04 October 2018
EMA/715149/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: hydrochlorothiazide / losartan

Procedure no.: PSUSA/00001655/201802

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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
Losartán/Hidroclorotiazida Aurobindo 100 mg/12,5 mg comprimidos recubiertos con película	PT/H/1010/002	78.072	LABORATORIOS AUROBINDO S.L.U.	ES
Losartan potassium/Hydrochlorothiazide 50 mg/12.5 mg film-coated tablet	not available	PL 40378/0086	APTIL PHARMA LIMITED	UK
Losartan potassium/Hydrochlorothiazide 100 mg/25 mg film-coated tablet	not available	PL 40378/0087	APTIL PHARMA LIMITED	UK
LOSARTAN HYDROCHLOROTHIAZIDE EG 100 mg/25 mg, comprimé pelliculé	not available	NL34722	EG LABO LABORATOIRES EUROGENERICS	FR
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	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
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530889	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530988	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5188230	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5627781	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5627682	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530780	MERCK SHARP & DOHME, LDA.	PT
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3124989	MERCK SHARP & DOHME, LDA.	PT
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3125085	MERCK SHARP & DOHME, LDA.	PT
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3124880	MERCK SHARP & DOHME, LDA.	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
FORTZAAR® 100 mg/25 mg filmom obložene tablete	not available	HR-H-433648826	MERCK SHARP & DOHME D.O.O.	HR
HYZAAR® 50 mg/12,5 mg filmom obložene tablete	not available	HR-H-667284558	MERCK SHARP & DOHME D.O.O.	HR
HYZAAR® 100 mg/12,5 mg filmom obložene tablete	not available	HR-H-883944358	MERCK SHARP & DOHME D.O.O.	HR
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	NEOPHARMED GENTILI S.R.L.	IT
Losartán/Hidroclorotiazida Krka 100 mg/12,5 mg comprimidos recubiertos con película EFG	CZ/H/0231/001	72937	KRKA, D.D., NOVO MESTO	ES
Losartan/HCT Krka 100 mg/12,5 mg Filmtabletten	CZ/H/0231/001	1-29766	KRKA, D.D., NOVO MESTO	AT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lorista HL, 100 mg + 12,5 mg, tabletki powlekane	CZ/H/0231/001	17486	KRKA, D.D., NOVO MESTO	PL
Лориста HL 100 mg/12,5 mg филмирани таблетки	CZ/H/0231/001	20100620	KRKA, D.D., NOVO MESTO	BG
Losartan/Hydrochlorothiazide Krka 100 mg/12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0231/001	020921	KRKA, D.D., NOVO MESTO	CY
Lorista H 100 mg/12,5 mg, õhukese polümeerikattega tabletid	CZ/H/0231/001	704910	KRKA, D.D., NOVO MESTO	EE
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/01	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/02	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/03	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/04	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/05	KRKA, D.D., NOVO MESTO	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
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/06	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/07	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/08	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/09	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/10	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/11	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/12	KRKA, D.D., NOVO MESTO	RO
Lorista HL 100 mg/12,5 mg comprimate filmate	CZ/H/0231/001	10425/2017/13	KRKA, D.D., NOVO MESTO	RO
Loortan® Plus 100 mg/12,5 mg, comprimés pelliculés	not available	2010040033	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
Loortan® Plus 50 mg/12,5 mg, comprimés pelliculés	not available	2004118438	MSD BELGIUM BVBA/SPRL	LU
Loortan® Plus Forte 100 mg/25 mg, comprimés pelliculés	not available	2004118439	MSD BELGIUM BVBA/SPRL	LU
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	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
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	NEOPHARMED GENTILI S.R.L.	IT
LOSAZID 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	031497047	ALFASIGMA S.P.A.	IT
LOSAZID 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	031497035	ALFASIGMA S.P.A.	IT
LOSAZID 50 mg + 12,5 mg compresse rivestite con film	NL/H/1458/001	031497050	ALFASIGMA S.P.A.	IT
LOSAZID 50 mg + 12,5 mg compresse rivestite con film	NL/H/1458/001	031497023	ALFASIGMA S.P.A.	IT
