15 January 2026
EMADOC-1700519818-2873421
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): valsartan, hydrochlorothiazide / valsartan

EURD list No. PSUSA/00010396/202504

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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Co-Vals 160 mg/12,5 mg comprimidos recubiertos con película	not available	65.797	ESTEVE PHARMACEUTICALS, S.A.	ES
Co-Vals 160 mg/12,5 mg comprimidos recubiertos con película	not available	65.797	ESTEVE PHARMACEUTICALS, S.A.	ES
AuroValsart HCT, 80 mg + 12,5 mg, tabletki powlekane	PT/H/1594/001/DC	23887	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
AuroValsart HCT, 160 mg + 12,5 mg, tabletki powlekane	PT/H/1594/002/DC	23888	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
AuroValsart HCT, 320 mg + 12,5 mg, tabletki powlekane	PT/H/1594/004	23890	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
Valsartan/Hydrochlorothiazide 160 mg/12.5 mg Film-coated Tablets	not available	PL 36687/0381	TORRENT PHARMA (UK) LTD.	XI
Valsartan/Hydrochlorothiazide 160 mg/25 mg Film-coated Tablets	not available	PL 36687/0382	TORRENT PHARMA (UK) LTD.	XI
Vals 80 mg comprimidos recubiertos con película.	not available	64.495	ESTEVE PHARMACEUTICALS, S.A.	ES
Vals 160 mg comprimidos recubiertos con película.	not available	64.494	ESTEVE PHARMACEUTICALS, S.A.	ES
Diovan 40 mg comprimidos revestidos por película	SE/H/0406/005	SE/H/0406/005	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 40 mg comprimidos revestidos por película	SE/H/0406/005	SE/H/0406/005	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 40 mg comprimidos revestidos	SE/H/0406/005	SE/H/0406/005	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.	
Diovan 40 mg comprimidos revestidos por película	SE/H/0406/005	SE/H/0406/005	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 40 mg comprimidos revestidos por película	SE/H/0406/005	SE/H/0406/005	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	SE/H/0406/004	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos	SE/H/0406/003	SE/H/0406/003	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.	
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	SE/H/0406/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	SE/H/0406/004	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	SE/H/0406/004	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	SE/H/0406/004	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	SE/H/0406/004	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	SE/H/0406/003	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	SE/H/0406/003	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	SE/H/0406/003	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	SE/H/0406/003	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	SE/H/0406/003	NOVARTIS PHARMA S.A.S.	FR
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	SE/H/0406/005	NOVARTIS PHARMA S.A.S.	FR
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	SE/H/0406/005	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	SE/H/0406/004	NOVARTIS PHARMA S.A.S.	FR
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV

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
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	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
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 160 mg filmtabletta	SE/H/0406/004	SE/H/0406/004	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan 80 mg filmtabletta	SE/H/0406/003	SE/H/0406/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovane 3 mg/ml solution buvable	SE/H/0406/007	2010060100	NOVARTIS PHARMA N.V.	LU
Diovane 80 mg comprimés pelliculés	SE/H/0406/003	2001120025	NOVARTIS PHARMA N.V.	LU
Diovane 160 mg comprimés pelliculés	SE/H/0406/004	2001120026	NOVARTIS PHARMA N.V.	LU
Diovane 320 mg comprimés pelliculés	SE/H/0406/006	2007040038	NOVARTIS PHARMA N.V.	LU
Diovane 160 mg Filmtabletten	SE/H/0406/004	2001120026	NOVARTIS PHARMA N.V.	LU
Diovane 3 mg/ml Lösung zum Einnehmen	SE/H/0406/007	2010060100	NOVARTIS PHARMA N.V.	LU
Diovane 320 mg Filmtabletten	SE/H/0406/006	2007040038	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Diovane 80 mg Filmtabletten	SE/H/0406/003	2001120025	NOVARTIS PHARMA N.V.	LU
Diovane 160 mg Filmtabletten	SE/H/0406/004	BE296423	NOVARTIS PHARMA N.V.	BE
Diovane 3 mg/ml Lösung zum Einnehmen	SE/H/0406/007	BE371007	NOVARTIS PHARMA N.V.	BE
Diovane 320 mg Filmtabletten	SE/H/0406/006	BE297604	NOVARTIS PHARMA N.V.	BE
Diovane 80 mg Filmtabletten	SE/H/0406/003	BE296414	NOVARTIS PHARMA N.V.	BE
Diovane 80 mg comprimés pelliculés	SE/H/0406/003	BE296414	NOVARTIS PHARMA N.V.	BE
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-02	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-03	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-04	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-06	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-07	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-08	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-09	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-10	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 80 mg filmom obložene tablete	SE/H/0406/003	HR-H-807143753-11	NOVARTIS HRVATSKA D.O.O.	HR
Diovan 3 mg/ml soluţie orală	SE/H/0406/007	9051/2016/01	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/066	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/068	NOVARTIS EUROPHARM LIMITED	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
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/069	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/080	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/081	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/082	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/083	NOVARTIS EUROPHARM LIMITED	SI
Diovan 320 mg filmsko obložene tablete	SE/H/0406/006	H/03/00476/096	NOVARTIS EUROPHARM LIMITED	SI
Diovan 320 mg filmsko obložene tablete	SE/H/0406/006	H/03/00476/099	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/006	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/019	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/020	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/021	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/023	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/036	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/038	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/039	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/049	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/050	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/051	NOVARTIS EUROPHARM LIMITED	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
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/053	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/054	NOVARTIS EUROPHARM LIMITED	SI
Diovan 40 mg filmsko obložene tablete	SE/H/0406/005	H/03/00476/001	NOVARTIS EUROPHARM LIMITED	SI
Diovan 160 mg filmsko obložene tablete	SE/H/0406/004	H/03/00476/061	NOVARTIS EUROPHARM LIMITED	SI
Diovan 3 mg/ml peroralna raztopina	SE/H/0406/007	H/03/00476/121	NOVARTIS EUROPHARM LIMITED	SI
Diovan 320 mg filmsko obložene tablete	SE/H/0406/006	H/03/00476/091	NOVARTIS EUROPHARM LIMITED	SI
Diovan 80 mg filmsko obložene tablete	SE/H/0406/003	H/03/00476/031	NOVARTIS EUROPHARM LIMITED	SI
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870414	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870415	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870416	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870417	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870425	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870426	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 160 mg	SE/H/0406/004	232870427	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με	SE/H/0406/004	232870428	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
λεπτό υμένιο δισκία 160 mg			S.A.C.I.	
