	34009 353 571 8 2	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 150 mg, comprimé pelliculé	DK/H/0168/001	34009 353 570 1 4	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 150 mg, comprimé pelliculé	DK/H/0168/001	34009 353 571 8 2	NOVARTIS PHARMA S.A.S.	FR
Trileptal 150 mg, comprimés pelliculés	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg, comprimés pelliculés	DK/H/0168/001	BE208993	NOVARTIS PHARMA N.V.	BE
Trileptal 150 mg, filmomhulde tabletten	DK/H/0168/001	RVG 24750	NOVARTIS PHARMA B.V.	NL
Trileptal 150 mg, filmomhulde tabletten	DK/H/0168/001	RVG 24750	NOVARTIS PHARMA B.V.	NL
Trileptal 300 mg apvalkotas tabletes	not available	99-0279	SIA NOVARTIS BALTICS	LV
Trileptal 300 mg apvalkotas tabletes	not available	99-0279	SIA NOVARTIS BALTICS	LV
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/01	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg	not available	9041/2016/02	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
comprimate filmate				
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/03	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/04	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/05	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/01	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/02	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/03	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/04	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 300 mg comprimate filmate	not available	9041/2016/05	NOVARTIS PHARMA GMBH	RO
Trileptal 300 mg comprimidos recubiertos con película.	DK/H/0168/002	63.093	NOVARTIS FARMACÉUTICA S.A.	ES
Trileptal 300 mg comprimidos recubiertos con película.	DK/H/0168/002	63.093	NOVARTIS FARMACÉUTICA S.A.	ES
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127586	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127685	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127784	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos	DK/H/0168/002	3127883	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
por película			FARMACÊUTICOS S.A.	
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127982	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127586	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127685	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127784	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127883	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg comprimidos revestidos por película	DK/H/0168/002	3127982	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 300 mg Film-coated tablets	not available	18463	NOVARTIS IRELAND LIMITED	CY
Trileptal 300 mg Film-coated tablets	not available	18463	NOVARTIS IRELAND LIMITED	CY
Trileptal 300 mg Film-coated tablets	DK/H/0168/002	PA0896/033/002	NOVARTIS IRELAND LIMITED	IE
Trileptal 300 mg Film-coated tablets	DK/H/0168/002	PA0896/033/002	NOVARTIS IRELAND LIMITED	IE
Trileptal 300 mg filmdragerade tabletter	DK/H/0168/002	15782	NOVARTIS SVERIGE AB	SE
Trileptal 300 mg filmdragerade tabletter	DK/H/0168/002	15782	NOVARTIS SVERIGE AB	SE
Trileptal 300 mg filmdrasjerte tabletter	not available	99-316	NOVARTIS NORGE AS	NO
Trileptal 300 mg	not available	99-316	NOVARTIS NORGE AS	NO

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filmdrasjerte tabletter				
Trileptal 300 mg filmom obalené tablety	not available	21/0336/00-S	NOVARTIS SLOVAKIA S.R.O.	SK
Trileptal 300 mg filmom obalené tablety	not available	21/0336/00-S	NOVARTIS SLOVAKIA S.R.O.	SK
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-02	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-03	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-04	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-05	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-02	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-03	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-04	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-05	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmom obložene tablete	DK/H/0168/002	HR-H-309957904-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 300 mg filmomhulde tabletten	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 300 mg filmomhulde tabletten	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 300 mg filmomhulde tabletten	DK/H/0168/002	RVG 24751	NOVARTIS PHARMA B.V.	NL
Trileptal 300 mg filmomhulde tabletten	DK/H/0168/002	RVG 24751	NOVARTIS PHARMA B.V.	NL