Losartán/Hidroclorotiazida TecniGen 100 mg/12,5 mg comprimidos recubiertos con película	not available	73.106	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392123	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite	UK/H/1685/002	041392135	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
con film				
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392147	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392150	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392162	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392174	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392186	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 e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392198	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392200	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392212	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392224	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392376	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite	UK/H/1685/002	041392364	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
con film				
Losartan e Idroclorotiazide Aurobindo 100 mg/12,5 mg compresse rivestite con film	UK/H/1685/002	041392414	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Losartán/Hidroclorotiazida ratio 100 mg/12,5 mg comprimidos recubiertos con película	UK/H/5001/001	76180	RATIOPHARM ESPANA SA	ES
Cozaar® Plus 50 mg/12,5 mg Filmtabletten	NL/H/1458/001	BE190915	MSD BELGIUM BVBA/SPRL	BE
Cozaar® Plus 50 mg/12,5 mg Filmtabletten	NL/H/1458/001	BE342754	MSD BELGIUM BVBA/SPRL	BE
Cozaar® Plus 100 mg/12,5 mg Filmtabletten	NL/H/1458/002	BE342781	MSD BELGIUM BVBA/SPRL	BE
Cozaar® Plus 100 mg/12,5 mg Filmtabletten	NL/H/1458/002	BE342772	MSD BELGIUM BVBA/SPRL	BE
Cozaar Comp 100 mg/12,5 mg filmdragerade tabletter	NL/H/1458/002	21023	MERCK SHARP & DOHME BV	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Comp Forte 100 mg/25 mg filmdragerade tabletter	NL/H/1458/003	13738	MERCK SHARP & DOHME BV	FI
Cozaar Comp 50 mg/12,5 mg filmdragerade tabletter	NL/H/1458/001	12140	MERCK SHARP & DOHME BV	FI
Hyzaar Forte 100 mg/25 mg filmtabletta	NL/H/1458/003	OGYI-T-6976/04	MSD PHARMA HUNGARY KFT.	HU
Cozaar® Plus Forte 100 mg/25 mg Filmtabletten	NL/H/1458/003	BE229302	MSD BELGIUM BVBA/SPRL	BE
Cozaar® Plus Forte 100 mg/25 mg Filmtabletten	NL/H/1458/003	BE342763	MSD BELGIUM BVBA/SPRL	BE
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/028	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/017	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Cosaar plus 50 mg/12,5 mg Filmtabletten	NL/H/1458/001	1-21713	MERCK SHARP & DOHME GES.M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Comp 50 mg/12,5 mg filmdragerade tabletter	NL/H/1458/001	12771	MERCK SHARP & DOHME BV	SE
Cozaar Comp., filmovertrukne tabletter	NL/H/1458/001	17535	MERCK SHARP & DOHME BV	DK
Hyzaar 50 mg/12,5 mg filmomhulde tabletten	NL/H/1458/001	RVG 19269	MERCK SHARP & DOHME BV	NL
Cozaar Comp 50 mg/12,5 mg filmuhúðaðar töflur	NL/H/1458/001	950133	MERCK SHARP & DOHME BV	IS
Cozaar Comp. 100 mg/12,5 mg, filmovertrukne tabletter	NL/H/1458/002	38633	MERCK SHARP & DOHME BV	DK
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530889	MERCK SHARP & DOHME, LDA.	PT
Cozaar Comp 50 mg/12,5 mg kalvopäällysteiset tabletit	NL/H/1458/001	12140	MERCK SHARP & DOHME BV	FI
HYZAAR 50 mg / 12.5 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/001	17142	MERCK SHARP & DOHME BV	CY
Cozaar Plus 50 mg/12,5 mg, comprimés pelliculés	NL/H/1458/001	BE190915	MSD BELGIUM BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Hyzaar 50 mg/12,5 mg filmtabletta	NL/H/1458/001	OGYI-T-6976/01	MSD PHARMA HUNGARY KFT.	HU
COZAAR PLUS 50mg/12,5 mg, filmomhulde tabletten	NL/H/1458/001	BE342754	MSD BELGIUM BVBA/SPRL	BE
Cozaar Plus 50 mg/12,5 mg, comprimés pelliculés	NL/H/1458/001	BE342754	MSD BELGIUM BVBA/SPRL	BE
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530988	MERCK SHARP & DOHME, LDA.	PT
HIZAAR 50 MG + 12,5 MG COMPRESSE RIVESTITE CON FILM	NL/H/1458/001	032079055	MSD ITALIA S.R.L.	IT
LORZAAR® PLUS 50/12,5 mg Filmtabletten Wirkstoffe: Losartan-Kalium und Hydrochlorothiazid	NL/H/1458/001	36017.00.00	MSD SHARP & DOHME GMBH	DE
LORZAAR® PLUS forte 100/12,5 mg Filmtabletten Wirkstoffe: Losartan-Kalium und Hydrochlorothiazid	NL/H/1458/002	64982.00.00	MSD SHARP & DOHME GMBH	DE
HYZAAR 100 mg / 12.5 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/002	21027	MERCK SHARP & DOHME BV	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Comp 100 mg/12,5 mg filmdragerade tabletter	NL/H/1458/002	22490	MERCK SHARP & DOHME BV	SE