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 320 mg	SE/H/0406/006	232870628	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 320 mg	SE/H/0406/006	232870629	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 320 mg	SE/H/0406/006	232870631	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870513	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870514	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870515	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870517	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870525	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 40 mg	SE/H/0406/005	232870526	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870313	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870314	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με	SE/H/0406/003	232870315	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
λεπτό υμένιο δισκία 80 mg			S.A.C.I.	
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870317	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870318	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870325	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870326	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870327	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan επικαλυμμένα με λεπτό υμένιο δισκία 80 mg	SE/H/0406/003	232870328	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan 3 mg/ml πόσιμο διάλυμα	SE/H/0406/007	232870701	NOVARTIS (HELLAS) S.A.C.I.	GR
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	3787884	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 160 mg comprimidos revestidos por película	SE/H/0406/004	3804184	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 3 mg/ml solução oral	SE/H/0406/007	5303441	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 40 mg comprimidos revestidos por película	SE/H/0406/005	5448980	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan 80 mg comprimidos revestidos	SE/H/0406/003	3787488	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.	
Diovan 80 mg comprimidos revestidos por película	SE/H/0406/003	3787587	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Tareg 3 mg/ml soluzione orale	SE/H/0406/007	033178423	NOVARTIS FARMA S.P.A.	IT
Tareg 320 mg compresse rivestite con film	SE/H/0406/006	033178346	NOVARTIS FARMA S.P.A.	IT
Tareg 320 mg compresse rivestite con film	SE/H/0406/006	033178359	NOVARTIS FARMA S.P.A.	IT
Tareg 320 mg compresse rivestite con film	SE/H/0406/006	033178373	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178260	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178272	NOVARTIS FARMA S.P.A.	IT
Diovan 160 mg apvalkotās tabletes	SE/H/0406/004	02-0392	SIA NOVARTIS BALTICS	LV
Diován 160 mg comprimidos recubiertos con película	SE/H/0406/004	64.473	NOVARTIS FARMACÉUTICA S.A.	ES
Diovan 160 mg filmdragerade tabletter	SE/H/0406/004	16709	NOVARTIS FINLAND OY	FI
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/011	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/044	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/045	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/046	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/048	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
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/049	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/050	SIA NOVARTIS BALTICS	LT
Diovan 160 mg plėvele dengtos tabletės	SE/H/0406/004	LT/1/01/1236/051	SIA NOVARTIS BALTICS	LT
Diovan 160 mg kalvopäällysteinen tabletti	SE/H/0406/004	16709	NOVARTIS FINLAND OY	FI
Diovan 3 mg/ml oraaliliuos	SE/H/0406/007	27966	NOVARTIS FINLAND OY	FI
Diovan 3 mg/ml šķīdums iekšķīgai lietošanai.	SE/H/0406/007	10-0320	SIA NOVARTIS BALTICS	LV
Diován 3 mg/ml solución oral	SE/H/0406/007	72.385	NOVARTIS FARMACÉUTICA S.A.	ES
Diovan, 3 mg/ml suukaudne lahus	SE/H/0406/007	687310	SIA NOVARTIS BALTICS	EE
Diovan 3 mg/ml oral lösning	SE/H/0406/007	27966	NOVARTIS FINLAND OY	FI
Diován 320 mg comprimidos recubiertos con película	SE/H/0406/006	68.817	NOVARTIS FARMACÉUTICA S.A.	ES
Diovan 320 mg filmdragerade tabletter	SE/H/0406/006	22668	NOVARTIS FINLAND OY	FI
Diovan 320 mg kalvopäällysteinen tabletti	SE/H/0406/006	22668	NOVARTIS FINLAND OY	FI
Diovan 40 mg filmdragerade tabletter	SE/H/0406/005	20510	NOVARTIS FINLAND OY	FI
Diovan 40 mg kalvopäällysteinen tabletti	SE/H/0406/005	20510	NOVARTIS FINLAND OY	FI
Diovan 80 mg apvalkotās tabletes	SE/H/0406/003	02-0391	SIA NOVARTIS BALTICS	LV
Diován 80 mg comprimidos recubiertos	SE/H/0406/003	64.454	NOVARTIS FARMACÉUTICA S.A.	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
con película				
Diovan 80 mg filmdragerade tabletter	SE/H/0406/003	16708	NOVARTIS FINLAND OY	FI
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/010	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/029	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/030	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/032	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/033	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/034	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/035	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/036	SIA NOVARTIS BALTICS	LT
Diovan 80 mg plėvele dengtos tabletės	SE/H/0406/003	LT/1/01/1236/037	SIA NOVARTIS BALTICS	LT
Diovan 80 mg kalvopäällysteinen tabletti	SE/H/0406/003	16708	NOVARTIS FINLAND OY	FI
Diován Cardio 40 mg comprimidos recubiertos con película	SE/H/0406/005	66.910	NOVARTIS FARMACÉUTICA S.A.	ES
Diovan, oral opløsning	SE/H/0406/007	46831	NOVARTIS HEALTHCARE A/S	DK
Diovan® 160 mg Filmtabletten	SE/H/0406/004	1-24276	NOVARTIS PHARMA GMBH	AT
Диован 160 mg филмирани таблетки	SE/H/0406/004	20020028	NOVARTIS EUROPHARM LIMITED	BG
Diovan 160 mg filmdragerade tabletter	SE/H/0406/004	17495	NOVARTIS SVERIGE AB	SE
