17 May 2018
EMA/309956/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): losartan

Procedure No.: PSUSA/00001912/201709

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LOSARTAN EG 100 mg, comprimé pelliculé	not available	NL34600	EG LABO LABORATOIRES EUROGENERICS	FR
Losartankalium "Actavis", filmovertrukne tabletter	DK/H/0922/002	38576	ACTAVIS GROUP PTC EHF.	DK
Losartan Actavis 25 mg filmdrasjerte tabletter	DK/H/0922/002	06-4246	ACTAVIS GROUP HF.	NO
Losartan Actavis 25 mg filmdragerade tabletter	DK/H/0922/002	24054	ACTAVIS GROUP HF.	SE
Losartan Actavis, 25 mg õhukese polümeerikattega tabletid	DK/H/0922/002	545407	ACTAVIS GROUP HF.	EE
COZAAR® 50 mg filmom obložene tablete	not available	HR-H-243903537	MERCK SHARP & DOHME D.O.O.	HR
COZAAR® 100 mg filmom obložene tablete	not available	HR-H-174024498	MERCK SHARP & DOHME D.O.O.	HR
COZAAR® 50 mg, comprimés pelliculés	NL/H/1457/002	1995040196	MSD BELGIUM BVBA/SPRL	LU
COZAAR® 100 mg, comprimés pelliculés	NL/H/1457/003	2007019153	MSD BELGIUM BVBA/SPRL	LU
COZAAR® 12,5 mg, comprimés pelliculés	NL/H/1457/001	1999060006	MSD BELGIUM BVBA/SPRL	LU
COZAAR® 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	2009030031	MSD BELGIUM BVBA/SPRL	LU
Kaliumlosartan Sandoz 25 mg, filmomhulde tabletten	not available	RVG 31868	SANDOZ B.V.	NL
Losarix 25 mg Filmtabletten	DE/H/1059/001	69470.00.00	HEXAL AG	DE
Losartan HEXAL 75 mg Filmtabletten	not available	60875.00.00	HEXAL AG	DE
Losartan 75 - 1 A Pharma	not available	60876.00.00	1 A PHARMA GMBH	DE
Losartan HEXAL 25 mg Filmtabletten	not available	60867.00.00	HEXAL AG	DE
Losartan 25 - 1 A Pharma	not available	60868.00.00	1 A PHARMA GMBH	DE
Lotanos 100mg Film-coated Tablets	not available	PA0711/123/005	ROWEX LTD	IE
Lotanos 50mg Film-coated Tablets	not available	PA0711/123/003	ROWEX LTD	IE
Losartan Aurobindo 25 mg filmdragerade tabletter	NL/H/3056/001	50998	AUROBINDO PHARMA (MALTA) LIMITED	SE
Losartan Aurobindo 25 mg compresse rivestite con film	NL/H/3056/001	043614015	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan Aurobindo 25 mg compresse rivestite con film	NL/H/3056/001	043614027	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan Aurobindo 25 mg compresse rivestite con film	NL/H/3056/001	043614039	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan Aurobindo 25 mg compresse rivestite con film	NL/H/3056/001	043614041	AUROBINDO PHARMA (ITALIA) S.R.L.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Losartan Aurobindo 25 mg compresse rivestite con film	NL/H/3056/001	043614054	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Cozaar® 25 mg film-coated tablets	not available	PL 0025/0336	MERCK SHARP & DOHME LTD.	UK
Neo-lotan 12,5 mg compresse rivestite con film	not available	029385034	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 12,5 mg compresse rivestite con film	not available	029385022	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 50 mg compresse rivestite con film	not available	029385010	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 100 mg compresse rivestite con film	not available	029385046	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 2.5 mg/ml polvere e solvente per sospensione orale	not available	029385059	NEOPHARMED GENTILI S.R.L.	IT
Lortaan IC 12,5 mg comprimidos revestidos por película	not available	3351582	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan IC 12,5 mg comprimidos revestidos por película	not available	3351681	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 50 mg comprimidos revestidos por película	not available	2356384	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 50 mg comprimidos revestidos por película	not available	2356483	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 50 mg comprimidos revestidos por película	not available	2356582	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 100 mg 100 mg comprimidos revestidos por película	not available	4103982	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 100 mg 100 mg comprimidos revestidos por película	not available	4104089	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 100 mg 100 mg comprimidos revestidos por película	not available	4104188	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Lortaan 100 mg 100 mg comprimidos revestidos por película	not available	4104287	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
COZAAR 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1457/002	16155	MERCK SHARP & DOHME BV	CY
COZAAR 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1457/003	19537	MERCK SHARP & DOHME BV	CY
COZAAR 12.5 mg επικαλυμμένα με λεπτό	NL/H/1457/001	20464	MERCK SHARP & DOHME BV	CY

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υμένιο δισκία				
Losartan Heumann 25 mg Filmtabletten	-	72064.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Losartan-ratiopharm® 25 mg Filmtabletten	DE/H/3505/002	86525.00.00	RATIOPHARM GMBH	DE
Losartan-ratiopharm® 12,5 mg Filmtabletten	DE/H/3505/001	2014090254	RATIOPHARM GMBH	LU
Losartan-ratiopharm® 50 mg Filmtabletten	DE/H/3505/003	2008040024	RATIOPHARM GMBH	LU