Trileptal 300 mg	not available	OGYI-T-6308/01	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
filmtabletta				
Trileptal 300 mg filmtabletta	not available	OGYI-T-6308/02	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 300 mg filmtabletta	not available	OGYI-T-6308/01	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 300 mg filmtabletta	not available	OGYI-T-6308/02	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 300 mg Filmtabletten	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 300 mg Filmtabletten	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 300 mg filmuhúðaðar töflur	DK/H/0168/002	IS/1/00/004/02	NOVARTIS HEALTHCARE A/S	IS
Trileptal 300 mg filmuhúðaðar töflur	DK/H/0168/002	IS/1/00/004/02	NOVARTIS HEALTHCARE A/S	IS
Trileptal 300 mg plėvele dengtos tabletės	not available	LT/1/97/1473/002	SIA NOVARTIS BALTICS	LT
Trileptal 300 mg plėvele dengtos tabletės	not available	LT/1/97/1473/002	SIA NOVARTIS BALTICS	LT
Trileptal 300 mg tabletki powlekane	not available	8256	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 300 mg tabletki powlekane	not available	8256	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 300 mg tablett, filmdragerad	DK/H/0168/002	14798	NOVARTIS FINLAND OY	FI
Trileptal 300 mg tablett, filmdragerad	DK/H/0168/002	14798	NOVARTIS FINLAND OY	FI
Trileptal 300 mg tabletti, kalvopäällysteinen	DK/H/0168/002	14798	NOVARTIS FINLAND OY	FI
Trileptal 300 mg tabletti, kalvopäällysteinen	DK/H/0168/002	14798	NOVARTIS FINLAND OY	FI
Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880401	NOVARTIS (HELLAS) S.A.C.I.	GR

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Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880402	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880403	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880402	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 300 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/002	198880403	NOVARTIS (HELLAS) S.A.C.I.	GR
TRILEPTAL 300 mg, comprimé pelliculé	DK/H/0168/002	3400935357243	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 300 mg, comprimé pelliculé	DK/H/0168/002	3400935357304	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 300 mg, comprimé pelliculé	DK/H/0168/002	3400935357243	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 300 mg, comprimé pelliculé	DK/H/0168/002	3400935357304	NOVARTIS PHARMA S.A.S.	FR
Trileptal 300 mg, comprimés pelliculés	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 300 mg, comprimés pelliculés	DK/H/0168/002	BE209002	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml mikstur, suspensjon	not available	00-8131	NOVARTIS NORGE AS	NO
Trileptal 60 mg/ml mikstur, suspensjon	not available	00-8131	NOVARTIS NORGE AS	NO
Trileptal 60 mg/ml mixtúra, dreifa	DK/H/0168/004	IS/1/01/039/01	NOVARTIS HEALTHCARE A/S	IS
Trileptal 60 mg/ml	DK/H/0168/004	IS/1/01/039/01	NOVARTIS HEALTHCARE A/S	IS

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mixtúra, dreifa				
Trileptal 60 mg/ml oraalisuspensio	DK/H/0168/004	16570	NOVARTIS FINLAND OY	FI
Trileptal 60 mg/ml oraalisuspensio	DK/H/0168/004	16570	NOVARTIS FINLAND OY	FI
Trileptal 60 mg/ml Oral Suspension	not available	19494	NOVARTIS IRELAND LIMITED	CY
Trileptal 60 mg/ml Oral Suspension	not available	19494	NOVARTIS IRELAND LIMITED	CY
Trileptal 60 mg/ml oral suspension	DK/H/0168/004	16570	NOVARTIS FINLAND OY	FI
Trileptal 60 mg/ml oral suspension	DK/H/0168/004	16570	NOVARTIS FINLAND OY	FI
Trileptal 60 mg/ml oral suspension	DK/H/0168/004	17348	NOVARTIS SVERIGE AB	SE
Trileptal 60 mg/ml oral suspension	DK/H/0168/004	17348	NOVARTIS SVERIGE AB	SE
Trileptal 60 mg/ml Oral Suspension.	DK/H/0168/004	PA0896/033/004	NOVARTIS IRELAND LIMITED	IE
Trileptal 60 mg/ml Oral Suspension.	DK/H/0168/004	PA0896/033/004	NOVARTIS IRELAND LIMITED	IE
Trileptal 60 mg/ml oralna suspenzija	DK/H/0168/004	HR-H-061545230-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 60 mg/ml oralna suspenzija	DK/H/0168/004	HR-H-061545230-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 60 mg/ml Suspensão oral	DK/H/0168/004	3732989	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 60 mg/ml Suspensão oral	DK/H/0168/004	3732989	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 60 mg/ml suspensión oral	DK/H/0168/004	64.398	NOVARTIS FARMACÉUTICA S.A.	ES
Trileptal 60 mg/ml	DK/H/0168/004	64.398	NOVARTIS FARMACÉUTICA	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
suspensión oral			S.A.	