Diovan 160 mg filmtabletta	SE/H/0406/004	OGYI-T-8484/13	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
DIOVAN® 160 mg protect Filmtabletten	SE/H/0406/004	50477.01.00	NOVARTIS PHARMA GMBH	DE
Diovan, 160 mg, tabletki powlekane	SE/H/0406/004	9292	NOVARTIS POLAND SP. Z O. O.	PL
Diovan 160, 160 mg filmomhulde tabletten	SE/H/0406/004	RVG 26940	NOVARTIS PHARMA B.V.	NL
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/01	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/02	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/03	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/04	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/05	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/06	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/07	NOVARTIS EUROPHARM LIMITED	RO
Diovan 160 mg comprimate filmate	SE/H/0406/004	3947/2011/08	NOVARTIS EUROPHARM LIMITED	RO
Diovan 3 mg/ml belsőleges oldat	SE/H/0406/007	OGYI-T-8484/15	NOVARTIS HUNGÁRIA KFT.	HU
Diovan® 3 mg/ml Lösung zum Einnehmen	SE/H/0406/007	82267.00.00	NOVARTIS PHARMA GMBH	DE
Diovan 3 mg/ml oral lösning	SE/H/0406/007	43158	NOVARTIS SVERIGE AB	SE
Диован 3 mg/ml перорален разтвор	SE/H/0406/007	20100328	NOVARTIS EUROPHARM LIMITED	BG
Diovan, 3 mg/ml, roztwór doustny	SE/H/0406/007	17043	NOVARTIS POLAND SP. Z O. O.	PL
Diovan 3 mg/ml drank	SE/H/0406/007	RVG 107481	NOVARTIS PHARMA B.V.	NL
Diovan 320 mg filmdragerade tabletter	SE/H/0406/006	22840	NOVARTIS SVERIGE AB	SE
DIOVAN® 320 mg forte	SE/H/0406/006	67298.00.00	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
Filmtabletten				
Diovan 320, 320 mg filmomhulde tabletten	SE/H/0406/006	RVG 34472	NOVARTIS PHARMA B.V.	NL
Diovan® 3 mg/ml Lösung zum Einnehmen	SE/H/0406/007	1-29223	NOVARTIS PHARMA GMBH	AT
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/01	NOVARTIS EUROPHARM LIMITED	RO
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/02	NOVARTIS EUROPHARM LIMITED	RO
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/03	NOVARTIS EUROPHARM LIMITED	RO
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/04	NOVARTIS EUROPHARM LIMITED	RO
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/05	NOVARTIS EUROPHARM LIMITED	RO
Diovan 40 mg comprimate filmate	SE/H/0406/005	3945/2011/06	NOVARTIS EUROPHARM LIMITED	RO
Диован 40 mg филмирани таблетки	SE/H/0406/005	20050206	NOVARTIS EUROPHARM LIMITED	BG
Diovan 40 mg filmdragerade tabletter	SE/H/0406/005	20464	NOVARTIS SVERIGE AB	SE
DIOVAN® 40 mg Filmtabletten	SE/H/0406/005	50477.02.00	NOVARTIS PHARMA GMBH	DE
Diovan® 80 mg Filmtabletten	SE/H/0406/003	1-24275	NOVARTIS PHARMA GMBH	AT
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/01	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/02	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/03	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/04	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/05	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg	SE/H/0406/003	3946/2011/06	NOVARTIS EUROPHARM	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			LIMITED	
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/07	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/08	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg comprimate filmate	SE/H/0406/003	3946/2011/09	NOVARTIS EUROPHARM LIMITED	RO
Diovan 80 mg filmdragerade tabletter	SE/H/0406/003	17494	NOVARTIS SVERIGE AB	SE
Diovan 80 mg filmtabletta	SE/H/0406/003	OGYI-T-8484/09	NOVARTIS HUNGÁRIA KFT.	HU
DIOVAN® 80 mg Filmtabletten	SE/H/0406/003	50477.00.00	NOVARTIS PHARMA GMBH	DE
Diovan, 80 mg, tabletki powlekane	SE/H/0406/003	9291	NOVARTIS POLAND SP. Z O. O.	PL
Diovan 80, 80 mg filmomhulde tabletten	SE/H/0406/003	RVG 26939	NOVARTIS PHARMA B.V.	NL
Diovan 160 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0406/004	19385	NOVARTIS IRELAND LIMITED	CY
Diovan 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0406/005	19635	NOVARTIS IRELAND LIMITED	CY
Diovan 80 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0406/003	19384	NOVARTIS IRELAND LIMITED	CY
Diovan πόσιμο διάλυμα 3 mg/ml	SE/H/0406/007	20694	NOVARTIS IRELAND LIMITED	CY
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	34009 381 551 8 1	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	34009 381 552 4 2	NOVARTIS PHARMA S.A.S.	FR
TAREG 160 mg, comprimé pelliculé	SE/H/0406/004	34009 381 555 3 2	NOVARTIS PHARMA S.A.S.	FR
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	34009 381 538 1 1	NOVARTIS PHARMA S.A.S.	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
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	34009 381 539 8 9	NOVARTIS PHARMA S.A.S.	FR
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	34009 381 540 6 1	NOVARTIS PHARMA S.A.S.	FR
TAREG 40 mg, comprimé pelliculé sécable	SE/H/0406/005	34009 381 543 5 1	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	34009 381 544 1 2	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	34009 381 545 8 0	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	34009 381 546 4 1	NOVARTIS PHARMA S.A.S.	FR
TAREG 80 mg, comprimé pelliculé	SE/H/0406/003	34009 381 549 3 1	NOVARTIS PHARMA S.A.S.	FR
Diovan 3 mg/ml perorální roztok	SE/H/0406/007	58/550/10-C	NOVARTIS S.R.O.,	CZ
Diovan 3 mg/ml perorálny roztok	SE/H/0406/007	58/0287/10-S	NOVARTIS SLOVAKIA S.R.O.	SK
TAREG 3 mg/ml, solution buvable	SE/H/0406/007	34009 491 474 8 9	NOVARTIS PHARMA S.A.S.	FR
Diovan 3 mg/ml mixtúra, lausn	SE/H/0406/007	IS/1/10/078/01	NOVARTIS HEALTHCARE A/S	IS
Diovan 3 mg/ml mikstur, oppløsning	SE/H/0406/007	10-7589	NOVARTIS NORGE AS	NO
Diovan 320 mg filmdrasjerte tabletter	SE/H/0406/006	05-3646	NOVARTIS NORGE AS	NO
Diovan 40 mg filmdrasjerte tabletter	SE/H/0406/005	04-2470	NOVARTIS NORGE AS	NO