Losartan-ratiopharm® 25 mg Filmtabletten	DE/H/3505/002	2014090255	RATIOPHARM GMBH	LU
Losartan-ratiopharm® 100 mg Filmtabletten	DE/H/3505/005	2015120220	RATIOPHARM GMBH	LU
Losartan potassium 25 mg film-coated tablets	not available	PL 21880/0140	MEDREICH PLC	UK
Losartan potassium 50 mg film-coated tablets	not available	PL 21880/0141	MEDREICH PLC	UK
Losartan potassium 100 mg film-coated tablets	not available	PL 21880/0142	MEDREICH PLC	UK
Losartan potassium 12.5 mg film-coated tablets	not available	PL21880/0139	MEDREICH PLC	UK
LOSARTAN EUROGENERICI 25 mg compresse rivestite con film	not available	039410016	EG S.P.A.	IT
Losartan-Kalium PUREN 25 mg Filmtabletten	PT/H/1543/001	96040.00.00	PUREN PHARMA GMBH & CO. KG	DE
Losartán NORMON 25 mg Comprimidos recubiertos con película	not available	67911	LABORATORIOS NORMON, S.A.	ES
Losanox 25 mg filmomhulde tabletten	DK/H/0915/001	RVG 33624	LABORATORIOS LICONSA, S.A.	NL
Cosaar 50 mg Filmtabletten	NL/H/1457/002	1-21306	MERCK SHARP & DOHME GES.M.B.H.	AT
Cosaar 100 mg Filmtabletten	NL/H/1457/003	1-24232	MERCK SHARP & DOHME GES.M.B.H.	AT
Cosaar 12,5 mg Filmtabletten	NL/H/1457/001	1-22927	MERCK SHARP & DOHME GES.M.B.H.	AT
Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317683	MERCK SHARP & DOHME, LDA.	PT

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Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317584	MERCK SHARP & DOHME, LDA.	PT
COZAAR 50 mg film-coated tablets	NL/H/1457/002	PA1286/4/2	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317782	MERCK SHARP & DOHME, LDA.	PT
Cozaar 50 mg filmomhulde tabletten	NL/H/1457/002	RVG 17617	MERCK SHARP & DOHME BV	NL
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982881	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982485	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982584	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982782	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982386	MERCK SHARP & DOHME, LDA.	PT
COZAAR 100 mg film-coated tablets	NL/H/1457/003	PA1286/4/3	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar 100 mg filmomhulde tabletten	NL/H/1457/003	RVG 26791	MERCK SHARP & DOHME BV	NL
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982683	MERCK SHARP & DOHME, LDA.	PT
Cozaar IC, 12,5 mg, comprimidos revestidos por película	NL/H/1457/001	3335080	MERCK SHARP & DOHME, LDA.	PT
Cozaar IC, 12,5 mg, comprimidos revestidos por película	NL/H/1457/001	3335189	MERCK SHARP & DOHME, LDA.	PT
COZAAR 12.5 mg film-coated tablets	NL/H/1457/001	PA1286/4/1	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar 12,5 mg filmomhulde tabletten	NL/H/1457/001	RVG 101836	MERCK SHARP & DOHME BV	NL
Cozaar 2,5 mg/ml pó e solvente para suspensão oral	NL/H/1457/004	5178207	MERCK SHARP & DOHME, LDA.	PT
Cozaar 2,5 mg/ml poeder en oplosmiddel voor orale suspensie	NL/H/1457/004	RVG 103001	MERCK SHARP & DOHME BV	NL
COZAAR 2.5 mg/ml powder and solvent for oral suspension	NL/H/1457/004	PA1286/4/4	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
COZAAR 50 mg, comprimé pelliculé	NL/H/1457/002	34009 343 350 9 9	MSD FRANCE	FR

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sécable				
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 558 452 0 1	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 492 3 0	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 338 524 2 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 459 1 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 082 4 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 449 6 4	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 461 6 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 083 0 5	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 491 7 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 085 3 4	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 087 6 3	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 458 5 5	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 495 2 0	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 088 2 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 447 3 5	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 358 495 8 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 450 4 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 494 6 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 084 7 3	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 574 496 9 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 095 9 3	MSD FRANCE	FR

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COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 091 3 5	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 093 6 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 563 580 3 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 090 7 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 089 9 2	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 574 497 5 9	MSD FRANCE	FR
COZAAR 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	34009 393 134 8 1	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 094 2 5	MSD FRANCE	FR