Trileptal 60 mg/ml Suspension zum Einnehmen	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml Suspension zum Einnehmen	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml zawiesina doustna	not available	7471	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 60 mg/ml zawiesina doustna	not available	7471	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 60 mg/ml Πόσιμο εναιώρημα	DK/H/0168/004	198880601	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 60 mg/ml Πόσιμο εναιώρημα	DK/H/0168/004	198880601	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 60 mg/ml, suspensie voor oraal gebruik	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml, suspensie voor oraal gebruik	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml, suspensie voor oraal gebruik	DK/H/0168/004	RVG 26830	NOVARTIS PHARMA B.V.	NL
Trileptal 60 mg/ml, suspensie voor oraal gebruik	DK/H/0168/004	RVG 26830	NOVARTIS PHARMA B.V.	NL
Trileptal 60 mg/ml, suspension buvable	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
Trileptal 60 mg/ml, suspension buvable	DK/H/0168/004	BE227342	NOVARTIS PHARMA N.V.	BE
TRILEPTAL 60 mg/ml, suspension buvable	DK/H/0168/004	3400935790125	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 60 mg/ml,	DK/H/0168/004	3400935790125	NOVARTIS PHARMA S.A.S.	FR

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suspension buvable				
Trileptal 600 mg apvalkotas tabletes	not available	99-0280	SIA NOVARTIS BALTICS	LV
Trileptal 600 mg apvalkotas tabletes	not available	99-0280	SIA NOVARTIS BALTICS	LV
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/01	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/02	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/03	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/04	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/05	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/01	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/02	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/03	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/04	NOVARTIS PHARMA GMBH	RO
TRILEPTAL 600 mg comprimate filmate	not available	9042/2016/05	NOVARTIS PHARMA GMBH	RO
Trileptal 600 mg comprimidos recubiertos con película	DK/H/0168/003	63.095	NOVARTIS FARMACÉUTICA S.A.	ES
Trileptal 600 mg comprimidos recubiertos con película	DK/H/0168/003	63.095	NOVARTIS FARMACÉUTICA S.A.	ES
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128089	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128188	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128287	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128386	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128485	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128089	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128188	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128287	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128386	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg comprimidos revestidos por película	DK/H/0168/003	3128485	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Trileptal 600 mg Film-coated tablets	not available	18464	NOVARTIS IRELAND LIMITED	CY
Trileptal 600 mg Film-coated tablets	not available	18464	NOVARTIS IRELAND LIMITED	CY
Trileptal 600 mg Film-coated tablets	DK/H/0168/003	PA0896/033/003	NOVARTIS IRELAND LIMITED	IE
Trileptal 600 mg Film-	DK/H/0168/003	PA0896/033/003	NOVARTIS IRELAND	IE

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
coated tablets			LIMITED	
Trileptal 600 mg filmdragerade tabletter	DK/H/0168/003	15783	NOVARTIS SVERIGE AB	SE
Trileptal 600 mg filmdragerade tabletter	DK/H/0168/003	15783	NOVARTIS SVERIGE AB	SE
Trileptal 600 mg filmdrasjerte tabletter	not available	99-317	NOVARTIS NORGE AS	NO
Trileptal 600 mg filmdrasjerte tabletter	not available	99-317	NOVARTIS NORGE AS	NO