Diovan 80 mg filmdrasjerte tabletter	SE/H/0406/003	00-6229	NOVARTIS NORGE AS	NO
Diovan 160 mg filmdrasjerte tabletter	SE/H/0406/004	00-6230	NOVARTIS NORGE AS	NO
Diovan, 160 mg õhukese polümeerikattega tabletid	SE/H/0406/004	375902	SIA NOVARTIS BALTICS	EE
Diovan, 80 mg õhukese	SE/H/0406/003	376002	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
polümeerikattega tabletid				
Diovane 160 mg filmomhulde tabletten	SE/H/0406/004	BE296423	NOVARTIS PHARMA N.V.	BE
Diovane 160 mg comprimés pelliculés	SE/H/0406/004	BE296423	NOVARTIS PHARMA N.V.	BE
Diovane 80 mg filmomhulde tabletten	SE/H/0406/003	BE296414	NOVARTIS PHARMA N.V.	BE
Tareg 160 mg compresse rivestite con film	SE/H/0406/004	033178930	NOVARTIS FARMA S.P.A.	IT
Tareg 160 mg compresse rivestite con film	SE/H/0406/004	033178942	NOVARTIS FARMA S.P.A.	IT
Tareg 160 mg compresse rivestite con film	SE/H/0406/004	033178955	NOVARTIS FARMA S.P.A.	IT
Tareg 160 mg compresse rivestite con film	SE/H/0406/004	033178967	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178548	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178551	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178563	NOVARTIS FARMA S.P.A.	IT
Tareg 40 mg compresse rivestite con film	SE/H/0406/005	033178599	NOVARTIS FARMA S.P.A.	IT
Tareg 80 mg compresse rivestite con film	SE/H/0406/003	033178880	NOVARTIS FARMA S.P.A.	IT
Tareg 80 mg compresse rivestite con film	SE/H/0406/003	033178892	NOVARTIS FARMA S.P.A.	IT
Tareg 80 mg compresse rivestite con film	SE/H/0406/003	033178904	NOVARTIS FARMA S.P.A.	IT
Tareg 80 mg compresse rivestite con film	SE/H/0406/003	033178916	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
Diovan 160 mg film-coated tablets	SE/H/0406/004	PA 0896/009/002	NOVARTIS IRELAND LIMITED	IE
Diovan 3 mg/ml oral solution	SE/H/0406/007	PA 0896/009/005	NOVARTIS IRELAND LIMITED	IE
Diovan 320 mg film-coated tablets	SE/H/0406/006	PA 0896/009/004	NOVARTIS IRELAND LIMITED	IE
Diovan 40 mg film-coated tablets	SE/H/0406/005	PA 0896/009/003	NOVARTIS IRELAND LIMITED	IE
Diovan 80 mg film-coated tablets	SE/H/0406/003	PA 0896/009/001	NOVARTIS IRELAND LIMITED	IE
Diovan 80 mg film-coated tablets	SE/H/0406/003	MA1249/02101	NOVARTIS IRELAND LIMITED	MT
Diovan 160 mg film-coated tablets	SE/H/0406/004	MA1249/02102	NOVARTIS IRELAND LIMITED	MT
Diovan 320 mg film-coated tablets	SE/H/0406/006	MA1249/02104	NOVARTIS IRELAND LIMITED	MT
Diovan 40 mg film-coated tablets	SE/H/0406/005	MA1249/02103	NOVARTIS IRELAND LIMITED	MT
Diovan 3 mg/ml oral solution	SE/H/0406/007	MA1249/02105	NOVARTIS IRELAND LIMITED	MT
Diovane 320 mg filmomhulde tabletten	SE/H/0406/006	BE297604	NOVARTIS PHARMA N.V.	BE
Diovane 320 mg comprimés pelliculés	SE/H/0406/006	BE297604	NOVARTIS PHARMA N.V.	BE
Diovane 3 mg/ml drank	SE/H/0406/007	BE371007	NOVARTIS PHARMA N.V.	BE
Diovane 3 mg/ml solution buvable	SE/H/0406/007	BE371007	NOVARTIS PHARMA N.V.	BE
Diovan 3 mg/ml geriamasis tirpalas	SE/H/0406/007	LT/1/01/1236/014	SIA NOVARTIS BALTICS	LT
Valpression 40 mg compresse rivestite con film	not available	033119239	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119203	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg	not available	033119215	A. MENARINI - INDUSTRIE	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 rivestite con film			FARMACEUTICHE RIUNITE - S.R.L.	
Valpression 40 mg compresse rivestite con film	not available	033119254	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119292	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119266	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119227	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119241	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119280	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119278	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119330	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119316	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119304	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119153	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg	not available	033119177	A. MENARINI - INDUSTRIE	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 rivestite con film			FARMACEUTICHE RIUNITE - S.R.L.	
Valpression 320 mg compresse rivestite con film	not available	033119328	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119342	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119140	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119138	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119191	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119165	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 40 mg compresse rivestite con film	not available	033119189	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 80 mg capsule	not available	033119013	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 160 mg capsule	not available	033119025	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Valpression 320 mg compresse rivestite con film	not available	033119355	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Vals 80 mg comprimidos recubiertos con película.	not available	64.495	ESTEVE PHARMACEUTICALS, S.A.	ES
Vals 160 mg comprimidos recubiertos	not available	64.494	ESTEVE PHARMACEUTICALS, S.A.	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
con película.				
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	SE/H/0565/002	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA -	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
mg/25 mg comprimidos revestidos por película			PRODUTOS FARMACÊUTICOS S.A.	
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	SE/H/0565/003	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA -	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
mg comprimidos revestidos por película			PRODUTOS FARMACÊUTICOS S.A.	