Losar Teva® 25 mg Filmtabletten Wirkstoff: Losartan-Kalium	DE/H/0848/002	66960.00.00	TEVA GMBH	DE
Losartan-AbZ 25 mg Filmtabletten	DE/H/3571/002	87048.00.00	ABZ-PHARMA GMBH	DE
Cozaar® 2,5 mg/ml Pulver und Lösungsmittel zur Herstellung einer Suspension zum Einnehmen	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 12,5 mg, Filmtabletten	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 12,5 mg, Filmtabletten	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 50 mg, Filmtabletten	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 50 mg, Filmtabletten	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg, Filmtabletten	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg, Filmtabletten	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 50 mg filmomhulde tabletten	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 50 mg filmomhulde tabletten	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 50 mg, comprimés pelliculés	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 50 mg, comprimés pelliculés	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 100 mg, comprimés pelliculés	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 100 mg, comprimés pelliculés	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg filmomhulde tabletten	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg filmomhulde tabletten	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 12,5 mg filmomhulde tabletten	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 12,5 mg, comprimés pelliculés	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 12,5 mg filmomhulde tabletten	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
Cozaar® 12,5 mg, comprimés pelliculés	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE

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COZAAR® 2,5 mg/ml poeder en oplosmiddel voor orale suspensie	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE
LOSAPREX 2,5 mg/ml polvere e solvente per sospensione orale	not available	029393055	ALFASIGMA S.P.A.	IT
LOSAPREX 100 mg compresse rivestite con film	NL/H/1457/003	029393042	ALFASIGMA S.P.A.	IT
LOSAPREX 12,5 mg compresse rivestite con film	NL/H/1457/001	029393030	ALFASIGMA S.P.A.	IT
LOSAPREX 12,5 mg compresse rivestite con film	NL/H/1457/001	029393028	ALFASIGMA S.P.A.	IT
LOSAPREX 50 mg compresse rivestite con film	NL/H/1457/002	029393016	ALFASIGMA S.P.A.	IT
Cozaar 100 mg comprimidos recubiertos con película	NL/H/1457/003	64.971	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Cozaar® 2,5 mg/ml Pulver und Lösungsmittel zur Herstellung einer Suspension zum Einnehmen	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 12,5 mg, Filmtabletten	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 12,5 mg, Filmtabletten	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmdragerade tabletter	NL/H/1457/003	16602	MERCK SHARP & DOHME BV	FI
Cozaar 12,5 mg filmdragerade tabletter	NL/H/1457/001	12916	MERCK SHARP & DOHME BV	FI
Cozaar 50 mg filmdragerade tabletter	NL/H/1457/002	11637	MERCK SHARP & DOHME BV	FI
COZAAR® 50 mg, Filmtabletten	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 50 mg, Filmtabletten	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg, Filmtabletten	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
COZAAR® 100 mg, Filmtabletten	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/030	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/011	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/029	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cosaar 50 mg Filmtabletten	NL/H/1457/002	1-21306	MERCK SHARP & DOHME GES.M.B.H.	AT

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Cosaar 100 mg Filmtabletten	NL/H/1457/003	1-24232	MERCK SHARP & DOHME GES.M.B.H.	AT
Cosaar 12,5 mg Filmtabletten	NL/H/1457/001	1-22927	MERCK SHARP & DOHME GES.M.B.H.	AT
COZAAR® 50 mg, comprimés pelliculés	NL/H/1457/002	1995040196	MSD BELGIUM BVBA/SPRL	LU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/01	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/02	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/03	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/04	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/05	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317683	MERCK SHARP & DOHME, LDA.	PT
Cozaar® 50 mg filmomhulde tabletten	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/06	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg tablett, filmdrasjert	NL/H/1457/002	00-8059	MERCK SHARP & DOHME BV	NO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/26	MSD PHARMA HUNGARY KFT.	HU
COZAAR, 50 mg, tabletki powlekane	NL/H/1457/002	R/6744	MSD POLSKA SP. Z O.O.	PL
Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317584	MERCK SHARP & DOHME, LDA.	PT
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/07	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/23	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/08	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/09	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg kalvopäällysteiset tabletit	NL/H/1457/002	11637	MERCK SHARP & DOHME BV	FI