Cozaar Comp 50 mg/12.5 mg film-coated tablets	NL/H/1458/001	PA 1286/1/2	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar Plus 50 mg/12,5 mg comprimidos recubiertos con película	NL/H/1458/001	61.600	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5188230	MERCK SHARP & DOHME, LDA.	PT
Cozaar Comp 100 mg/12,5 mg kalvopäällysteiset tabletit	NL/H/1458/002	21023	MERCK SHARP & DOHME BV	FI
Cozaar Comp 100 mg/12,5 mg filmdrasjerte tabletter	NL/H/1458/002	05-3601	MERCK SHARP & DOHME BV	NO
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5627781	MERCK SHARP & DOHME, LDA.	PT
HIZAAR 50MG + 12.5MG COMPRESSE RIVESTITE CON FILM	NL/H/1458/001	032079028	MSD ITALIA S.R.L.	IT
Cozaar Plus 50 mg/12,5 mg, filmomhulde tabletten	NL/H/1458/001	BE190915	MSD BELGIUM BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Plus 100 mg/12,5 mg, comprimés pelliculés	NL/H/1458/002	2010010009	MSD BELGIUM BVBA/SPRL	LU
Cozaar Plus 100 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/002	5627682	MERCK SHARP & DOHME, LDA.	PT
Cozaar Plus 50 mg/12,5 mg comprimidos revestidos por película	NL/H/1458/001	2530780	MERCK SHARP & DOHME, LDA.	PT
Cozaar Comp® 50 mg/12.5 mg film-coated tablets	NL/H/1458/001	PL 00025/0338	MERCK SHARP & DOHME LTD.	UK
HYZAAR, 50 mg + 12,5 mg, tabletki powlekane	NL/H/1458/001	4336	MSD POLSKA SP. Z O.O.	PL
Cozaar Comp 100 mg/12.5 mg film-coated tablets	NL/H/1458/002	PA 1286/1/1	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Cozaar Plus 100 mg/12,5 mg, comprimés pelliculés	NL/H/1458/002	BE342781	MSD BELGIUM BVBA/SPRL	BE
HYZAAR, 100 mg + 12,5 mg, tabletki powlekane	NL/H/1458/002	18673	MSD POLSKA SP. Z O.O.	PL
Cozaar Comp Forte 100 mg/25 mg filmuhúðaðar töflur	NL/H/1458/003	IS/1/05/101/01	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 Plus 100 mg/12,5 mg, filmomhulde tabletten	NL/H/1458/002	BE342781	MSD BELGIUM BVBA/SPRL	BE
HIZAAR 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	032079030	MSD ITALIA S.R.L.	IT
Cozaar Plus 100 mg/12,5 mg, comprimés pelliculés	NL/H/1458/002	BE342772	MSD BELGIUM BVBA/SPRL	BE
Cozaar Plus Forte 100 mg/25 mg filmomhulde tabletten	NL/H/1458/003	BE229302	MSD BELGIUM BVBA/SPRL	BE
Cozaar Plus 100 mg/12,5 mg filmomhulde tabletten	NL/H/1458/002	RVG 32433	MERCK SHARP & DOHME BV	NL
Hyzaar Forte 100 mg/25 mg filmtabletta	NL/H/1458/003	OGYI-T-6976/03	MSD PHARMA HUNGARY KFT.	HU
Cozaar Plus 100 mg/12,5 mg, filmomhulde tabletten	NL/H/1458/002	BE342772	MSD BELGIUM BVBA/SPRL	BE
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3124989	MERCK SHARP & DOHME, LDA.	PT
Fortzaar 100 mg/25 mg Filmtabletten	NL/H/1458/003	1-23960	MERCK SHARP & DOHME GES.M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Comp. Forte, filmovertrukne tabletter	NL/H/1458/003	32905	MERCK SHARP & DOHME BV	DK
Cozaar Comp® 100 mg/25 mg film-coated tablets	NL/H/1458/003	PL 00025/0374	MERCK SHARP & DOHME LTD.	UK
Cozaar Comp 100 mg/25 mg film-coated tablets	NL/H/1458/003	PA 1286/1/3	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3125085	MERCK SHARP & DOHME, LDA.	PT
HIZAAR 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	032079042	MSD ITALIA S.R.L.	IT
Cozaar Plus Forte 100 mg/25 mg, comprimés pelliculés	NL/H/1458/003	BE229302	MSD BELGIUM BVBA/SPRL	BE
Cozaar Comp Forte 100 mg/25 mg filmdragerade tabletter	NL/H/1458/003	14654	MERCK SHARP & DOHME BV	SE
Cozaar Comp 100 mg/12,5 mg filmuhúðaðar töflur	NL/H/1458/002	IS/1/06/117/01	MERCK SHARP & DOHME BV	IS
FORTZAAR® 100/25 mg Filmtabletten|Wirkstoffe: Losartan-Kalium und Hydrochlorothiazid	NL/H/1458/003	50798.00.00	MSD SHARP & DOHME 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
Cozaar Comp Forte 100 mg/25 mg kalvopäällysteiset tabletit	NL/H/1458/003	13738	MERCK SHARP & DOHME BV	FI
Cozaar Plus Forte 100 mg/25 mg filmomhulde tabletten	NL/H/1458/003	BE342763	MSD BELGIUM BVBA/SPRL	BE
Cozaar Comp® 100 mg/12.5 mg film-coated tablets	NL/H/1458/002	PL 00025/0473	MERCK SHARP & DOHME LTD.	UK
Cozaar Comp Forte 100 mg/25 mg filmdrasjerte tabletter	NL/H/1458/003	98-5073	MERCK SHARP & DOHME BV	NO
Cozaar Plus 50 mg/12,5 mg, comprimés pelliculés	NL/H/1458/001	1997120929	MSD BELGIUM BVBA/SPRL	LU
Cozaar Plus Forte 100 mg/25 mg, comprimés pelliculés	NL/H/1458/003	2002040024	MSD BELGIUM BVBA/SPRL	LU
Fortzaar 100 mg/25 mg comprimidos recubiertos con película	NL/H/1458/003	63.473	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Fortzaar 100 mg/25 mg comprimidos revestidos por película	NL/H/1458/003	3124880	MERCK SHARP & DOHME, LDA.	PT