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	SE/H/0565/001	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	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
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	SE/H/0565/002	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	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
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	SE/H/0565/003	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	SE/H/0565/001	NOVARTIS HUNGÁRIA KFT.	HU
Co-Diovane 160 mg/25 mg, comprimés pelliculés	SE/H/0565/003	2005010014	NOVARTIS PHARMA N.V.	LU
Co-Diovane 80 mg/12,5 mg, comprimés pelliculés	SE/H/0565/001	2002107201	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Co-Diovane 160 mg/12,5 mg, comprimés pelliculés	SE/H/0565/002	2003100011	NOVARTIS PHARMA N.V.	LU
Co-Diovane 160 mg /25 mg Filmtabletten	SE/H/0565/003	2005010014	NOVARTIS PHARMA N.V.	LU
Co-Diovane 160 mg /12,5 mg Filmtabletten	SE/H/0565/002	2003100011	NOVARTIS PHARMA N.V.	LU
Co-Diovane 80 mg /12,5 mg Filmtabletten	SE/H/0565/001	2002107201	NOVARTIS PHARMA N.V.	LU
Co-Diovane 80 mg /12,5 mg Filmtabletten	SE/H/0565/001	BE192893	NOVARTIS PHARMA N.V.	BE
Co-Diovane 160 mg /12,5 mg Filmtabletten	SE/H/0565/002	BE258517	NOVARTIS PHARMA N.V.	BE
Co-Diovane 160 mg /25 mg Filmtabletten	SE/H/0565/003	BE271747	NOVARTIS PHARMA N.V.	BE
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-02	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-03	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-04	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-05	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-06	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-07	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-08	NOVARTIS HRVATSKA D.O.O.	HR

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
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-02	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-03	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-04	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-05	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-06	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-07	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-08	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-09	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-02	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-03	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-04	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-05	NOVARTIS HRVATSKA D.O.O.	HR

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
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-06	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-07	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-08	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-02	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-03	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-04	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-05	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-06	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-07	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-08	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-09	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-02	NOVARTIS HRVATSKA D.O.O.	HR

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
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-03	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-04	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-05	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-06	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-07	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-08	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-09	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/019	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/001	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/035	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/051	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/060	NOVARTIS EUROPHARM LIMITED	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
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114037	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114049	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114417	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114429	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114431	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114443	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114052	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114381	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114393	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114405	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114506	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114518	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
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114520	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114532	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114544	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114456	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114468	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114470	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114482	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114494	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114557	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114569	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114571	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114583	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
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114595	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114607	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114619	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114621	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114633	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114645	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114658	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114660	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114672	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114684	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114696	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114708	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
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114064	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114076	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114088	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114090	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114102	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114114	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/12,5 mg compresse rivestite con film	SE/H/0565/002	034114126	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114138	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114140	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114153	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114165	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114177	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
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114189	NOVARTIS FARMA S.P.A.	IT
Cotareg 160 mg/25 mg compresse rivestite con film	SE/H/0565/003	034114191	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114203	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114215	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114227	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114239	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114241	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114254	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114266	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114278	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/12,5 mg compresse rivestite con film	SE/H/0565/004	034114367	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114280	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
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114292	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114304	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114316	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114328	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114330	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114342	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114355	NOVARTIS FARMA S.P.A.	IT
Cotareg 320 mg/25 mg compresse rivestite con film	SE/H/0565/005	034114379	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114013	NOVARTIS FARMA S.P.A.	IT
Cotareg 80 mg/12,5 mg compresse rivestite con film	SE/H/0565/001	034114025	NOVARTIS FARMA S.P.A.	IT