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/18	MSD PHARMA HUNGARY KFT.	HU
Cozaar® 50 mg filmomhulde tabletten	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/21	MSD PHARMA HUNGARY 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
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/14	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/10	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/15	MSD PHARMA HUNGARY KFT.	HU
Cozaar® 50 mg, comprimés pelliculés	NL/H/1457/002	BE173074	MSD BELGIUM BVBA/SPRL	BE
Lortaan 50 mg compresse rivestite con film	NL/H/1457/002	029384017	MSD ITALIA S.R.L.	IT
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/11	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/13	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/12	MERCK SHARP & DOHME ROMANIA SRL	RO
LORZAAR® PROTECT 50 mg Filmtabletten Wirkstoff: Losartan-Kalium	NL/H/1457/002	41546.01.00	MSD SHARP & DOHME GMBH	DE
Cozaar® 50 mg film-coated tablets	NL/H/1457/002	PL 0025/0324	MERCK SHARP & DOHME LTD.	UK
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/12	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/13	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/17	MSD PHARMA HUNGARY KFT.	HU
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/20	MSD PHARMA HUNGARY KFT.	HU
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/16	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg film-coated tablets	NL/H/1457/002	PA1286/4/2	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/14	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmdragerade tabletter	NL/H/1457/002	12209	MERCK SHARP & DOHME BV	SE
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/15	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar, 50 mg, comprimidos revestidos por película	NL/H/1457/002	2317782	MERCK SHARP & DOHME, LDA.	PT
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/24	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/16	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/19	MSD PHARMA HUNGARY KFT.	HU
Cozaar® 50 mg, comprimés pelliculés	NL/H/1457/002	BE341047	MSD BELGIUM BVBA/SPRL	BE
COZAAR 50 mg filmuhúðaðar töflur	NL/H/1457/002	930306	MERCK SHARP & DOHME BV	IS

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/17	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg filmomhulde tabletten	NL/H/1457/002	RVG 17617	MERCK SHARP & DOHME BV	NL
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/25	MSD PHARMA HUNGARY KFT.	HU
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/22	MSD PHARMA HUNGARY KFT.	HU
COZAAR 50 mg film-coated tablets	NL/H/1457/002	MA058/01302	MERCK SHARP & DOHME LTD.	MT
COZAAR 50 mg comprimate filmate	NL/H/1457/002	2977/2010/18	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 50 mg comprimidos recubiertos con película	NL/H/1457/002	60.913	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
COZAAR, 100 mg, tabletki powlekane	NL/H/1457/003	10734	MSD POLSKA SP. Z O.O.	PL
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982881	MERCK SHARP & DOHME, LDA.	PT
COZAAR 100 mg film-coated tablets	NL/H/1457/003	MA058/01303	MERCK SHARP & DOHME LTD.	MT
Cozaar 100 mg filmdragerade tabletter	NL/H/1457/003	17386	MERCK SHARP & DOHME BV	SE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/34	MSD PHARMA HUNGARY KFT.	HU
Lortaan 100 mg compresse rivestite con film	NL/H/1457/003	029384043	MSD ITALIA S.R.L.	IT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982485	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/31	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/37	MSD PHARMA HUNGARY KFT.	HU
Cozaar® 100 mg, comprimés pelliculés	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/27	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982584	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg kalvopäällysteiset tabletit	NL/H/1457/003	16602	MERCK SHARP & DOHME BV	FI
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982782	MERCK SHARP & DOHME, LDA.	PT
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982386	MERCK SHARP & DOHME, LDA.	PT
COZAAR 100 mg film-coated tablets	NL/H/1457/003	PA1286/4/3	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/30	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/40	MSD PHARMA HUNGARY 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
Cozaar® 100 mg, comprimés pelliculés	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/38	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/35	MSD PHARMA HUNGARY KFT.	HU
COZAAR 100 mg filmuhúðaðar töflur	NL/H/1457/003	IS/1/02/103/01	MERCK SHARP & DOHME BV	IS
COZAAR® 100 mg filmomhulde tabletten	NL/H/1457/003	BE341056	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/39	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/29	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/36	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg tablett, filmdrasjert	NL/H/1457/003	01-6335	MERCK SHARP & DOHME BV	NO