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-02	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-03	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-04	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-05	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-02	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-03	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-04	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-05	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmom obložene tablete	DK/H/0168/003	HR-H-007294503-01	NOVARTIS HRVATSKA D.O.O.	HR
Trileptal 600 mg filmomhulde tabletten	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal 600 mg filmomhulde tabletten	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal 600 mg	DK/H/0168/003	RVG 24752	NOVARTIS PHARMA 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
filmomhulde tabletten				
Trileptal 600 mg filmomhulde tabletten	DK/H/0168/003	RVG 24752	NOVARTIS PHARMA B.V.	NL
Trileptal 600 mg filmtabletta	not available	OGYI-T-6308/03	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 600 mg filmtabletta	not available	OGYI-T-6308/04	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 600 mg filmtabletta	not available	OGYI-T-6308/03	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 600 mg filmtabletta	not available	OGYI-T-6308/04	NOVARTIS HUNGÁRIA KFT.	HU
Trileptal 600 mg Filmtabletten	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal 600 mg Filmtabletten	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal 600 mg filmuhúðaðar töflur	DK/H/0168/003	IS/1/00/004/03	NOVARTIS HEALTHCARE A/S	IS
Trileptal 600 mg filmuhúðaðar töflur	DK/H/0168/003	IS/1/00/004/03	NOVARTIS HEALTHCARE A/S	IS
Trileptal 600 mg plėvele dengtos tabletės	not available	LT/1/97/1473/003	SIA NOVARTIS BALTICS	LT
Trileptal 600 mg plėvele dengtos tabletės	not available	LT/1/97/1473/003	SIA NOVARTIS BALTICS	LT
Trileptal 600 mg tabletki powlekane	not available	8257	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 600 mg tabletki powlekane	not available	8257	NOVARTIS POLAND SP. Z O. O.	PL
Trileptal 600 mg tablett, filmdragerad	DK/H/0168/003	14799	NOVARTIS FINLAND OY	FI
Trileptal 600 mg tablett, filmdragerad	DK/H/0168/003	14799	NOVARTIS FINLAND OY	FI
Trileptal 600 mg tabletti, kalvopäällysteinen	DK/H/0168/003	14799	NOVARTIS FINLAND OY	FI
Trileptal 600 mg tabletti,	DK/H/0168/003	14799	NOVARTIS FINLAND 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
kalvopäällysteinen				
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880502	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880503	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880502	NOVARTIS (HELLAS) S.A.C.I.	GR
Trileptal 600 mg Επικαλυμένα με λεπτό υμένιο δισκία	DK/H/0168/003	198880503	NOVARTIS (HELLAS) S.A.C.I.	GR
TRILEPTAL 600 mg, comprimé pelliculé	DK/H/0168/003	3400935357472	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 600 mg, comprimé pelliculé	DK/H/0168/003	3400935357533	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 600 mg, comprimé pelliculé	DK/H/0168/003	3400935357472	NOVARTIS PHARMA S.A.S.	FR
TRILEPTAL 600 mg, comprimé pelliculé	DK/H/0168/003	3400935357533	NOVARTIS PHARMA S.A.S.	FR
Trileptal 600 mg, comprimés pelliculés	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal 600 mg, comprimés pelliculés	DK/H/0168/003	BE209011	NOVARTIS PHARMA N.V.	BE
Trileptal, 150 mg õhukese polümeerikattega	not available	346401	SIA "NOVARTIS BALTICS"	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
tabletid				
Trileptal, 150 mg õhukese polümeerikattega tabletid	not available	346401	SIA "NOVARTIS BALTICS"	EE
Trileptal, 300 mg õhukese polümeerikattega tabletid	not available	346501	SIA "NOVARTIS BALTICS"	EE
Trileptal, 300 mg õhukese polümeerikattega tabletid	not available	346501	SIA "NOVARTIS BALTICS"	EE
Trileptal, 600 mg õhukese polümeerikattega tabletid	not available	346601	SIA "NOVARTIS BALTICS"	EE
Trileptal, 600 mg õhukese polümeerikattega tabletid	not available	346601	SIA "NOVARTIS BALTICS"	EE
Trileptal, filmovertrukne tabletter	DK/H/0168/001	30413	NOVARTIS HEALTHCARE A/S	DK
Trileptal, filmovertrukne tabletter	DK/H/0168/002	30414	NOVARTIS HEALTHCARE A/S	DK
Trileptal, filmovertrukne tabletter	DK/H/0168/003	30415	NOVARTIS HEALTHCARE A/S	DK
Trileptal, filmovertrukne tabletter	DK/H/0168/001	30413	NOVARTIS HEALTHCARE A/S	DK
Trileptal, filmovertrukne tabletter	DK/H/0168/002	30414	NOVARTIS HEALTHCARE A/S	DK
Trileptal, filmovertrukne tabletter	DK/H/0168/003	30415	NOVARTIS HEALTHCARE A/S	DK
Trileptal, oral suspension	DK/H/0168/004	31743	NOVARTIS HEALTHCARE A/S	DK

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
Trileptal, oral suspension	DK/H/0168/004	31743	NOVARTIS HEALTHCARE A/S	DK
Trileptal® 150 mg – Filmtabletten	DK/H/0168/001	1-23489	NOVARTIS PHARMA GMBH	AT
Trileptal® 150 mg – Filmtabletten	DK/H/0168/001	1-23489	NOVARTIS PHARMA GMBH	AT
Trileptal® 150 mg Film-coated Tablets	DK/H/0168/001	PL 00101/0581	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 150 mg Film-coated Tablets	DK/H/0168/001	PL 00101/0581	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 150 mg Filmtabletten	DK/H/0168/001	47357.00.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 150 mg Filmtabletten	DK/H/0168/001	47357.00.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 300 mg – Filmtabletten	DK/H/0168/002	1-23490	NOVARTIS PHARMA GMBH	AT
Trileptal® 300 mg – Filmtabletten	DK/H/0168/002	1-23490	NOVARTIS PHARMA GMBH	AT
Trileptal® 300 mg Film-coated Tablets	DK/H/0168/002	PL 00101/0582	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 300 mg Film-coated Tablets	DK/H/0168/002	PL 00101/0582	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 300 mg Filmtabletten	DK/H/0168/002	47357.01.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 300 mg Filmtabletten	DK/H/0168/002	47357.01.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 60 mg/ml - Suspension zum Einnehmen	DK/H/0168/004	1-24351	NOVARTIS PHARMA GMBH	AT
Trileptal® 60 mg/ml - Suspension zum	DK/H/0168/004	1-24351	NOVARTIS PHARMA 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
Einnehmen				
Trileptal® 60 mg/ml Oral Suspension	DK/H/0168/004	PL 00101/0631	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 60 mg/ml Oral Suspension	DK/H/0168/004	PL 00101/0631	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 60 mg/ml Suspension zum Einnehmen	DK/H/0168/004	51794.00.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 60 mg/ml Suspension zum Einnehmen	DK/H/0168/004	51794.00.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 600 mg – Filmtabletten	DK/H/0168/003	1-23491	NOVARTIS PHARMA GMBH	AT
Trileptal® 600 mg – Filmtabletten	DK/H/0168/003	1-23491	NOVARTIS PHARMA GMBH	AT
Trileptal® 600 mg Film-coated Tablets	DK/H/0168/003	PL 00101/0583	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 600 mg Film-coated Tablets	DK/H/0168/003	PL 00101/0583	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Trileptal® 600 mg Filmtabletten	DK/H/0168/003	47357.02.00	NOVARTIS PHARMA GMBH	DE
Trileptal® 600 mg Filmtabletten	DK/H/0168/003	47357.02.00	NOVARTIS PHARMA GMBH	DE
Трилептал 150 mg филмирани таблетки	not available	20010441	NOVARTIS PHARMA GMBH	BG
Трилептал 150 mg филмирани таблетки	not available	20010441	NOVARTIS PHARMA GMBH	BG
Трилептал 300 mg филмирани таблетки	not available	20010387	NOVARTIS PHARMA GMBH	BG
Трилептал 300 mg	not available	20010387	NOVARTIS PHARMA GMBH	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
филмирани таблетки				
Трилептал 60 mg/ml перорална суспензия	not available	20020486	NOVARTIS PHARMA GMBH	BG
Трилептал 60 mg/ml перорална суспензия	not available	20020486	NOVARTIS PHARMA GMBH	BG
Трилептал 600 mg филмирани таблетки	not available	20010386	NOVARTIS PHARMA GMBH	BG
Трилептал 600 mg филмирани таблетки	not available	20010386	NOVARTIS PHARMA GMBH	BG