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/002	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/003	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/004	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/005	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/006	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/007	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/008	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/009	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/010	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/011	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/012	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/020	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/021	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/022	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/023	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/013	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/014	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/015	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/016	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/017	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 80 mg/12,5 mg filmsko obložene tablete	SE/H/0565/001	H/03/00406/018	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/024	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/025	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/026	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/027	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/028	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/029	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/030	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/031	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/032	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/033	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/12,5 mg filmsko obložene tablete	SE/H/0565/002	H/03/00406/034	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/036	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/037	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/038	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/039	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/040	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/041	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/042	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/043	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/044	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/045	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/046	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/047	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/048	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/049	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 160 mg/25 mg filmsko obložene tablete	SE/H/0565/003	H/03/00406/050	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/052	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/053	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/054	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/055	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/056	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/057	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/058	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/12,5 mg filmsko obložene tablete	SE/H/0565/004	H/03/00406/059	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/061	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/062	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/063	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/064	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/065	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/066	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/067	NOVARTIS EUROPHARM LIMITED	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
Co-Diovan 320 mg/25 mg filmsko obložene tablete	SE/H/0565/005	H/03/00406/068	NOVARTIS EUROPHARM LIMITED	SI
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890409	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890509	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890107	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890108	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890201	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890202	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890203	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890204	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890205	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890206	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	238890207	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
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890301	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890302	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890303	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890304	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890305	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890306	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890307	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890308	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890309	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (160+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	238890310	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890401	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890402	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
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890403	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890404	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890405	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890406	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890407	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/004	238890408	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890501	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890502	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890503	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890504	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890505	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890506	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
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890507	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (320+25) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/005	238890508	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890101	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890102	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890103	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890104	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890105	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan (80+12,5) mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	238890106	NOVARTIS (HELLAS) S.A.C.I.	GR
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4804688	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4804787	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4804886	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
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4804985	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4805081	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4805180	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	4805289	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	5022769	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/12,5 mg 160 mg/ 12,5 mg comprimidos revestidos por película	SE/H/0565/002	5022777	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	2644383	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	2644482	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	2644581	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	5022744	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
Co-Diovan 80 mg /12,5 mg comprimidos revestidos por película	SE/H/0565/001	5022751	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5022801	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5022819	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5273388	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5273487	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5273586	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan Forte 160 mg/25 mg comprimidos revestidos por película	SE/H/0565/003	5273685	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Co-Diovan 160 mg/25 mg plėvele dengtos tabletės	SE/H/0565/003	LT/1/01/1235/003	SIA NOVARTIS BALTICS	LT
Co-Diovan 160 mg/12,5 mg plėvele dengtos tabletės	SE/H/0565/002	LT/1/01/1235/002	SIA NOVARTIS BALTICS	LT
Co-Diovan 160 mg/12,5 mg apvalkotās tabletes	SE/H/0565/002	02-0393	SIA NOVARTIS BALTICS	LV
Co-Diovan 160 mg/25 mg apvalkotās tabletes	SE/H/0565/003	02-0394	SIA NOVARTIS BALTICS	LV
Co-Diovan 80 mg/12,5 mg plėvele dengtos tabletės	SE/H/0565/001	LT/1/01/1235/001	SIA NOVARTIS BALTICS	LT
Co-Diovan 80 mg/12,5 mg apvalkotās tabletes	SE/H/0565/001	98–0310	SIA NOVARTIS BALTICS	LV

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
Co-Diovan, 80/12,5 mg õhukese polümeerikattega tabletid	SE/H/0565/001	246499	SIA NOVARTIS BALTICS	EE
Diovan Comp 160 mg/12,5 mg kalvopäällysteiset tabletit	SE/H/0565/002	18231	NOVARTIS FINLAND OY	FI
Diovan Comp 160 mg/25 mg kalvopäällysteiset tabletit	SE/H/0565/003	19618	NOVARTIS FINLAND OY	FI
Diovan Comp 160 mg/12,5 mg filmdragerade tabletter	SE/H/0565/002	18231	NOVARTIS FINLAND OY	FI
Diovan Comp 160 mg/25 mg filmdragerade tabletter	SE/H/0565/003	19618	NOVARTIS FINLAND OY	FI