Cozaar 100 mg filmomhulde tabletten	NL/H/1457/003	RVG 26791	MERCK SHARP & DOHME BV	NL
Cozaar® 100 mg film-coated tablets	NL/H/1457/003	PL 0025/0416	MERCK SHARP & DOHME LTD.	UK
Cozaar 100 mg, 100 mg, comprimidos revestidos por película	NL/H/1457/003	3982683	MERCK SHARP & DOHME, LDA.	PT
LORZAAR® PROTECT 100 mg Filmtabletten Wirkstoff: Losartan-Kalium	NL/H/1457/003	41546.02.00	MSD SHARP & DOHME GMBH	DE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/28	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/32	MSD PHARMA HUNGARY KFT.	HU
COZAAR® 100 mg filmomhulde tabletten	NL/H/1457/003	BE235627	MSD BELGIUM BVBA/SPRL	BE
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/33	MSD PHARMA HUNGARY KFT.	HU
COZAAR® 100 mg, comprimés pelliculés	NL/H/1457/003	2007019153	MSD BELGIUM BVBA/SPRL	LU
Cozaar IC, 12,5 mg, comprimidos revestidos por película	NL/H/1457/001	3335080	MERCK SHARP & DOHME, LDA.	PT
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/01	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/04	MSD PHARMA HUNGARY KFT.	HU
КОЗАР 12,5 mg филмирани таблетки	NL/H/1457/001	20090151	MERCK SHARP & DOHME BULGARIA EOOD	BG
Cozaar® 12,5 mg filmomhulde tabletten	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
Lortaan 12,5 mg compresse rivestite con film	NL/H/1457/001	029384031	MSD ITALIA S.R.L.	IT
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/02	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/03	MERCK SHARP & DOHME ROMANIA SRL	RO
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/04	MERCK SHARP & DOHME	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
			ROMANIA SRL	
Cozaar® 12,5 mg, comprimés pelliculés	NL/H/1457/001	BE201144	MSD BELGIUM BVBA/SPRL	BE
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/05	MERCK SHARP & DOHME ROMANIA SRL	RO
Lortaan 12,5 mg compresse rivestite con film	NL/H/1457/001	029384029	MSD ITALIA S.R.L.	IT
Cozaar 12,5 mg kalvopäällysteiset tabletit	NL/H/1457/001	12916	MERCK SHARP & DOHME BV	FI
COZAAR 12.5 mg film-coated tablets	NL/H/1457/001	MA058/01305	MERCK SHARP & DOHME LTD.	MT
Cozaar® 12,5 mg filmomhulde tabletten	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
COZAAR 12,5 mg filmuhúðaðar töflur	NL/H/1457/001	970239	MERCK SHARP & DOHME BV	IS
Cozaar® 12,5 mg, comprimés pelliculés	NL/H/1457/001	BE341031	MSD BELGIUM BVBA/SPRL	BE
COZAAR, 12,5 mg, tabletki powlekane	NL/H/1457/001	9978	MSD POLSKA SP. Z O.O.	PL
Cozaar 12,5 mg filmdragerade tabletter	NL/H/1457/001	13718	MERCK SHARP & DOHME BV	SE
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/06	MERCK SHARP & DOHME ROMANIA SRL	RO
LORZAAR® START 12,5 mg Filmtabletten Wirkstoff: Losartan-Kalium	NL/H/1457/001	41542.00.00	MSD SHARP & DOHME GMBH	DE
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/07	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar, filmovertrukne tabletter	NL/H/1457/001	18978	MERCK SHARP & DOHME BV	DK
Cozaar IC, 12,5 mg, comprimidos revestidos por película	NL/H/1457/001	3335189	MERCK SHARP & DOHME, LDA.	PT
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/07	MSD PHARMA HUNGARY KFT.	HU
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/06	MSD PHARMA HUNGARY KFT.	HU
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/01	MSD PHARMA HUNGARY KFT.	HU
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/08	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/10	MSD PHARMA HUNGARY KFT.	HU
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/09	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/05	MSD PHARMA HUNGARY KFT.	HU
COZAAR 12,5 mg comprimate filmate	NL/H/1457/001	2976/2010/10	MERCK SHARP & DOHME ROMANIA SRL	RO
Cozaar® 12.5 mg film-coated tablets	NL/H/1457/001	PL 0025/0515	MERCK SHARP & DOHME LTD.	UK
COZAAR® 12,5 mg, comprimés pelliculés	NL/H/1457/001	1999060006	MSD BELGIUM BVBA/SPRL	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
COZAAR 12.5 mg film-coated tablets	NL/H/1457/001	PA1286/4/1	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/11	MSD PHARMA HUNGARY KFT.	HU
Cozaar 12,5 mg INICIO comprimidos recubiertos con película	NL/H/1457/001	62.117	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Cozaar 12,5 mg filmomhulde tabletten	NL/H/1457/001	RVG 101836	MERCK SHARP & DOHME BV	NL
Cozaar 12,5 mg tablett, filmdrasjert	NL/H/1457/001	97-2003	MERCK SHARP & DOHME BV	NO
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/08	MSD PHARMA HUNGARY KFT.	HU
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/09	MSD PHARMA HUNGARY KFT.	HU
COZAAR® 2.5 mg/ml powder and solvent for oral suspension	NL/H/1457/004	PL 00025/0530	MERCK SHARP & DOHME LTD.	UK
COZAAR® 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	2009030031	MSD BELGIUM BVBA/SPRL	LU
Cozaar 2,5 mg/ml pó e solvente para suspensão oral	NL/H/1457/004	5178207	MERCK SHARP & DOHME, LDA.	PT
Cozaar 2,5 mg/ml poeder en oplosmiddel voor orale suspensie	NL/H/1457/004	RVG 103001	MERCK SHARP & DOHME BV	NL
COZAAR 2.5 mg/ml powder and solvent for oral suspension	NL/H/1457/004	PA1286/4/4	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
COZAAR® 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE
LORTAAN 2,5 mg/ml polvere e solvente per sospensione orale	NL/H/1457/004	029384056	MSD ITALIA S.R.L.	IT
LORZAAR® 2,5 mg/ml Pulver und Lösungsmittel zur Herstellung einer Suspension zum Einnehmen Wirkstoff: Losartan-Kalium	NL/H/1457/004	77187.00.00	MSD SHARP & DOHME GMBH	DE
COZAAR 2,5 mg/ml mixtúruduft og leysir, dreifa	NL/H/1457/004	IS/1/09/063/01	MERCK SHARP & DOHME BV	IS
Cozaar 2,5 mg/ml pulver og væske til mikstur, suspensjon	NL/H/1457/004	09-6574	MERCK SHARP & DOHME BV	NO
COZAAR® 2,5 mg/ml poeder en oplosmiddel voor orale suspensie	NL/H/1457/004	BE334521	MSD BELGIUM BVBA/SPRL	BE
Cozaar, filmovertrukne tabletter	NL/H/1457/002	15844	MERCK SHARP & DOHME BV	DK
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 343 350 9 9	MSD FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 558 452 0 1	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 492 3 0	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 338 524 2 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 459 1 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 082 4 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 449 6 4	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 461 6 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 083 0 5	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 491 7 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 085 3 4	MSD FRANCE	FR
Cozaar, filmovertrukne tabletter	NL/H/1457/003	32842	MERCK SHARP & DOHME BV	DK
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 087 6 3	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 371 458 5 5	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 495 2 0	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 088 2 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 447 3 5	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 358 495 8 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 371 450 4 6	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 574 494 6 9	MSD FRANCE	FR
COZAAR 50 mg, comprimé pelliculé sécable	NL/H/1457/002	34009 392 084 7 3	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 574 496 9 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 095 9 3	MSD FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 091 3 5	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 093 6 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 563 580 3 8	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 090 7 4	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 089 9 2	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 574 497 5 9	MSD FRANCE	FR
COZAAR 2,5 mg/ml, poudre et solvant pour suspension buvable	NL/H/1457/004	34009 393 134 8 1	MSD FRANCE	FR
COZAAR 100 mg, comprimé pelliculé	NL/H/1457/003	34009 392 094 2 5	MSD FRANCE	FR
COZAAR® 50 mg, potahované tablety	NL/H/1457/002	58/607/08-C	MERCK SHARP & DOHME BV	CZ
COZAAR® 100 mg, potahované tablety	NL/H/1457/003	58/608/08-C	MERCK SHARP & DOHME BV	CZ
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/002	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/003	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/004	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/005	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/006	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/007	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/008	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/009	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 12,5 mg filmsko obložene tablete	NL/H/1457/001	H/95/00429/010	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/012	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/013	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/014	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/015	MERCK SHARP & DOHME	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
			INOVATIVNA ZDRAVILA D.O.O.	
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/016	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/017	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/018	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/019	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/020	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/021	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/022	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/023	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/024	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/025	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/026	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/027	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 50 mg filmsko obložene tablete	NL/H/1457/002	H/95/00429/028	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/031	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/032	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/033	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/034	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/035	MERCK SHARP & DOHME	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
			INOVATIVNA ZDRAVILA D.O.O.	