Fortzaar 100 mg/25 mg filmomhulde tabletten	NL/H/1458/003	RVG 23597	MERCK SHARP & DOHME BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cozaar Plus Forte 100 mg/25 mg, comprimés pelliculés	NL/H/1458/003	BE342763	MSD BELGIUM BVBA/SPRL	BE
Cozaar Comp 50 mg/12,5 mg filmdrasjerte tabletter	NL/H/1458/001	96-1305	MERCK SHARP & DOHME BV	NO
FORTZAAR 100 mg / 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/003	19292	MERCK SHARP & DOHME BV	CY
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 371 445 0 6	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 098 8 3	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 096 5 4	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 343 353 8 9	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 101 9 3	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 102 5 4	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
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 100 2 5	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 338 520 7 8	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 371 446 7 4	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 392 097 1 5	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 371 444 4 5	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 558 453 7 9	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 377 487 7 3	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 351 862 5 6	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 392 103 1 5	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
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 377 488 3 4	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 371 407 1 3	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 392 105 4 4	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 377 484 8 3	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 392 108 3 4	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 574 499 8 8	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 371 406 5 2	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 574 500 6 9	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 561 782 8 5	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
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 351 861 9 5	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 574 502 9 8	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 392 104 8 3	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 392 110 8 4	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 574 501 2 0	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 371 408 8 1	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 377 486 0 5	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 392 106 0 5	MSD FRANCE	FR
FORTZAAR 100 mg/12,5 mg, comprimé pelliculé	NL/H/1458/002	34009 377 485 4 4	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
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 574 503 5 9	MSD FRANCE	FR
FORTZAAR 100 mg/25 mg, comprimé pelliculé	NL/H/1458/003	34009 392 107 7 3	MSD FRANCE	FR
Hyzaar 50 mg/12,5 mg, comprimé pelliculé	NL/H/1458/001	34009 574 498 1 0	MSD FRANCE	FR
Forzaar 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	034310019	MSD ITALIA S.R.L.	IT
Forzaar 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	034310021	MSD ITALIA S.R.L.	IT
HYZAAR Forte, 100 mg + 25 mg, tabletki powlekane	NL/H/1458/003	9705	MSD POLSKA SP. Z O.O.	PL
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/002	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/003	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/004	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	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
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/005	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/006	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/007	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/008	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/009	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/010	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/011	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/012	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/013	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	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
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/014	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/015	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 50 mg/12,5 mg filmsko obložene tablete	NL/H/1458/001	H/97/00745/016	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/018	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/019	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/020	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/021	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/022	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/023	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	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