Diovan Comp 80 mg/12,5 mg kalvopäällysteiset tabletit	SE/H/0565/001	13181	NOVARTIS FINLAND OY	FI
Diovan Comp 80 mg/12,5 mg filmdragerade tabletter	SE/H/0565/001	13181	NOVARTIS FINLAND OY	FI
Co-Diován 160 mg/12,5 mg comprimidos recubiertos con película	SE/H/0565/002	65.794	NOVARTIS FARMACÉUTICA S.A.	ES
Co-Diován 320 mg/12,5 mg comprimidos recubiertos con película	SE/H/0565/004	69.930	NOVARTIS FARMACÉUTICA S.A.	ES
Co-Diován 80 mg/12,5 mg comprimidos recubiertos con película	SE/H/0565/001	62.206	NOVARTIS FARMACÉUTICA S.A.	ES
Co-Diován Forte 160 mg/25 mg comprimidos recubiertos con película	SE/H/0565/003	66.526	NOVARTIS FARMACÉUTICA S.A.	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
Co-Diován Forte 320 mg/25 mg comprimidos recubiertos con película	SE/H/0565/005	69.942	NOVARTIS FARMACÉUTICA S.A.	ES
CoDiovan® 160 mg/12,5 mg Filmtabletten	SE/H/0565/002	52884.00.00	NOVARTIS PHARMA GMBH	DE
Co-Diovan 160 mg + 25 mg tabletki powlekane	SE/H/0565/003	10629	NOVARTIS POLAND SP. Z O. O.	PL
Ко-Диован 160 mg/12,5 mg филмирани таблетки	SE/H/0565/002	20020026	NOVARTIS EUROPHARM LIMITED	BG
Ко-Диован 160 mg/25 mg филмирани таблетки	SE/H/0565/003	20040085	NOVARTIS EUROPHARM LIMITED	BG
Co-Diovan 160/12,5, 160 mg/12,5 mg filmomhulde tabletten	SE/H/0565/002	RVG 29491	NOVARTIS PHARMA B.V.	NL
Co-Diovan 160 mg/12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/002	18977	NOVARTIS IRELAND LIMITED	CY
Co-Diovan 160/25, 160 mg/25 mg filmomhulde tabletten	SE/H/0565/003	RVG 31122	NOVARTIS PHARMA B.V.	NL
Co-Diovan 160 mg/25 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/003	19477	NOVARTIS IRELAND LIMITED	CY
CoDiovan® 320 mg/12,5 mg Filmtabletten	SE/H/0565/004	69620.00.00	NOVARTIS PHARMA GMBH	DE
Co-Diovan 320/12,5, 320 mg/12,5 mg filmomhulde tabletten	SE/H/0565/004	RVG 100453	NOVARTIS PHARMA B.V.	NL
Co-Diovan 320/25, 320 mg/25 mg filmomhulde tabletten	SE/H/0565/005	RVG 100454	NOVARTIS PHARMA B.V.	NL
Co-Diovan® 80 mg/12,5 mg – Filmtabletten	SE/H/0565/001	1 - 22463	NOVARTIS PHARMA GMBH	AT
CoDiovan® 80 mg/12,5	SE/H/0565/001	40763.00.00	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
mg Filmtabletten				
Co-Diovan 80/12,5, 80 mg/12,5 mg filmomhulde tabletten	SE/H/0565/001	RVG 22365	NOVARTIS PHARMA B.V.	NL
Co-Diovan 80 mg/12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0565/001	17851	NOVARTIS IRELAND LIMITED	CY
CoDiovan® forte 160 mg/25 mg Filmtabletten	SE/H/0565/003	52884.01.00	NOVARTIS PHARMA GMBH	DE
Co-Diovan forte® 160 mg/12,5 mg – Filmtabletten	SE/H/0565/002	1 – 25101	NOVARTIS PHARMA GMBH	AT
CoDiovan® forte 320 mg/25 mg Filmtabletten	SE/H/0565/005	69621.00.00	NOVARTIS PHARMA GMBH	DE
Co-Diovan fortissimum®160 mg/25 mg - Filmtabletten	SE/H/0565/003	1-25735	NOVARTIS PHARMA GMBH	AT
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 363 107 2 8	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 371 254 0 6	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 372 013 7 7	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 372 014 3 8	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 565 9 1	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 566 5 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 567 1 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 568 8 1	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 569 4 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5	SE/H/0565/002	34009 381 570 2 4	NOVARTIS PHARMA S.A.S.	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
mg, comprimé pelliculé				
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 381 571 9 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 564 976 8 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 564 977 4 4	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/12,5 mg, comprimé pelliculé	SE/H/0565/002	34009 564 978 0 5	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 359 232 0 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 371 394 7 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 371 395 3 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 371 397 6 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 572 5 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 573 1 4	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 574 8 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 575 4 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 576 0 4	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 577 7 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 381 578 3 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 563 683 7 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25 mg, comprimé pelliculé	SE/H/0565/003	34009 563 684 3 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 160 mg/25	SE/H/0565/003	34009 563 686 6 2	NOVARTIS PHARMA S.A.S.	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
mg, comprimé pelliculé				
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 344 300 5 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 344 301 1 4	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 344 302 8 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 344 303 4 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 371 961 9 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 371 962 5 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 371 963 1 4	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 557 6 1	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 558 2 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 559 9 0	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 560 7 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 561 3 3	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 563 6 2	NOVARTIS PHARMA S.A.S.	FR
COTAREG 80 mg/12,5 mg, comprimé pelliculé	SE/H/0565/001	34009 381 564 2 3	NOVARTIS PHARMA S.A.S.	FR
Diovan Comp 160 mg/12,5 mg filmdragerade tabletter	SE/H/0565/002	19525	NOVARTIS SVERIGE AB	SE
Diovan Comp 160 mg/25 mg filmdragerade tabletter	SE/H/0565/003	21204	NOVARTIS SVERIGE AB	SE
Diovan Comp 320	SE/H/0565/004	23641	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/12,5 mg filmdragerade tabletter				
Diovan Comp 320 mg/25 mg filmdragerade tabletter	SE/H/0565/005	23642	NOVARTIS SVERIGE AB	SE
Diovan Comp 80 mg/12,5 mg filmdragerade tabletter	SE/H/0565/001	14098	NOVARTIS SVERIGE AB	SE
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	OGYI-T-6552/05	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	OGYI-T-6552/06	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	OGYI-T-6552/07	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/12,5 mg filmtabletta	SE/H/0565/002	OGYI-T-6552/08	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	OGYI-T-6552/09	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	OGYI-T-6552/10	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	OGYI-T-6552/11	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 160/25 mg filmtabletta	SE/H/0565/003	OGYI-T-6552/12	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	OGYI-T-6552/01	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	OGYI-T-6552/02	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	OGYI-T-6552/03	NOVARTIS HUNGÁRIA KFT.	HU
Diovan HCT 80/12,5 mg filmtabletta	SE/H/0565/001	OGYI-T-6552/04	NOVARTIS HUNGÁRIA KFT.	HU
Diovan Comp 160 mg/25 mg filmdrasjerte tabletter	SE/H/0565/003	01-10080	NOVARTIS NORGE AS	NO
Diovan Comp 320	SE/H/0565/004	06-4074	NOVARTIS NORGE AS	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
mg/12,5 mg filmdrasjerte tabletter				
Diovan Comp 320 mg/25 mg filmdrasjerte tabletter	SE/H/0565/005	06-4075	NOVARTIS NORGE AS	NO
Diovan Comp 160 mg/12,5 mg filmdrasjerte tabletter	SE/H/0565/002	01-10079	NOVARTIS NORGE AS	NO
Diovan Comp 80 mg/12,5 mg filmdrasjerte tabletter	SE/H/0565/001	97-2716	NOVARTIS NORGE AS	NO
Co-Diovane 160 mg/12,5 mg, comprimés pelliculés	SE/H/0565/002	BE258517	NOVARTIS PHARMA N.V.	BE
Co-Diovane 160 mg/12,5 mg filmomhulde tabletten	SE/H/0565/002	BE258517	NOVARTIS PHARMA N.V.	BE
Co-Diovane 80 mg/12,5 mg filmomhulde tabletten	SE/H/0565/001	BE192893	NOVARTIS PHARMA N.V.	BE
Co-Diovane 80 mg/12,5 mg, comprimés pelliculés	SE/H/0565/001	BE192893	NOVARTIS PHARMA N.V.	BE
Co-Diovan 160 mg/12.5 mg film-coated tablets	SE/H/0565/002	PA 0896/007/002	NOVARTIS IRELAND LIMITED	IE
Co-Diovan 160 mg/25 mg film-coated tablets	SE/H/0565/003	PA 0896/007/003	NOVARTIS IRELAND LIMITED	IE
Co-Diovan 320 mg/12.5 mg film-coated tablets	SE/H/0565/004	PA 0896/007/004	NOVARTIS IRELAND LIMITED	IE
Co-Diovan 320 mg/25 mg film-coated tablets	SE/H/0565/005	PA 0896/007/005	NOVARTIS IRELAND LIMITED	IE
Co-Diovan 80 mg/12.5 mg film-coated tablets	SE/H/0565/001	PA 0896/007/001	NOVARTIS IRELAND LIMITED	IE
Co-Diovan 160 mg/25 mg film-coated tablets	SE/H/0565/003	MA1249/02003	NOVARTIS IRELAND LIMITED	MT
Co-Diovan 80 mg/12.5	SE/H/0565/001	MA1249/02001	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
mg film-coated tablets			LIMITED	
Co-Diovan 160 mg/12.5 mg film-coated tablets	SE/H/0565/002	MA1249/02002	NOVARTIS IRELAND LIMITED	MT
Co-Diovan 320 mg/12.5 mg film-coated tablets	SE/H/0565/004	MA1249/02004	NOVARTIS IRELAND LIMITED	MT
Co-Diovan 320 mg/25 mg film-coated tablets	SE/H/0565/005	MA1249/02005	NOVARTIS IRELAND LIMITED	MT
Co-Diovane 160 mg/25 mg filmomhulde tabletten	SE/H/0565/003	BE271747	NOVARTIS PHARMA N.V.	BE