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/036	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/037	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/038	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/039	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/040	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/041	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/042	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/043	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/044	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cozaar 100 mg filmsko obložene tablete	NL/H/1457/003	H/95/00429/045	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
COZAAR 12.5 mg επικαλυμμένα με λεπτό υμένιο δισκία.	NL/H/1457/001	40422/31-05-2013	VIANEX S.A.	GR
COZAAR 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1457/002	38346/31-05-2013	VIANEX S.A.	GR
COZAAR 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1457/003	43376/ 31-05-2013	VIANEX S.A.	GR
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/43	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/47	MSD PHARMA HUNGARY KFT.	HU
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/44	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/45	MSD PHARMA HUNGARY KFT.	HU
Cozaar 12,5 mg filmtabletta	NL/H/1457/001	OGYI-T-6454/41	MSD PHARMA HUNGARY KFT.	HU
Cozaar 100 mg filmtabletta	NL/H/1457/003	OGYI-T-6454/46	MSD PHARMA HUNGARY KFT.	HU
Cozaar 50 mg filmtabletta	NL/H/1457/002	OGYI-T-6454/42	MSD PHARMA HUNGARY KFT.	HU
Losartan-CT 25 mg Filmtabletten	DE/H/3570/002	87044.00.00	ABZ-PHARMA GMBH	DE
Losartán ratio 25 mg comprimidos	DE/H/3570/002	78.475	RATIOPHARM ESPANA SA	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				
Losartán ratio 100 mg comprimidos recubiertos con película EFG	DE/H/3570/005	79923	RATIOPHARM ESPANA SA	ES
Losatrix 50 mg filmuhúðaðar töflur.	UK/H/0925/003	IS/1/13/093/02	RATIOPHARM GMBH	IS
Losatrix 25 mg filmuhúðaðar töflur.	UK/H/0925/002	IS/1/13/093/01	RATIOPHARM GMBH	IS
Losatrix 100 mg filmuhúðaðar töflur.	UK/H/0925/004	IS/1/13/093/03	RATIOPHARM GMBH	IS
Arbartan 100 mg filmtabletta	UK/H/0925/004	OGYI-T-20536/07	TEVA GYÓGYSZERGYÁR ZRT	HU
Arbartan 25 mg filmtabletta	UK/H/0925/002	OGYI-T-20536/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Arbartan 50 mg filmtabletta	UK/H/0925/003	OGYI-T-20536/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan ratiopharm 100 mg Filmtabletten	UK/H/0925/004	1-27322	TEVA B.V	AT
Losartan ratiopharm 25 mg Filmtabletten	UK/H/0925/002	1-27320	TEVA B.V	AT
Losartan ratiopharm 50 mg Filmtabletten	UK/H/0925/003	1-27321	TEVA B.V	AT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038112064	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038112052	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038112049	TEVA ITALIA S.R.L.	IT
Losartan Teva 100 mg compresse rivestite con film	UK/H/0925/004	038112254	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038112203	TEVA ITALIA S.R.L.	IT
Losatrix 100 mg kalvopäällysteiset tabletit	UK/H/0925/004	22360	RATIOPHARM GMBH	FI
Losartan Teva 100 mg compresse rivestite con film	UK/H/0925/004	038112278	TEVA ITALIA S.R.L.	IT
Losartan Teva 100 mg compresse rivestite con film	UK/H/0925/004	038112266	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038112215	TEVA ITALIA S.R.L.	IT
Losatrix 50 mg kalvopäällysteiset tabletit	UK/H/0925/003	22359	RATIOPHARM GMBH	FI
Losatrix 25 mg kalvopäällysteiset tabletit	UK/H/0925/002	22358	RATIOPHARM GMBH	FI
Losartan Teva 50 mg compresse rivestite con film	UK/H/0925/003	038112241	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite	UK/H/0925/002	038112191	TEVA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
con film				
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	38112049	TEVA ITALIA S.R.L.	IT
Losartan Teva 50 mg compresse rivestite con film	UK/H/0925/003	038112227	TEVA ITALIA S.R.L.	IT
Losartan Teva 50 mg compresse rivestite con film	UK/H/0925/003	038112239	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098226	TEVA ITALIA S.R.L.	IT
Losartan Teva 100 mg tabletter, filmdrasjerte	UK/H/0925/004	06-4223	TEVA SWEDEN AB	NO
Losartan potassium 25 mg Film-coated Tablets	UK/H/0925/002	PL 00289/0963	TEVA UK LIMITED	UK
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098012	TEVA ITALIA S.R.L.	IT
Losartan Teva 50 mg tabletter, filmdrasjerte	UK/H/0925/003	06-4222	TEVA SWEDEN AB	NO
Losatrix 100 mg kalvopäällysteiset tabletit	UK/H/0925/004	22360	RATIOPHARM GMBH	FI
Losartankalium ”Teva”, filmovertrukne tabletter	UK/H/0925/003	39586	TEVA DENMARK A/S	DK
Losatrix 25 mg kalvopäällysteiset tabletit	UK/H/0925/002	22358	RATIOPHARM GMBH	FI
Losartan Teva 25 mg tabletter, filmdrasjerte	UK/H/0925/002	06-4221	TEVA SWEDEN AB	NO
Losartan Teva 50 mg filmdragerade tabletter	UK/H/0925/003	24078	TEVA SWEDEN AB	SE
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098125	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098024	TEVA ITALIA S.R.L.	IT
LOSARTAN TEVA 100 mg, comprimé pelliculé sécable	UK/H/0925/004	NL33084	TEVA SANTÉ	FR
Losartan Teva 100 mg filmdragerade tabletter	UK/H/0925/004	24079	TEVA SWEDEN AB	SE
Losartankalium ”Teva”, filmovertrukne tabletter	UK/H/0925/004	39587	TEVA DENMARK A/S	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098214	TEVA ITALIA S.R.L.	IT
LOSARTAN TEVA 50 mg, comprimé pelliculé sécable	UK/H/0925/003	NL33083	TEVA SANTÉ	FR
Losatrix 50 mg kalvopäällysteiset tabletit	UK/H/0925/003	22359	RATIOPHARM GMBH	FI
Losartan Potassium 100 mg Film-coated Tablets	UK/H/0925/004	PL 00289/0965	TEVA UK LIMITED	UK
Losartan ratiopharm, 50 mg, comprimidos revestidos por película	UK/H/0925/003	5089701	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098277	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089651	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089669	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098137	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098190	TEVA ITALIA S.R.L.	IT
Losartan potassium 50 mg Film-coated Tablets	UK/H/0925/003	PL 00289/0964	TEVA UK LIMITED	UK
Losartankalium 25 mg PCH, filmomhulde tabletten	UK/H/0925/002	RVG 34188	PHARMACHEMIE B.V	NL
Losartankalium 50 mg PCH, filmomhulde tabletten	UK/H/0925/003	RVG 34189	PHARMACHEMIE B.V	NL
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098149	TEVA ITALIA S.R.L.	IT
Losartankalium 100 mg PCH, filmomhulde tabletten	UK/H/0925/004	RVG 34190	PHARMACHEMIE B.V	NL
Arbartan 25 mg filmtabletta	UK/H/0925/002	OGYI-T-20536/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098063	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 50 mg,	UK/H/0925/003	5089644	RATIOPHARM-COMERCIO E	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
comprimidos revestidos por película			INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098036	TEVA ITALIA S.R.L.	IT
Arbartan 50 mg filmtabletta	UK/H/0925/003	OGYI-T-20536/06	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098238	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089727	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098289	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098101	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089610	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098291	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098099	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098152	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 50 mg, comprimidos revestidos por película	UK/H/0925/003	5089602	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan ratiopharm, 50 mg, comprimidos revestidos por película	UK/H/0925/003	5089578	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098048	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 50 mg, comprimidos revestidos por película	UK/H/0925/003	5089677	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098253	TEVA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098075	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098051	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098176	TEVA ITALIA S.R.L.	IT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089719	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan ratiopharm, 100 mg, comprimidos revestidos por película	UK/H/0925/004	5089628	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098265	TEVA ITALIA S.R.L.	IT
Arbartan 50 mg filmtabletta	UK/H/0925/003	OGYI-T-20536/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Arbartan 100 mg filmtabletta	UK/H/0925/004	OGYI-T-20536/09	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan ratiopharm, 50 mg, comprimidos revestidos por película	UK/H/0925/003	5089636	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098087	TEVA ITALIA S.R.L.	IT
Arbartan 25 mg filmtabletta	UK/H/0925/002	OGYI-T-20536/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098303	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098240	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098164	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098113	TEVA ITALIA S.R.L.	IT
Arbartan 100 mg filmtabletta	UK/H/0925/004	OGYI-T-20536/08	TEVA GYÓGYSZERGYÁR ZRT	HU
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098188	TEVA ITALIA S.R.L.	IT
Losartan Teva 25 mg compresse rivestite con film	UK/H/0925/002	038098202	TEVA ITALIA S.R.L.	IT
Neo-lotan 12,5 mg compresse rivestite	not available	029385034	NEOPHARMED GENTILI S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
con film				
Neo-lotan 12,5 mg compresse rivestite con film	not available	029385022	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 50 mg compresse rivestite con film	not available	029385010	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 100 mg compresse rivestite con film	not available	029385046	NEOPHARMED GENTILI S.R.L.	IT
Neo-lotan 2.5 mg/ml polvere e solvente per sospensione orale	not available	029385059	NEOPHARMED GENTILI S.R.L.	IT
Losartan "Aurobindo", filmovertrukne tabletter	NL/H/3056/001	54075	AUROBINDO PHARMA (MALTA) LIMITED	DK
Losartankalium Aurobindo 25 mg, filmomhulde tabletten	NL/H/3056/001	RVG 105815	AUROBINDO PHARMA B.V.	NL
Losartán Aurobindo 25 mg comprimidos recubiertos con película	NL/H/3056/001	79.245	LABORATORIOS AUROBINDO S.L.U.	ES
LOSAPREX 2,5 mg/ml polvere e solvente per sospensione orale	not available	029393055	ALFASIGMA S.P.A.	IT
LOSAPREX 100 mg compresse rivestite con film	NL/H/1457/003	029393042	ALFASIGMA S.P.A.	IT
LOSAPREX 12,5 mg compresse rivestite con film	NL/H/1457/001	029393030	ALFASIGMA S.P.A.	IT
LOSAPREX 12,5 mg compresse rivestite con film	NL/H/1457/001	029393028	ALFASIGMA S.P.A.	IT
LOSAPREX 50 mg compresse rivestite con film	NL/H/1457/002	029393016	ALFASIGMA S.P.A.	IT