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/024	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/025	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/026	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Hyzaar 100 mg/12,5 mg filmsko obložene tablete	NL/H/1458/002	H/97/00745/027	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/029	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/030	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/031	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/032	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/033	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	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
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/034	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/035	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/036	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/037	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/038	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/039	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/040	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Fortzaar 100 mg/25 mg filmsko obložene tablete	NL/H/1458/003	H/97/00745/041	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
LOSAZID 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	031497047	ALFASIGMA 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
LOSAZID 100 mg + 25 mg compresse rivestite con film	NL/H/1458/003	031497035	ALFASIGMA S.P.A.	IT
LOSAZID 50 mg + 12,5 mg compresse rivestite con film	NL/H/1458/001	031497050	ALFASIGMA S.P.A.	IT
LOSAZID 50 mg + 12,5 mg compresse rivestite con film	NL/H/1458/001	031497023	ALFASIGMA S.P.A.	IT
HYZAAR 50 mg/12.5 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/001	39222/31-05-2013	VIANEX S.A.	GR
HYZAAR Forte 100 mg/12.5 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/002	39665/09-06-2015	VIANEX S.A.	GR
HYZAAR Extra Forte 100 mg/25 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1458/003	43379/31-05-2013	VIANEX S.A.	GR
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	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
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	NEOPHARMED GENTILI S.R.L.	IT
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 492 2 1	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 485 6 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 482 7 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 471 5 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 497 4 0	MSD FRANCE	FR
Entrizen/HCT 50 mg/12,5 mg filmomhulde tabletten	NL/H/1606/001	RVG 103818	MERCK SHARP & DOHME BV	NL
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé	NL/H/1606/003	34009 396 498 0 1	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
pelliculé				
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 575 757 0 0	MSD FRANCE	FR
Entrizen/HCT 100 mg/12,5 mg filmomhulde tabletten	NL/H/1606/002	RVG 103819	MERCK SHARP & DOHME BV	NL
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 575 760 1 1	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 476 7 8	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 399 357 9 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 575 756 4 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé	NL/H/1606/002	34009 396 488 5 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
pelliculé				
Entrizen/HCT 100 mg/25 mg filmomhulde tabletten	NL/H/1606/003	RVG 103820	MERCK SHARP & DOHME BV	NL
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 472 1 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 479 6 8	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 502 8 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 494 5 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 486 2 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé	NL/H/1606/001	34009 396 475 0 0	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
pelliculé				
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 474 4 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 499 7 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 501 1 1	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 495 1 1	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 489 1 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 487 9 8	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
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 473 8 8	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 575 755 8 8	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 481 0 1	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 480 4 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 575 759 3 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 491 6 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 496 8 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
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 575 758 7 8	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 50 mg/12,5 mg, comprimé pelliculé	NL/H/1606/001	34009 396 477 3 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 493 9 9	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/12,5 mg, comprimé pelliculé	NL/H/1606/002	34009 396 483 3 0	MSD FRANCE	FR
LOSARTAN/HYDROCHLOROTHIAZIDE ZENTIVA 100 mg/25 mg, comprimé pelliculé	NL/H/1606/003	34009 396 500 5 0	MSD FRANCE	FR
Losartan/HCT MSD 100 mg/25 mg Filmtabletten	NL/H/1606/003	1-28602	MERCK SHARP & DOHME GES.M.B.H.	AT
Losartan/HCT MSD 50 mg/12,5 mg Filmtabletten	NL/H/1606/001	1-28601	MERCK SHARP & DOHME GES.M.B.H.	AT
Losartan + Hidroclorotiazida ratiopharm, 100 mg +	PT/H/0240/001	5299334	RATIOPHARM LDA	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
12,5 mg, Comprimidos revestidos por película				
Losartan/HCT ratiopharm 100 mg/12,5 mg Filmtabletten	PT/H/0240/001	1-29685	TEVA B.V	AT
Losartan Hydrochlorotiazyd Krka, 100 mg + 12,5 mg, tabletki powlekane	NL/H/1930/001	18108	KRKA, D.D., NOVO MESTO	PL
Lavestra HCT 100 mg/12,5 mg comprimidos recubiertos con película	NL/H/1930/001	73846	KRKA, D.D., NOVO MESTO	ES
Loortan® Plus 50 mg/12,5 mg, Filmtabletten	not available	BE358102	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 50 mg/12,5 mg, Filmtabletten	not available	BE195246	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, Filmtabletten	not available	BE358127	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, Filmtabletten	not available	BE358111	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus Forte 100 mg/25 mg, Filmtabletten	not available	BE229503	MSD BELGIUM BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Loortan® Plus Forte 100 mg/25 mg, Filmtabletten	not available	BE358136	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, filmomhulde tabletten	not available	BE358127	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 50 mg/12,5 mg, comprimés pelliculés	not available	BE358102	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 50 mg/12,5 mg, filmomhulde tabletten	not available	BE358102	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 50 mg/12,5 mg, filmomhulde tabletten	not available	BE195246	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, comprimés pelliculés	not available	BE358111	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, filmomhulde tabletten	not available	BE358111	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 100 mg/12,5 mg, comprimés pelliculés	not available	BE358127	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus 50 mg/12,5 mg, comprimés pelliculés	not available	BE195246	MSD BELGIUM BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Loortan® Plus Forte 100 mg/25 mg, comprimés pelliculés	not available	BE229503	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus Forte 100 mg/25 mg, comprimés pelliculés	not available	BE358136	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus Forte 100 mg/25 mg, filmomhulde tabletten	not available	BE229503	MSD BELGIUM BVBA/SPRL	BE
Loortan® Plus Forte 100 mg/25 mg, filmomhulde tabletten	not available	BE358136	MSD BELGIUM BVBA/SPRL	BE
Neo-Lotan Plus 50 mg + 12.5 mg compresse rivestite con film	not available	032993014	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 50 mg + 12,5 mg compresse rivestite con film	not available	032993053	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993038	NEOPHARMED GENTILI S.R.L.	IT
Neo-Lotan Plus 100 mg + 25 mg compresse rivestite con film	not available	032993040	NEOPHARMED GENTILI S.R.L.	IT
Losartan potassium/hydrochlorothiazide 50 mg/12.5 mg film-coated tablets	not available	PL 21880/0143	MEDREICH PLC	UK

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 potassium/hydrochlorothiazide 100 mg/25 mg film-coated tablets	not available	PL 21880/0144	MEDREICH PLC	UK