Co-Diovane 160 mg/25 mg, comprimés pelliculés	SE/H/0565/003	BE271747	NOVARTIS PHARMA N.V.	BE
Co-Diovan 160 mg/12,5 mg filmom obložene tablete	SE/H/0565/002	HR-H-819796318-01	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 160 mg/25 mg filmom obložene tablete	SE/H/0565/003	HR-H-133986302-01	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/12,5 mg filmom obložene tablete	SE/H/0565/004	HR-H-174157013-01	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 320 mg/25 mg filmom obložene tablete	SE/H/0565/005	HR-H-063837675-01	NOVARTIS HRVATSKA D.O.O.	HR
Co-Diovan 80 mg/12,5 mg filmom obložene tablete	SE/H/0565/001	HR-H-035681755-01	NOVARTIS HRVATSKA D.O.O.	HR
Kalpress 160 mg comprimidos recubiertos con película	SE/H/0407/002	64.452	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress 320 mg comprimidos recubiertos con película	SE/H/0407/004	69.517	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress 80 mg comprimidos recubiertos	SE/H/0407/001	64.453	NOVARTIS FARMACÉUTICA S.A.	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
con película				
Angiosan, 160 mg filmdragerad tablett	SE/H/0407/002	19956	NOVARTIS SVERIGE AB	SE
Angiosan, 320 mg filmdragerad tablett	SE/H/0407/004	22841	NOVARTIS SVERIGE AB	SE
Angiosan, 40 mg filmdragerad tablett	SE/H/0407/003	20465	NOVARTIS SVERIGE AB	SE
Angiosan, 80 mg filmdragerad tablett	SE/H/0407/001	19955	NOVARTIS SVERIGE AB	SE
Valsartan 40 mg film-coated tablets	not available	PL 17907/0360	BRISTOL LABORATORIES LIMITED	XI
Valsartan 80 mg film-coated tablets	not available	PL 17907/0361	BRISTOL LABORATORIES LIMITED	XI
Valsartan 160 mg film-coated tablets	not available	PL 17907/0362	BRISTOL LABORATORIES LIMITED	XI
Valsartan 320 mg film-coated tablets	not available	PL 17907/0363	BRISTOL LABORATORIES LIMITED	XI
Co-Vals Forte 160 mg/25 mg comprimidos recubiertos con película	not available	66.676	ESTEVE PHARMACEUTICALS, S.A.	ES
Co-Vals Forte 320 mg/25 mg comprimidos recubiertos con película	not available	70.363	ESTEVE PHARMACEUTICALS, S.A.	ES
Valsartan 40mg capsules	not available	PL 20416/0503	CRESCENT PHARMA LIMITED	XI
Valsartan 80mg capsules	not available	PL 20416/0504	CRESCENT PHARMA LIMITED	XI
Valsartan 160mg capsules	not available	PL 20416/0505	CRESCENT PHARMA LIMITED	XI
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/03	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/12	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/02	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160	not available	OGYI-T-21875/10	ACTAVIS GROUP PTC EHF.	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
mg/12.5 mg filmtabletta				
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/06	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/05	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/04	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/11	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/16	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/17	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/20	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/21	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/18	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/19	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/13	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/07	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/09	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/14	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/08	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 160 mg/25 mg filmtabletta	not available	OGYI-T-21875/15	ACTAVIS GROUP PTC EHF.	HU
Valsocard HCT 80 mg/12.5 mg filmtabletta	not available	OGYI-T-21875/01	ACTAVIS GROUP PTC EHF.	HU
Co-Vals Forte 160 mg/25	not available	66.676	ESTEVE	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
mg comprimidos recubiertos con película			PHARMACEUTICALS, S.A.	
Co-Vals Forte 320 mg/25 mg comprimidos recubiertos con película	not available	70.363	ESTEVE PHARMACEUTICALS, S.A.	ES
Vals 80 mg comprimidos recubiertos con película.	not available	64.495	ESTEVE PHARMACEUTICALS, S.A.	ES
Vals 160 mg comprimidos recubiertos con película.	not available	64.494	ESTEVE PHARMACEUTICALS, S.A.	ES
Vals 320 mg comprimidos recubiertos con película.	not available	69.345	ESTEVE PHARMACEUTICALS, S.A.	ES
Co-Vals 320 mg/12,5 mg comprimidos recubiertos con película	not available	70.361	ESTEVE PHARMACEUTICALS, S.A.	ES
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/12	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/11	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/10	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/09	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/14	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/13	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/16	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/17	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/19	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/18	ACTAVIS GROUP PTC EHF.	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
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/20	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/21	ACTAVIS GROUP PTC EHF.	HU
Valsocard 80 mg fimtabletta	not available	OGYI-T-21874/08	ACTAVIS GROUP PTC EHF.	HU
Valsocard 160 mg fimtabletta	not available	OGYI-T-21874/15	ACTAVIS GROUP PTC EHF.	HU
Kalpress Plus 160 mg/12,5 mg comprimidos recubiertos con película	ES/H/0895/002	65.898	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress Plus 320 mg/12,5 mg comprimidos recubiertos con película	ES/H/0895/004	70.058	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress Plus 80mg/12,5 mg comprimidos recubiertos con película	ES/H/0895/001	62.562	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress Plus Forte 160 mg/25 mg comprimidos recubiertos con película	ES/H/0895/003	66.744	NOVARTIS FARMACÉUTICA S.A.	ES
Kalpress Plus Forte 320 mg/25 mg comprimidos recubiertos con película	ES/H/0895/005	70.060	NOVARTIS FARMACÉUTICA S.A.	ES
Troval 80 mg Film-coated tablets	not available	ML 21333	DELORBIS PHARMACEUTICALS LIMITED	CY
Troval 160 mg Film-coated tablets	not available	ML 21334	DELORBIS PHARMACEUTICALS LIMITED	CY
Co-Vals Forte 160 mg/25 mg comprimidos recubiertos con película	not available	66.676	ESTEVE PHARMACEUTICALS, S.A.	ES
Co-Vals Forte 320 mg/25 mg comprimidos	not available	70.363	ESTEVE PHARMACEUTICALS, S.A.	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
recubiertos con película				
Valsartan ratiopharm 120 mg filmdragerade tabletter	SE/H/0747/003	26041	RATIOPHARM GMBH	SE
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134078	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134155	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134054	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134039	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134092	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134080	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134041	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134167	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134116	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134104	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse	not available	034134128	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE -	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
rivestite con film			S.R.L.	
COMBISARTAN 80 mg/12,5 mg compresse rivestite con film	not available	034134015	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134142	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 80 mg/12,5 mg compresse rivestite con film	not available	034134027	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/12,5 mg compresse rivestite con film	not available	034134066	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 160 mg/25 mg compresse rivestite con film	not available	034134130	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134332	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134229	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134205	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134217	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134179	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134243	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse	not available	034134181	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE -	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
rivestite con film			S.R.L.	
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134231	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/12,5 mg compresse rivestite con film	not available	034134193	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134268	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134256	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134294	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134320	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134344	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134306	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134270	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134318	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
COMBISARTAN 320 mg/25 mg compresse rivestite con film	not available	034134282	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT