30 November 2017
EMA/812634/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: Etoricoxib

Procedure no.: PSUSA/00001334/201703

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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
ACOXXEL 120 mg comprimidos recubiertos con película	UK/H/3190/004	71.590	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
ACOXXEL 30 mg comprimidos recubiertos con película	UK/H/3190/001	71.587	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
ACOXXEL 60 mg comprimidos recubiertos con película	UK/H/3190/002	71.588	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
ACOXXEL 90 mg comprimidos recubiertos con película	UK/H/3190/003	71.589	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
ACOXXEL® 30 mg film-coated tablets	UK/H/3190/001	PL 0025/0524	MERCK SHARP & DOHME LTD.	UK
ACOXXEL® 60 mg film-coated tablets	UK/H/3190/002	PL 0025/0525	MERCK SHARP & DOHME LTD.	UK
ACOXXEL® 90 mg film-coated tablets	UK/H/3190/003	PL 0025/0526	MERCK SHARP & DOHME LTD.	UK
ACOXXEL®120 mg film-coated tablets	UK/H/3190/004	PL 0025/0527	MERCK SHARP & DOHME LTD.	UK
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821293	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821317	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821329	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821331	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821343	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821356	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
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821368	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821370	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821382	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821394	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/001-004	035821406	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821418	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821420	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821305	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 30 mg compresse rivestite con film	UK/H/0533/001	035821444	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 30 mg compresse rivestite con film	UK/H/0533/001	035821432	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821014	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821026	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821038	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821040	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
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821053	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821077	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821089	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821091	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821103	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821115	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821127	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821139	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821141	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821065	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821154	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821166	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821178	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821180	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
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821192	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821216	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821228	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821230	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821242	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821255	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821267	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821279	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821281	NEOPHARMED GENTILI S.R.L.	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821204	NEOPHARMED GENTILI S.R.L.	IT
ARCOXIA 120 mg apvalkotās tabletes	UK/H/0532/003	03-0005	MERCK SHARP & DOHME LATVIJA SIA	LV
ARCOXIA 120 mg comprimate filmate	not available	854/2008/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 120 mg comprimate filmate	not available	854/2008/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 120 mg comprimate filmate	not available	854/2008/01	MERCK SHARP & DOHME ROMANIA SRL	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
ARCOXIA 120 mg comprimate filmate	not available	854/2008/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 120 mg comprimidos recubiertos con película	UK/H/0532/003	64.930	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4107280	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4242780	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4107181	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106985	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4107082	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106084	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106886	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106381	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106787	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4107389	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106480	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106282	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
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4242889	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106183	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106589	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	UK/H/0532/003	4106688	MERCK SHARP & DOHME, LDA.	PT
ARCOXIA 120 mg film-coated tablets	UK/H/0532/003	19446	MERCK SHARP & DOHME BV	CY
Arcoxia 120 mg Film-coated tablets	UK/H/0532/003	PA 1997/1/4	MERCK SHARP & DOHME BV	IE
ARCOXIA 120 mg film-coated tablets	UK/H/0532/003	MA224/00303	MERCK SHARP & DOHME BV	MT
Arcoxia 120 mg filmdragerade tabletter	UK/H/0532/003	17275	MERCK SHARP & DOHME BV	FI
Arcoxia 120 mg filmdragerade tabletter	UK/H/0532/003	18367	MERCK SHARP & DOHME BV	SE
ARCOXIA 120 mg filmdrasjerte tabletter	UK/H/0532/003	02-1052	MERCK SHARP & DOHME BV	NO
ARCOXIA 120 mg filmom obalené tablety	UK/H/0532/003	29/0005/03-S	MERCK SHARP & DOHME BV	SK
ARCOXIA 120 mg filmomhulde tabletten	UK/H/0532/003	RVG 27707	MERCK SHARP & DOHME BV	NL
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/042	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
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/043	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/044	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/045	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/046	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/047	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/048	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/049	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/050	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/051	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
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/052	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/053	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/054	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/055	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/056	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/057	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 120 mg filmsko obložene tablete	UK/H/0532/003	H/02/00207/058	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Arcoxia 120 mg Filmtabletten	UK/H/0532/003	1-24676	MERCK SHARP & DOHME BV	AT
ARCOXIA 120 mg Filmtabletten	UK/H/0532/003	59863.02.00	MERCK SHARP & DOHME BV	DE
ARCOXIA 120 mg filmuhúðaðar töflur	UK/H/0532/003	IS/1/02/025/03	MERCK SHARP & DOHME BV	IS
ARCOXIA 120 mg plėvele dengtos tabletės	UK/H/0532/003	LT/1/08/1419/011	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 120 mg plėvele dengtos tabletės	UK/H/0532/003	LT/1/08/1419/009	MERCK SHARP & DOHME, UAB	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
ARCOXIA 120 mg plėvele dengtos tabletės	UK/H/0532/003	LT/1/08/1419/010	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 120 mg plėvele dengtos tabletės	UK/H/0532/003	LT/1/08/1419/008	MERCK SHARP & DOHME, UAB	LT
Arcoxia 120 mg tabletti, kalvopäällysteinen	UK/H/0532/003	17275	MERCK SHARP & DOHME BV	FI
ARCOXIA 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/003	19446	MERCK SHARP & DOHME BV	CY
ARCOXIA 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/003	71849/06-11-2013	VIANEX S.A.	GR
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820378	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820380	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820327	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820341	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820291	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820339	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820303	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820392	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820354	MSD 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
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820366	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820315	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820416	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820428	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	UK/H/0532/003	035820404	MSD ITALIA S.R.L.	IT
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/58	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/12	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/16	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/01	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/10	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/14	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/11	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/04	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/03	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
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/13	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/15	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/18	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/02	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/17	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/20	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/19	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	UK/H/0532/003	OGYI-T-8825/21	MSD PHARMA HUNGARY KFT.	HU
ARCOXIA 30 mg apvalkotās tabletes	UK/H/0532/004	07-0353	MERCK SHARP & DOHME LATVIJA SIA	LV
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 30 mg comprimidos recubiertos con película	UK/H/0532/004	69.376	MERCK SHARP & DOHME DE ESPAÑA, 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
Arcoxia 30 mg comprimidos revestidos por película	UK/H/0532/004	5118625	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 30 mg comprimidos revestidos por película	UK/H/0532/004	5063037	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 30 mg comprimidos revestidos por película	UK/H/0532/004	5715032	MERCK SHARP & DOHME, LDA.	PT
ARCOXIA 30 mg film-coated tablets	UK/H/0532/004	20338	MERCK SHARP & DOHME BV	CY
Arcoxia 30 mg Film-coated tablets	UK/H/0532/004	PA 1997/1/1	MERCK SHARP & DOHME BV	IE
Arcoxia 30 mg filmdragerade tabletter	UK/H/0532/004	22403	MERCK SHARP & DOHME BV	FI
ARCOXIA 30 mg filmdragerade tabletter	UK/H/0532/004	24178	MERCK SHARP & DOHME BV	SE
ARCOXIA 30 mg filmdrasjerte tabletter	UK/H/0532/004	06-4274	MERCK SHARP & DOHME BV	NO
ARCOXIA 30 mg filmomhulde tabletten	UK/H/0532/004	RVG 34279	MERCK SHARP & DOHME BV	NL
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/002	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/003	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/004	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
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/005	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/006	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/007	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 30 mg filmsko obložene tablete	UK/H/0532/004	H/02/00207/059	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Arcoxia 30 mg Filmtabletten	UK/H/0532/004	1-27325	MERCK SHARP & DOHME BV	AT
ARCOXIA 30 mg Filmtabletten	UK/H/0532/004	66968.00.00	MERCK SHARP & DOHME BV	DE
ARCOXIA 30 mg filmuhúðaðar töflur	UK/H/0532/004	IS/1/06/050/01	MERCK SHARP & DOHME BV	IS
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/013	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/001	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/015	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/014	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/012	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/016	MERCK SHARP & DOHME, UAB	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
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/017	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 30 mg plėvele dengtos tabletės	UK/H/0532/004	LT/1/08/1419/018	MERCK SHARP & DOHME, UAB	LT
Arcoxia 30 mg tabletti, kalvopäällysteinen	UK/H/0532/004	22403	MERCK SHARP & DOHME BV	FI
ARCOXIA 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/004	20338	MERCK SHARP & DOHME BV	CY
ARCOXIA 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/004	71850/06-11-2013	VIANEX S.A.	GR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 387 977 7 0	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 387 978 3 1	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 387 980 8 1	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 387 976 0 2	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 573 530 9 4	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 387 974 8 0	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, comprimé pelliculé	UK/H/0532/004	34009 300 043 0 2	MERCK SHARP & DOHME BV	FR
ARCOXIA 30 mg, filmom obalené tablety	UK/H/0532/004	29/0125/08-S	MERCK SHARP & DOHME BV	SK
ARCOXIA 30 mg compresse rivestite con film	UK/H/0532/004	035820442	MSD 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
ARCOXIA 30 mg compresse rivestite con film	UK/H/0532/004	035820430	MSD ITALIA S.R.L.	IT
ARCOXIA 30 mg compresse rivestite con film	UK/H/0532/004	035820568	MSD ITALIA S.R.L.	IT
ARCOXIA 30 mg film-coated tablets	UK/H/0532/004	MA224/00304	MERCK SHARP & DOHME BV	MT
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/55	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/25	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/26	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/23	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/24	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/05	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/22	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	UK/H/0532/004	OGYI-T-8825/59	MSD PHARMA HUNGARY KFT.	HU
ARCOXIA 60 mg apvalkotās tabletes	UK/H/0532/001	03-0003	MERCK SHARP & DOHME LATVIJA SIA	LV
ARCOXIA 60 mg comprimate filmate	not available	852/2008/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/01	MERCK SHARP & DOHME ROMANIA SRL	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
ARCOXIA 60 mg comprimate filmate	not available	852/2008/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 60 mg comprimidos recubiertos con película	UK/H/0532/001	64.928	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108189	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107587	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108783	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108585	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108288	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108387	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4242483	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107686	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107488	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107983	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4242384	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
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108080	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108684	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4108486	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107884	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	UK/H/0532/001	4107785	MERCK SHARP & DOHME, LDA.	PT
ARCOXIA 60 mg film-coated tablets	UK/H/0532/001	19444	MERCK SHARP & DOHME BV	CY
Arcoxia 60 mg Film-coated tablets	UK/H/0532/001	PA 1997/1/2	MERCK SHARP & DOHME BV	IE
Arcoxia 60 mg filmdragerade tabletter	UK/H/0532/001	17273	MERCK SHARP & DOHME BV	FI
Arcoxia 60 mg filmdragerade tabletter	UK/H/0532/001	18365	MERCK SHARP & DOHME BV	SE
ARCOXIA 60 mg filmdrasjerte tabletter	UK/H/0532/001	02-1050	MERCK SHARP & DOHME BV	NO
ARCOXIA 60 mg filmom obalené tablety	UK/H/0532/001	29/0003/03-S	MERCK SHARP & DOHME BV	SK
ARCOXIA 60 mg filmomhulde tabletten	UK/H/0532/001	RVG 27705	MERCK SHARP & DOHME BV	NL
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/010	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/009	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
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/008	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/011	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/012	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/013	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/014	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/015	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/016	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/017	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/018	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
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/019	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/020	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/021	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/022	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/023	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/024	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 60 mg filmsko obložene tablete	UK/H/0532/001	H/02/00207/060	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Arcoxia 60 mg Filmtabletten	UK/H/0532/001	1-24674	MERCK SHARP & DOHME BV	AT
ARCOXIA 60 mg Filmtabletten	UK/H/0532/001	59863.00.00	MERCK SHARP & DOHME BV	DE
ARCOXIA 60 mg filmuhúðaðar töflur	UK/H/0532/001	IS/1/02/025/01	MERCK SHARP & DOHME BV	IS
ARCOXIA 60 mg plėvele dengtos tabletės	UK/H/0532/001	LT/1/08/1419/002	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 60 mg plėvele dengtos tabletės	UK/H/0532/001	LT/1/08/1419/003	MERCK SHARP & DOHME, UAB	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
ARCOXIA 60 mg plėvele dengtos tabletės	UK/H/0532/001	LT/1/08/1419/004	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 60 mg plėvele dengtos tabletės	UK/H/0532/001	LT/1/08/1419/019	MERCK SHARP & DOHME, UAB	LT
Arcoxia 60 mg tabletti, kalvopäällysteinen	UK/H/0532/001	17273	MERCK SHARP & DOHME BV	FI
ARCOXIA 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/001	19444	MERCK SHARP & DOHME BV	CY
ARCOXIA 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/001	71847/06-11-2013	VIANEX S.A.	GR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 512 0 5	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 503 1 4	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 947 0 0	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 949 3 9	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 511 4 4	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 501 9 2	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 502 5 3	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 948 7 8	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 513 7 3	MERCK SHARP & DOHME BV	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
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 573 510 8 3	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 950 1 1	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 925 7 7	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 928 6 7	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 923 4 8	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 926 3 8	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 387 924 0 9	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg, comprimé pelliculé	UK/H/0532/001	34009 275 698 9 0	MERCK SHARP & DOHME BV	FR
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820063	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820051	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820048	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820024	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820012	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820036	MSD 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
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820075	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820099	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820101	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820113	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820125	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820137	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820149	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820087	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	UK/H/0532/001	035820570	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg film-coated tablets	UK/H/0532/001	MA224/00302	MERCK SHARP & DOHME BV	MT
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/56	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/29	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/37	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/06	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
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/38	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/34	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/32	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/27	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/35	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/28	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/31	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/30	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/36	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/39	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/33	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/07	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/40	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	UK/H/0532/001	OGYI-T-8825/60	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
ARCOXIA 90 mg apvalkotās tabletes	UK/H/0532/002	03-0004	MERCK SHARP & DOHME LATVIJA SIA	LV
ARCOXIA 90 mg comprimate filmate	not available	853/2008/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ARCOXIA 90 mg comprimidos recubiertos con película	UK/H/0532/002	64.929	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109385	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109484	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4110086	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109187	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109781	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109088	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4242681	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4108882	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
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109880	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109583	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4242582	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4108981	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4110185	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109989	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109286	MERCK SHARP & DOHME, LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	UK/H/0532/002	4109682	MERCK SHARP & DOHME, LDA.	PT
ARCOXIA 90 mg film-coated tablets	UK/H/0532/002	19445	MERCK SHARP & DOHME BV	CY
Arcoxia 90 mg Film-coated tablets	UK/H/0532/002	PA 1997/1/3	MERCK SHARP & DOHME BV	IE
ARCOXIA 90 mg film-coated tablets	UK/H/0532/002	MA224/00301	MERCK SHARP & DOHME BV	MT
Arcoxia 90 mg filmdragerade tabletter	UK/H/0532/002	17274	MERCK SHARP & DOHME BV	FI
Arcoxia 90 mg filmdragerade tabletter	UK/H/0532/002	18366	MERCK SHARP & DOHME BV	SE
ARCOXIA 90 mg filmdrasjerte tabletter	UK/H/0532/002	02-1051	MERCK SHARP & DOHME BV	NO
ARCOXIA 90 mg filmom obalené tablety	UK/H/0532/002	29/0004/03-S	MERCK SHARP & DOHME BV	SK

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
ARCOXIA 90 mg filmomhulde tabletten	UK/H/0532/002	RVG 27706	MERCK SHARP & DOHME BV	NL
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/025	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/026	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/027	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/028	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/029	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/030	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/031	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/032	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/033	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
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/034	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/035	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/036	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/037	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/038	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/039	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/040	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
ARCOXIA 90 mg filmsko obložene tablete	UK/H/0532/002	H/02/00207/041	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Arcoxia 90 mg Filmtabletten	UK/H/0532/002	1-24675	MERCK SHARP & DOHME BV	AT
ARCOXIA 90 mg Filmtabletten	UK/H/0532/002	59863.01.00	MERCK SHARP & DOHME BV	DE
ARCOXIA 90 mg filmuhúðaðar töflur	UK/H/0532/002	IS/1/02/025/02	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
ARCOXIA 90 mg plėvele dengtos tabletės	UK/H/0532/002	LT/1/08/1419/005	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 90 mg plėvele dengtos tabletės	UK/H/0532/002	LT/1/08/1419/007	MERCK SHARP & DOHME, UAB	LT
ARCOXIA 90 mg plėvele dengtos tabletės	UK/H/0532/002	LT/1/08/1419/006	MERCK SHARP & DOHME, UAB	LT
Arcoxia 90 mg tabletti, kalvopäällysteinen	UK/H/0532/002	17274	MERCK SHARP & DOHME BV	FI
ARCOXIA 90 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/002	19445	MERCK SHARP & DOHME BV	CY
ARCOXIA 90 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0532/002	71848/06-11-2013	VIANEX S.A.	GR
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820164	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820152	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820176	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820188	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820253	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820289	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820238	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820190	MSD 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
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820226	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820277	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820214	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820265	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820240	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	UK/H/0532/002	035820202	MSD ITALIA S.R.L.	IT
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/57	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/46	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/53	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/42	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/51	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/54	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/52	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/50	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
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/49	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/47	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/44	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/41	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/43	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/48	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/08	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/09	MSD PHARMA HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	UK/H/0532/002	OGYI-T-8825/45	MSD PHARMA HUNGARY KFT.	HU
Arcoxia, 120 mg õhukese polümeerikattega tabletid	UK/H/0532/003	393002	MERCK SHARP & DOHME OU	EE
ARCOXIA, 120 mg, tabletki powlekane	UK/H/0532/003	11468	MSD POLSKA SP. Z O.O.	PL
Arcoxia, 30 mg õhukese polümeerikattega tabletid	UK/H/0532/004	572308	MERCK SHARP & DOHME OU	EE
ARCOXIA, 30 mg, tabletki powlekane	UK/H/0532/004	22580	MSD POLSKA SP. Z O.O.	PL
Arcoxia, 60 mg õhukese polümeerikattega tabletid	UK/H/0532/001	393202	MERCK SHARP & DOHME OU	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
ARCOXIA, 60 mg, tabletki powlekane	UK/H/0532/001	11466	MSD POLSKA SP. Z O.O.	PL
Arcoxia, 90 mg õhukese polümeerikattega tabletid	UK/H/0532/002	393102	MERCK SHARP & DOHME OU	EE
ARCOXIA, 90 mg, tabletki powlekane	UK/H/0532/002	11467	MSD POLSKA SP. Z O.O.	PL
Arcoxia, filmovertrukne tabletter	UK/H/0532/001	33573	MERCK SHARP & DOHME BV	DK
Arcoxia, filmovertrukne tabletter	UK/H/0532/002	33574	MERCK SHARP & DOHME BV	DK
Arcoxia, filmovertrukne tabletter	UK/H/0532/003	33575	MERCK SHARP & DOHME BV	DK
Arcoxia, filmovertrukne tabletter	UK/H/0532/004	39698	MERCK SHARP & DOHME BV	DK
ARCOXIA® 120 mg comprimés pelliculés	UK/H/0532/003	BE239556	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg comprimés pelliculés	UK/H/0532/003	BE387405	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg comprimés pelliculés	UK/H/0532/003	2008019636	MERCK SHARP & DOHME BV	LU
ARCOXIA® 120 mg film-coated tablets	UK/H/0532/003	PL 04900/0004	MERCK SHARP & DOHME BV	UK
ARCOXIA® 120 mg filmom obložene tablete	not available	UP/I-530-09/12-02/63	MERCK SHARP & DOHME D.O.O.	HR
ARCOXIA® 120 mg filmomhulde tabletten	UK/H/0532/003	BE387405	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg filmomhulde tabletten	UK/H/0532/003	BE239556	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg Filmtabletten	UK/H/0532/003	BE387405	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg Filmtabletten	UK/H/0532/003	BE239556	MERCK SHARP & DOHME BV	BE
ARCOXIA® 120 mg, potahované tablety	UK/H/0532/003	29/078/03-C	MERCK SHARP & DOHME BV	CZ

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
ARCOXIA® 30 mg comprimés pelliculés	UK/H/0532/004	BE303886	MERCK SHARP & DOHME BV	BE
ARCOXIA® 30 mg comprimés pelliculés	UK/H/0532/004	2008020011	MERCK SHARP & DOHME BV	LU
ARCOXIA® 30 mg film-coated tablets	UK/H/0532/004	PL 04900/0001	MERCK SHARP & DOHME BV	UK
ARCOXIA® 30 mg filmom obložene tablete	not available	UP/I-530-09/12-02/60	MERCK SHARP & DOHME D.O.O.	HR
ARCOXIA® 30 mg filmomhulde tabletten	UK/H/0532/004	BE303886	MERCK SHARP & DOHME BV	BE
ARCOXIA® 30 mg Filmtabletten	UK/H/0532/004	BE303886	MERCK SHARP & DOHME BV	BE
ARCOXIA® 30 mg, potahované tablety	UK/H/0532/004	29/617/08-C	MERCK SHARP & DOHME BV	CZ
ARCOXIA® 60 mg comprimés pelliculés	UK/H/0532/001	BE387387	MERCK SHARP & DOHME BV	BE
ARCOXIA® 60 mg comprimés pelliculés	UK/H/0532/001	BE239531	MERCK SHARP & DOHME BV	BE
ARCOXIA® 60 mg comprimés pelliculés	UK/H/0532/001	2008019634	MERCK SHARP & DOHME BV	LU
ARCOXIA® 60 mg film-coated tablets	UK/H/0532/001	PL 04900/0002	MERCK SHARP & DOHME BV	UK
ARCOXIA® 60 mg filmom obložene tablete	not available	UP/I-530-09/12-02/61	MERCK SHARP & DOHME D.O.O.	HR
ARCOXIA® 60 mg filmomhulde tabletten	UK/H/0532/001	BE239531	MERCK SHARP & DOHME BV	BE
ARCOXIA® 60 mg filmomhulde tabletten	UK/H/0532/001	BE387387	MERCK SHARP & DOHME BV	BE
ARCOXIA® 60 mg Filmtabletten	UK/H/0532/001	BE239531	MERCK SHARP & DOHME BV	BE
ARCOXIA® 60 mg Filmtabletten	UK/H/0532/001	BE387387	MERCK SHARP & DOHME BV	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
ARCOXIA® 60 mg, potahované tablety	UK/H/0532/001	29/076/03-C	MERCK SHARP & DOHME BV	CZ
ARCOXIA® 90 mg comprimés pelliculés	UK/H/0532/002	BE239547	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg comprimés pelliculés	UK/H/0532/002	BE387396	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg comprimés pelliculés	UK/H/0532/002	2008019635	MERCK SHARP & DOHME BV	LU
ARCOXIA® 90 mg film-coated tablets	UK/H/0532/002	PL 04900/0003	MERCK SHARP & DOHME BV	UK
ARCOXIA® 90 mg filmom obložene tablete	not available	UP/I-530-09/12-02/62	MERCK SHARP & DOHME D.O.O.	HR
ARCOXIA® 90 mg filmomhulde tabletten	UK/H/0532/002	BE239547	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg filmomhulde tabletten	UK/H/0532/002	BE387396	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg Filmtabletten	UK/H/0532/002	BE239547	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg Filmtabletten	UK/H/0532/002	BE387396	MERCK SHARP & DOHME BV	BE
ARCOXIA® 90 mg, potahované tablety	UK/H/0532/002	29/077/03-C	MERCK SHARP & DOHME BV	CZ
AUXIB® 120 mg film-coated tablets	UK/H/0533/004	PL 0025/0427	MERCK SHARP & DOHME LTD.	UK
AUXIB® 30 mg film-coated tablets	UK/H/0533/001	PL 0025/0479	MERCK SHARP & DOHME LTD.	UK
AUXIB® 60 mg film-coated tablets	UK/H/0533/002	PL 0025/0425	MERCK SHARP & DOHME LTD.	UK
AUXIB® 90 mg film-coated tablets	UK/H/0533/003	PL 0025/0426	MERCK SHARP & DOHME LTD.	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
Bericox 120 mg apvalkotās tabletes	HU/H/0435/004	16-0213	KRKA, D.D., NOVO MESTO	LV
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/033	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/034	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/035	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/036	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/037	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/038	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/039	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/040	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/041	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/042	KRKA, D.D., NOVO MESTO	LT
Bericox 120 mg plėvele dengtos tabletės	HU/H/0435/004	LT/1/16/3998/043	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg apvalkotās tabletes	HU/H/0435/001	16-0210	KRKA, D.D., NOVO MESTO	LV
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/001	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/002	KRKA, D.D., NOVO MESTO	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
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/003	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/004	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/005	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/006	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/007	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/008	KRKA, D.D., NOVO MESTO	LT
Bericox 30 mg plėvele dengtos tabletės	HU/H/0435/001	LT/1/16/3998/009	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg apvalkotās tabletes	HU/H/0435/002	16-0211	KRKA, D.D., NOVO MESTO	LV
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/010	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/011	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/012	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/013	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/014	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/015	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/016	KRKA, D.D., NOVO MESTO	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
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/017	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/018	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/019	KRKA, D.D., NOVO MESTO	LT
Bericox 60 mg plėvele dengtos tabletės	HU/H/0435/002	LT/1/16/3998/020	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg apvalkotās tabletes	HU/H/0435/003	16-0212	KRKA, D.D., NOVO MESTO	LV
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/021	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/022	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/023	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/024	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/025	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/026	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/027	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/028	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/029	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/030	KRKA, D.D., NOVO MESTO	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
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/031	KRKA, D.D., NOVO MESTO	LT
Bericox 90 mg plėvele dengtos tabletės	HU/H/0435/003	LT/1/16/3998/032	KRKA, D.D., NOVO MESTO	LT
Caretor 120 mg Filmtabletten	NL/H/3269/004	93089.00.00	SYNTHON BV	DE
Caretor 30 mg Filmtabletten	NL/H/3269/001	93086.00.00	SYNTHON BV	DE
Caretor 60 mg Filmtabletten	NL/H/3269/002	93087.00.00	SYNTHON BV	DE
Caretor 90 mg Filmtabletten	NL/H/3269/003	93088.00.00	SYNTHON BV	DE
Coxerit 120 mg filmuhúðaðar töflur	NL/H/3269/004	IS/1/17/004/04	WILLIAMS & HALLS EHF	IS
Coxerit 30 mg filmuhúðaðar töflur	NL/H/3269/001	IS/1/17/004/01	WILLIAMS & HALLS EHF	IS
Coxerit 60 mg filmuhúðaðar töflur	NL/H/3269/002	IS/1/17/004/02	WILLIAMS & HALLS EHF	IS
Coxerit 90 mg filmuhúðaðar töflur	NL/H/3269/003	IS/1/17/004/03	WILLIAMS & HALLS EHF	IS
Coxeta 120 mg filmom obložene tablete	DE/H/5031/004	HR-H-189772606	PLIVA HRVATSKA D.O.O.	HR
Coxeta 120 mg filmsko obložene tablete	DE/H/5031/004	H/16/02155/031	TEVA B.V	SI
Coxeta 30 mg filmom obložene tablete	DE/H/5031/001	HR-H-547635349	PLIVA HRVATSKA D.O.O.	HR
Coxeta 30 mg filmsko obložene tablete	DE/H/5031/001	H/16/02155/002	TEVA B.V	SI
Coxeta 30 mg filmsko obložene tablete	DE/H/5031/001	H/16/02155/009	TEVA B.V	SI
Coxeta 60 mg filmom obložene tablete	DE/H/5031/002	HR-H-072514188	PLIVA HRVATSKA D.O.O.	HR
Coxeta 60 mg filmsko obložene tablete	DE/H/5031/002	H/16/02155/019	TEVA B.V	SI
Coxeta 90 mg filmom obložene tablete	DE/H/5031/003	HR-H-328272058	PLIVA 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
Coxient 120 mg comprimate filmate	IS/H/0244/004	9285/2016/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 120 mg comprimate filmate	IS/H/0244/004	9285/2016/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 120 mg comprimate filmate	IS/H/0244/004	9285/2016/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 120 mg comprimate filmate	IS/H/0244/004	9285/2016/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 120 mg comprimate filmate	IS/H/0244/004	9285/2016/05	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
COXIENT 120 mg filmuhúðaðar töflur	IS/H/0244/004	IS/1/16/070/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 120 mg filmuhúðaðar töflur	IS/H/0244/004	IS/1/16/070/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 120 mg filmuhúðaðar töflur	IS/H/0244/004	IS/1/16/070/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 120 mg filmuhúðaðar töflur	IS/H/0244/004	IS/1/16/070/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 120 mg filmuhúðaðar töflur	IS/H/0244/004	IS/1/16/070/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
Coxient 30 mg comprimate filmate	IS/H/0244/001	9282/2016/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 30 mg comprimate filmate	IS/H/0244/001	9282/2016/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 30 mg comprimate filmate	IS/H/0244/001	9282/2016/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 30 mg comprimate filmate	IS/H/0244/001	9282/2016/05	ALVOGEN MALTA OPERATIONS (ROW) LTD	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
Coxient 30 mg comprimate filmate	IS/H/0244/001	9282/2016/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
COXIENT 30 mg filmuhúðaðar töflur	IS/H/0244/001	IS/1/16/070/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 30 mg filmuhúðaðar töflur	IS/H/0244/001	IS/1/16/070/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 30 mg filmuhúðaðar töflur	IS/H/0244/001	IS/1/16/070/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 30 mg filmuhúðaðar töflur	IS/H/0244/001	IS/1/16/070/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 30 mg filmuhúðaðar töflur	IS/H/0244/001	IS/1/16/070/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
Coxient 60 mg comprimate filmate	IS/H/0244/002	9283/2016/05	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 60 mg comprimate filmate	IS/H/0244/002	9283/2016/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 60 mg comprimate filmate	IS/H/0244/002	9283/2016/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 60 mg comprimate filmate	IS/H/0244/002	9283/2016/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 60 mg comprimate filmate	IS/H/0244/002	9283/2016/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
COXIENT 60 mg filmuhúðaðar töflur	IS/H/0244/002	IS/1/16/070/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 60 mg filmuhúðaðar töflur	IS/H/0244/002	IS/1/16/070/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 60 mg filmuhúðaðar töflur	IS/H/0244/002	IS/1/16/070/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	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**
COXIENT 60 mg filmuhúðaðar töflur	IS/H/0244/002	IS/1/16/070/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 60 mg filmuhúðaðar töflur	IS/H/0244/002	IS/1/16/070/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
Coxient 90 mg comprimate filmate	IS/H/0244/003	9284/2016/05	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 90 mg comprimate filmate	IS/H/0244/003	9284/2016/04	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 90 mg comprimate filmate	IS/H/0244/003	9284/2016/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 90 mg comprimate filmate	IS/H/0244/003	9284/2016/02	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
Coxient 90 mg comprimate filmate	IS/H/0244/003	9284/2016/01	ALVOGEN MALTA OPERATIONS (ROW) LTD	RO
COXIENT 90 mg filmuhúðaðar töflur	IS/H/0244/003	IS/1/16/070/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 90 mg filmuhúðaðar töflur	IS/H/0244/003	IS/1/16/070/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 90 mg filmuhúðaðar töflur	IS/H/0244/003	IS/1/16/070/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 90 mg filmuhúðaðar töflur	IS/H/0244/003	IS/1/16/070/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
COXIENT 90 mg filmuhúðaðar töflur	IS/H/0244/003	IS/1/16/070/03	ALVOGEN MALTA OPERATIONS (ROW) LTD	IS
Coxiloc 120 mg tabletti, kalvopäällysteinen	NL/H/3269/004	32553	AVANSOR PHARMA OY	FI
Coxiloc 30 mg tabletti, kalvopäällysteinen	NL/H/3269/001	32550	AVANSOR PHARMA OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Coxiloc 60 mg tabletti, kalvopäällysteinen	NL/H/3269/002	32551	AVANSOR PHARMA OY	FI
Coxiloc 90 mg tabletti, kalvopäällysteinen	NL/H/3269/003	32552	AVANSOR PHARMA OY	FI
COXITOR	DE/H/3908/001	869415	SANDOZ PHARMACEUTICALS D.D.	EE
COXITOR	DE/H/3908/002	869515	SANDOZ PHARMACEUTICALS D.D.	EE
COXITOR	DE/H/3908/003	869215	SANDOZ PHARMACEUTICALS D.D.	EE
COXITOR	DE/H/3908/004	869315	SANDOZ PHARMACEUTICALS D.D.	EE
Coxitor 120 mg apvalkotās tabletes	DE/H/3908/004	15-0104	SANDOZ PHARMACEUTICALS D.D.	LV
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/097	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/098	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/099	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/100	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/101	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/102	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/103	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/104	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/105	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/106	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/107	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/109	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/110	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/111	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/112	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/113	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/114	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/115	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/116	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/119	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/118	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/120	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/121	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/122	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/123	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/124	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/125	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/126	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/127	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/128	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/117	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 120 mg plėvelė dengtos tabletės	DE/H/3908/004	LT/1/15/3721/108	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/001	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/002	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/003	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/004	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/005	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/006	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/007	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/008	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/012	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/010	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/009	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/013	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/014	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/011	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/015	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/016	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/017	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/018	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/019	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/020	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/021	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/022	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/023	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/024	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/025	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/026	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/027	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/028	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/029	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/031	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/030	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 30 mg plėvelė dengtos tabletės	DE/H/3908/001	LT/1/15/3721/032	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg apvalkotās tabletes	DE/H/3908/002	15-0102	SANDOZ PHARMACEUTICALS D.D.	LV
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/033	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/034	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/035	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/036	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/037	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/038	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/039	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/040	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/041	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/042	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/043	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/044	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/045	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/046	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/047	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/048	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/049	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/050	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/051	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/052	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/053	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/054	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/055	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/056	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/058	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/059	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/057	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/060	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/061	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/062	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/063	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 60 mg plėvelė dengtos tabletės	DE/H/3908/002	LT/1/15/3721/064	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg apvalkotās tabletes	DE/H/3908/003	15-0103	SANDOZ PHARMACEUTICALS D.D.	LV
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/065	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/072	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/073	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/074	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/075	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/076	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/077	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/078	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/079	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/080	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/081	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/082	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/083	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/084	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/085	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/086	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/087	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/088	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/089	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/090	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/091	SANDOZ PHARMACEUTICALS D.D.	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
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/092	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/093	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/094	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/095	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/096	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/066	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/067	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/068	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/069	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/070	SANDOZ PHARMACEUTICALS D.D.	LT
Coxitor 90 mg plėvelė dengtos tabletės	DE/H/3908/003	LT/1/15/3721/071	SANDOZ PHARMACEUTICALS D.D.	LT
Coxolin, 120 mg filmdragerade tabletter	NL/H/3271/004	55246	SYNTHON BV	SE
Coxolin, 30 mg filmdragerade tabletter	NL/H/3271/001	55243	SYNTHON BV	SE
Coxolin, 60 mg filmdragerade tabletter	NL/H/3271/002	55244	SYNTHON BV	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
Coxolin, 90 mg filmdragerade tabletter	NL/H/3271/003	55245	SYNTHON BV	SE
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/04	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/03	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/05	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/02	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/01	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/04	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/03	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/05	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/02	ZENTIVA, K.S.	RO
Doloxib 120 mg comprimate filmate	EE/H/0211/004	8956/2016/01	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/02	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/01	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/03	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/02	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/01	ZENTIVA, K.S.	RO
Doloxib 30 mg comprimate filmate	EE/H/0211/001	8953/2016/03	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/04	ZENTIVA, K.S.	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
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/02	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/03	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/05	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/06	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/01	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/04	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/02	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/03	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/05	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/06	ZENTIVA, K.S.	RO
Doloxib 60 mg comprimate filmate	EE/H/0211/002	8954/2016/01	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/04	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/05	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/02	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/03	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/06	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/01	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/04	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/05	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/02	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/03	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/06	ZENTIVA, K.S.	RO
Doloxib 90 mg comprimate filmate	EE/H/0211/003	8955/2016/01	ZENTIVA, K.S.	RO
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL

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
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 120 mg, tabletki powlekane	EE/H/0211/004	23384	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL
Doloxib, 30 mg, tabletki powlekane	EE/H/0211/001	23381	ZENTIVA, K.S.	PL

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
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 60 mg, tabletki powlekane	EE/H/0211/002	23382	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL

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
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Doloxib, 90 mg, tabletki powlekane	EE/H/0211/003	23383	ZENTIVA, K.S.	PL
Ecoxyton 120 mg tabletten	NL/H/3269/004	RVG 116021	SYNTHON BV	NL
Ecoxyton 30 mg tabletten	NL/H/3269/001	RVG 116018	SYNTHON BV	NL
Ecoxyton 60 mg tabletten	NL/H/3269/002	RVG 116019	SYNTHON BV	NL
Ecoxyton 90 mg tabletten	NL/H/3269/003	RVG 116020	SYNTHON BV	NL
Etori - 1 A Pharma 120 mg Filmtabletten	DE/H/3908/004	90438.00.00	1 A PHARMA GMBH	DE
Etori - 1 A Pharma 30 mg Filmtabletten	DE/H/3908/001	90435.00.00	1 A 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
Etori - 1 A Pharma 60 mg Filmtabletten	DE/H/3908/002	90436.00.00	1 A PHARMA GMBH	DE
Etori - 1 A Pharma 90 mg Filmtabletten	DE/H/3908/003	90437.00.00	1 A PHARMA GMBH	DE
Etoriax 120 mg filmtabletta	HU/H/0443/004	OGYI-T-23100/04	KRKA, D.D., NOVO MESTO	HU
Etoriax 30 mg filmtabletta	HU/H/0443/001	OGYI-T-23100/01	KRKA, D.D., NOVO MESTO	HU
Etoriax 60 mg filmtabletta	HU/H/0443/002	OGYI-T-23100/02	KRKA, D.D., NOVO MESTO	HU
Etoriax 90 mg filmtabletta	HU/H/0443/003	OGYI-T-23100/03	KRKA, D.D., NOVO MESTO	HU
Etoriax® 120 mg Filmtabletten	HU/H/0435/004	96358.00.00	TAD PHARMA GMBH	DE
Etoriax® 30 mg Filmtabletten	HU/H/0435/001	96355.00.00	TAD PHARMA GMBH	DE
Etoriax® 60 mg Filmtabletten	HU/H/0435/002	96356.00.00	TAD PHARMA GMBH	DE
Etoriax® 90 mg Filmtabletten	HU/H/0435/003	96357.00.00	TAD PHARMA GMBH	DE
Etorican® 120 mg Filmtabletten	DE/H/4691/004	97259.00.00	HORMOSAN PHARMA GMBH	DE
Etorican® 30 mg Filmtabletten	DE/H/4691/001	97256.00.00	HORMOSAN PHARMA GMBH	DE
Etorican® 60 mg Filmtabletten	DE/H/4691/002	97257.00.00	HORMOSAN PHARMA GMBH	DE
Etorican® 90 mg Filmtabletten	DE/H/4691/003	97258.00.00	HORMOSAN PHARMA GMBH	DE
Etoricox-AbZ 120 mg Filmtabletten	DE/H/5031/004	92037.00.00	ABZ-PHARMA GMBH	DE
Etoricox-AbZ 30 mg Filmtabletten	DE/H/5031/001	92034.00.00	ABZ-PHARMA GMBH	DE
Etoricox-AbZ 60 mg Filmtabletten	DE/H/5031/002	92035.00.00	ABZ-PHARMA GMBH	DE
Etoricox-AbZ 90 mg Filmtabletten	DE/H/5031/003	92036.00.00	ABZ-PHARMA GMBH	DE
Etoricox-HEXAL 120 mg Filmtabletten	DE/H/3909/004	90486.00.00	HEXAL AG	DE
Etoricox-HEXAL 30 mg Filmtabletten	DE/H/3909/001	90483.00.00	HEXAL AG	DE
Etoricox-HEXAL 60 mg Filmtabletten	DE/H/3909/002	90484.00.00	HEXAL AG	DE
Etoricox-HEXAL 90 mg Filmtabletten	DE/H/3909/003	90485.00.00	HEXAL AG	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
Etoricoxib "Krka", filmovertrukne tabletter	HU/H/0435/001	56836	KRKA, D.D., NOVO MESTO	DK
Etoricoxib "Krka", filmovertrukne tabletter	HU/H/0435/002	56837	KRKA, D.D., NOVO MESTO	DK
Etoricoxib "Krka", filmovertrukne tabletter	HU/H/0435/003	56838	KRKA, D.D., NOVO MESTO	DK
Etoricoxib "Krka", filmovertrukne tabletter	HU/H/0435/004	56839	KRKA, D.D., NOVO MESTO	DK
Etoricoxib 120 mg Film-coated Tablets	ES/H/0264/004	PA 0688/032/004	CHANELLE MEDICAL	IE
Etoricoxib 120 mg Film-coated Tablets	NL/H/3151/004	PA0405/140/004	GENERICS [UK] LIMITED	IE
Etoricoxib 120 mg Film-coated tablets	DE/H/3908/004	PA0711/232/004	ROWEX LTD	IE
Etoricoxib 120 mg Film-coated Tablets	ES/H/0264/004	PA 0688/032/004	CHANELLE MEDICAL	IE
Etoricoxib 120 mg Film-coated Tablets	ES/H/0264/004	MA1026/00104	CHANELLE MEDICAL	MT
Etoricoxib 120 mg Film-coated Tablets	ES/H/0264/004	MA1026/00104	CHANELLE MEDICAL	MT
Etoricoxib 120 mg film-coated tablets	NL/H/3461/004	PL 42357/0170	AMNEAL PHARMA EUROPE LIMITED	UK
Etoricoxib 120 mg film-coated tablets	DE/H/3469/004	PL 31596/0004	ZYDUS FRANCE	UK
Etoricoxib 120 mg film-coated tablets	HU/H/0435/004	PL 01656/0217	KRKA, D.D., NOVO MESTO	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	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
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	PT/H/1603/004	PL 16363/0488	MILPHARM LIMITED	UK
Etoricoxib 120 mg Film-coated Tablets	DE/H/5092/004	PL 36687/0213	TORRENT PHARMA LTD.	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg film-coated tablets	EE/H/0211/004	PL 17780/0727	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 120 mg Film-coated Tablets	NL/H/3576/004	PL 25258/0202	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Etoricoxib 120 mg Film-coated Tablets	DE/H/5031/004	PL 00289/1937	TEVA UK LIMITED	UK
Etoricoxib 120mg Film-coated Tablets	SE/H/1574/004	PL 30306/0771	ACTAVIS GROUP PTC EHF.	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
Etoricoxib 30 mg Film-coated Tablets	ES/H/0264/001	PA 0688/032/001	CHANELLE MEDICAL	IE
Etoricoxib 30 mg Film-coated Tablets	NL/H/3151/001	PA0405/104/001	GENERICS [UK] LIMITED	IE
Etoricoxib 30 mg Film-coated tablets	DE/H/3908/001	PA0711/232/001	ROWEX LTD	IE
Etoricoxib 30 mg Film-coated Tablets	ES/H/0264/001	PA 0688/032/001	CHANELLE MEDICAL	IE
Etoricoxib 30 mg Film-coated Tablets	ES/H/0264/001	MA1026/00101	CHANELLE MEDICAL	MT
Etoricoxib 30 mg Film-coated Tablets	ES/H/0264/001	MA1026/00101	CHANELLE MEDICAL	MT
Etoricoxib 30 mg film-coated tablets	NL/H/3461/001	PL 42357/0167	AMNEAL PHARMA EUROPE LIMITED	UK
Etoricoxib 30 mg film-coated tablets	DE/H/3469/001	PL 31596/0001	ZYDUS FRANCE	UK
Etoricoxib 30 mg film-coated tablets	HU/H/0435/001	PL 01656/0214	KRKA, D.D., NOVO MESTO	UK
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg film-coated tablets	PT/H/1603/001	PL 16363/0485	MILPHARM LIMITED	UK
Etoricoxib 30 mg Film-coated Tablets	DE/H/5092/001	PL 36687/0210	HEUMANN PHARMA GMBH & CO. GENERICA KG	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
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg film-coated tablets	EE/H/0211/001	PL 17780/0724	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 30 mg Film-coated Tablets	NL/H/3576/001	PL 25258/0199	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Etoricoxib 30 mg Film-coated Tablets	DE/H/5031/001	PL 00289/1934	TEVA UK LIMITED	UK
Etoricoxib 30mg Film-coated Tablets	SE/H/1574/001	PL 30306/0768	ACTAVIS GROUP PTC EHF.	UK
Etoricoxib 60 mg Film-coated Tablets	ES/H/0264/002	PA 0688/032/002	CHANELLE MEDICAL	IE
Etoricoxib 60 mg Film-coated Tablets	NL/H/3151/002	PA0405/140/002	GENERICS [UK] LIMITED	IE
Etoricoxib 60 mg Film-coated tablets	DE/H/3908/002	PA0711/232/002	ROWEX LTD	IE
Etoricoxib 60 mg Film-coated Tablets	ES/H/0264/002	PA 0688/032/002	CHANELLE MEDICAL	IE
Etoricoxib 60 mg Film-coated Tablets	ES/H/0264/002	MA1026/00102	CHANELLE MEDICAL	MT
Etoricoxib 60 mg Film-coated Tablets	ES/H/0264/002	MA1026/00102	CHANELLE MEDICAL	MT
Etoricoxib 60 mg film-coated tablets	NL/H/3461/002	PL 42357/0168	AMNEAL PHARMA EUROPE LIMITED	UK
Etoricoxib 60 mg film-coated tablets	DE/H/3469/002	PL 31596/0002	ZYDUS FRANCE	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
Etoricoxib 60 mg film-coated tablets	HU/H/0435/002	PL 01656/0215	KRKA, D.D., NOVO MESTO	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	PT/H/1603/002	PL 16363/0486	MILPHARM LIMITED	UK
Etoricoxib 60 mg Film-coated Tablets	DE/H/5092/002	PL 36687/0211	TORRENT PHARMA LTD.	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	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
Etoricoxib 60 mg film-coated tablets	EE/H/0211/002	PL 17780/0725	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 60 mg Film-coated Tablets	NL/H/3576/002	PL 25258/0200	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Etoricoxib 60 mg Film-coated Tablets	DE/H/5031/003	PL 00289/1935	TEVA UK LIMITED	UK
Etoricoxib 60mg Film-coated Tablets	SE/H/1574/002	PL 30306/0769	ACTAVIS GROUP PTC EHF.	UK
Etoricoxib 90 mg Film-coated Tablets	ES/H/0264/003	PA 0688/032/003	CHANELLE MEDICAL	IE
Etoricoxib 90 mg Film-coated Tablets	NL/H/3151/003	PA0405/140/003	GENERICS [UK] LIMITED	IE
Etoricoxib 90 mg Film-coated tablets	DE/H/3908/003	PA0711/232/003	ROWEX LTD	IE
Etoricoxib 90 mg Film-coated Tablets	ES/H/0264/003	PA 0688/032/003	CHANELLE MEDICAL	IE
Etoricoxib 90 mg Film-coated Tablets	ES/H/0264/003	MA1026/00103	CHANELLE MEDICAL	MT
Etoricoxib 90 mg Film-coated Tablets	ES/H/0264/003	MA1026/00103	CHANELLE MEDICAL	MT
Etoricoxib 90 mg film-coated tablets	NL/H/3461/003	PL 42357/0169	AMNEAL PHARMA EUROPE LIMITED	UK
Etoricoxib 90 mg film-coated tablets	DE/H/3469/003	PL 31596/0003	ZYDUS FRANCE	UK
Etoricoxib 90 mg film-coated tablets	HU/H/0435/003	PL 01656/0216	KRKA, D.D., NOVO MESTO	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	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
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	PT/H/1603/003	PL 16363/0487	MILPHARM LIMITED	UK
Etoricoxib 90 mg Film-coated Tablets	DE/H/5092/003	PL 36687/0212	TORRENT PHARMA LTD.	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg film-coated tablets	EE/H/0211/003	PL 17780/0726	WINTHROP PHARMACEUTICALS UK LTD	UK
Etoricoxib 90 mg Film-coated Tablets	NL/H/3576/003	PL 25258/0201	GLENMARK PHARMACEUTICALS EUROPE LIMITED	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
Etoricoxib 90 mg Film-coated Tablets	DE/H/5031/003	PL 00289/1936	TEVA UK LIMITED	UK
Etoricoxib 90mg Film-coated Tablets	SE/H/1574/003	PL 30306/0770	ACTAVIS GROUP PTC EHF.	UK
Etoricoxib Actavis 120 mg apvalkotās tabletes	SE/H/1574/004	17-0037	ACTAVIS GROUP PTC EHF.	LV
Etoricoxib Actavis 120 mg film-coated tablets	SE/H/1574/004	MA628/15703	ACTAVIS GROUP PTC EHF.	MT
Etoricoxib Actavis 120 mg filmdragerade tabletter	SE/H/1574/004	53664	ACTAVIS GROUP PTC EHF.	SE
Etoricoxib Actavis 120 mg filmuhúðaðar töflur	SE/H/1574/004	IS/1/17/001/04	ACTAVIS GROUP PTC EHF.	IS
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/034	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/038	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/033	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/043	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/044	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/036	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/037	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/042	ACTAVIS GROUP PTC EHF.	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
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/035	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/040	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/039	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 120 mg plėvele dengtos tabletės	SE/H/1574/004	LT/1/17/4026/041	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg apvalkotās tabletes	SE/H/1574/001	17-0034	ACTAVIS GROUP PTC EHF.	LV
Etoricoxib Actavis 30 mg film-coated tablets	SE/H/1574/001	MA628/15704	ACTAVIS GROUP PTC EHF.	MT
Etoricoxib Actavis 30 mg filmdragerade tabletter	SE/H/1574/001	53661	ACTAVIS GROUP PTC EHF.	SE
Etoricoxib Actavis 30 mg filmuhúðaðar töflur	SE/H/1574/001	IS/1/17/001/01	ACTAVIS GROUP PTC EHF.	IS
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/004	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/005	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/003	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/002	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/001	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 30 mg plėvele dengtos tabletės	SE/H/1574/001	LT/1/17/4026/006	ACTAVIS GROUP PTC EHF.	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
Etoricoxib Actavis 60 mg apvalkotās tabletes	SE/H/1574/002	17-0035	ACTAVIS GROUP PTC EHF.	LV
Etoricoxib Actavis 60 mg film-coated tablets	SE/H/1574/002	MA628/15701	ACTAVIS GROUP PTC EHF.	MT
Etoricoxib Actavis 60 mg filmdragerade tabletter	SE/H/1574/002	53662	ACTAVIS GROUP PTC EHF.	SE
Etoricoxib Actavis 60 mg filmuhúðaðar töflur	SE/H/1574/002	IS/1/17/001/02	ACTAVIS GROUP PTC EHF.	IS
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/007	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/010	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/012	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/009	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/019	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/016	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/011	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/018	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/020	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/014	ACTAVIS GROUP PTC EHF.	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
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/015	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/017	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/008	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 60 mg plėvele dengtos tabletės	SE/H/1574/002	LT/1/17/4026/013	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg apvalkotās tabletes	SE/H/1574/003	17-0036	ACTAVIS GROUP PTC EHF.	LV
Etoricoxib Actavis 90 mg film-coated tablets	SE/H/1574/003	MA628/15702	ACTAVIS GROUP PTC EHF.	MT
Etoricoxib Actavis 90 mg filmdragerade tabletter	SE/H/1574/003	53663	ACTAVIS GROUP PTC EHF.	SE
Etoricoxib Actavis 90 mg filmuhúðaðar töflur	SE/H/1574/003	IS/1/17/001/03	ACTAVIS GROUP PTC EHF.	IS
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/022	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/030	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/021	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/027	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/026	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/028	ACTAVIS GROUP PTC EHF.	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
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/031	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/024	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/025	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/023	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/032	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis 90 mg plėvele dengtos tabletės	SE/H/1574/003	LT/1/17/4026/029	ACTAVIS GROUP PTC EHF.	LT
Etoricoxib Actavis, 120 mg õhukese polümeerikattega tabletid	SE/H/1574/004	931517	ACTAVIS GROUP PTC EHF.	EE
Etoricoxib Actavis, 30 mg õhukese polümeerikattega tabletid	SE/H/1574/001	931217	ACTAVIS GROUP PTC EHF.	EE
Etoricoxib Actavis, 60 mg õhukese polümeerikattega tabletid	SE/H/1574/002	931317	ACTAVIS GROUP PTC EHF.	EE
Etoricoxib Actavis, 90 mg õhukese polümeerikattega tabletid	SE/H/1574/003	931417	ACTAVIS GROUP PTC EHF.	EE
Etoricoxib AL 120 mg Filmtabletten	DE/H/4251/004	93721.00.00	ALIUD PHARMA GMBH	DE
Etoricoxib AL 30 mg Filmtabletten	DE/H/4251/001	93718.00.00	ALIUD PHARMA GMBH	DE
Etoricoxib AL 60 mg Filmtabletten	DE/H/4251/002	93719.00.00	ALIUD PHARMA GMBH	DE
Etoricoxib AL 90 mg Filmtabletten	DE/H/4251/003	93720.00.00	ALIUD PHARMA GMBH	DE
Etoricoxib Alchemia 120 mg filmdragerad tablett	SE/H/1575/04	53668	ALCHEMIA LIMITED	SE
Etoricoxib Alchemia 120 mg Filmtabletten	SE/H/1575/004	96699.00.00	ALCHEMIA LIMITED	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
Etoricoxib Alchemia 30 mg filmdragerad tablett	SE/H/1575/01	53665	ALCHEMIA LIMITED	SE
Etoricoxib Alchemia 30 mg Filmtabletten	SE/H/1575/001	96696.00.00	ALCHEMIA LIMITED	DE
Etoricoxib Alchemia 60 mg filmdragerad tablett	SE/H/1575/02	53666	ALCHEMIA LIMITED	SE
Etoricoxib Alchemia 60 mg Filmtabletten	SE/H/1575/002	96697.00.00	ALCHEMIA LIMITED	DE
Etoricoxib Alchemia 90 mg filmdragerad tablett	SE/H/1575/03	53667	ALCHEMIA LIMITED	SE
Etoricoxib Alchemia 90 mg Filmtabletten	SE/H/1575/003	96698.00.00	ALCHEMIA LIMITED	DE
Etoricoxib Alembic 120 mg comprimidos recubiertos con película EFG	DE/H/5092/003	81465	ALEMBIC PHARMACEUTICALS EUROPE LTD	ES
Etoricoxib Alembic 30 mg comprimidos recubiertos con película EFG	DE/H/5092/001	81463	ALEMBIC PHARMACEUTICALS EUROPE LTD	ES
Etoricoxib Alembic 60 mg comprimidos recubiertos con película EFG	DE/H/5092/002/DC	81462	ALEMBIC PHARMACEUTICALS EUROPE LTD	ES
Etoricoxib Alembic 90 mg comprimidos recubiertos con película EFG	DE/H/5092/003/DC	81464	ALEMBIC PHARMACEUTICALS EUROPE LTD	ES
Etoricoxib Alter 120 mg comprimidos recubiertos con película EFG	ES/H/0411/004	81.827	LABORATORIOS ALTER, 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
Etoricoxib Alter 30 mg comprimidos recubiertos con película EFG	ES/H/0411/001	81.824	LABORATORIOS ALTER, S.A.	ES
Etoricoxib Alter 60 mg comprimidos recubiertos con película EFG	ES/H/0411/002	81.825	LABORATORIOS ALTER, S.A.	ES
Etoricoxib Alter 90 mg comprimidos recubiertos con película EFG	ES/H/0411/003	81.826	LABORATORIOS ALTER, S.A.	ES
Etoricoxib Amneal 120 mg comprimidos recubiertos con película EFG	NL/H/3461/004	81.201	AMNEAL PHARMA EUROPE LIMITED	ES
Etoricoxib Amneal 120 mg filmdragerade tabletter	NL/H/3461/004	52506	AMNEAL PHARMA EUROPE LIMITED	SE
Etoricoxib Amneal 120 mg filmdrasjerte tabletter	NL/H/3461/004	15-10555	AMNEAL PHARMA EUROPE LIMITED	NO
Etoricoxib Amneal 120 mg Filmtabletten	NL/H/3461/004	94327.00.00	AMNEAL PHARMA EUROPE LIMITED	DE
Etoricoxib Amneal 120 mg tabletti, kalvopäällysteinen	NL/H/3461/004	33114	AMNEAL PHARMA EUROPE LIMITED	FI
Etoricoxib Amneal 120 mg, filmomhulde tabletten	NL/H/3461/004	RVG 116935	AMNEAL PHARMA EUROPE LIMITED	NL
Etoricoxib Amneal 30 mg comprimidos recubiertos con película EFG	NL/H/3461/001	81.198	AMNEAL PHARMA EUROPE LIMITED	ES
Etoricoxib Amneal 30 mg filmdragerade tabletter	NL/H/3461/001	52503	AMNEAL PHARMA EUROPE LIMITED	SE
Etoricoxib Amneal 30 mg filmdrasjerte tabletter	NL/H/3461/001	15-10552	AMNEAL PHARMA EUROPE LIMITED	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
Etoricoxib Amneal 30 mg Filmtabletten	NL/H/3461/001	94324.00.00	AMNEAL PHARMA EUROPE LIMITED	DE
Etoricoxib Amneal 30 mg tabletti, kalvopäällysteinen	NL/H/3461/001	33111	AMNEAL PHARMA EUROPE LIMITED	FI
Etoricoxib Amneal 30 mg, filmomhulde tabletten	NL/H/3461/001	RVG 116932	AMNEAL PHARMA EUROPE LIMITED	NL
Etoricoxib Amneal 60 mg comprimidos recubiertos con película EFG	NL/H/3461/002	81.199	AMNEAL PHARMA EUROPE LIMITED	ES
Etoricoxib Amneal 60 mg filmdragerade tabletter	NL/H/3461/002	52504	AMNEAL PHARMA EUROPE LIMITED	SE
Etoricoxib Amneal 60 mg filmdrasjerte tabletter	NL/H/3461/002	15-10553	AMNEAL PHARMA EUROPE LIMITED	NO
Etoricoxib Amneal 60 mg Filmtabletten	NL/H/3461/002	94325.00.00	AMNEAL PHARMA EUROPE LIMITED	DE
Etoricoxib Amneal 60 mg tabletti, kalvopäällysteinen	NL/H/3461/002	33112	AMNEAL PHARMA EUROPE LIMITED	FI
Etoricoxib Amneal 60 mg, filmomhulde tabletten	NL/H/3461/002	RVG 116933	AMNEAL PHARMA EUROPE LIMITED	NL
Etoricoxib Amneal 90 mg comprimidos recubiertos con película EFG	NL/H/3461/003	81.200	AMNEAL PHARMA EUROPE LIMITED	ES
Etoricoxib Amneal 90 mg filmdragerade tabletter	NL/H/3461/003	52505	AMNEAL PHARMA EUROPE LIMITED	SE
Etoricoxib Amneal 90 mg filmdrasjerte tabletter	NL/H/3461/003	15-10554	AMNEAL PHARMA EUROPE LIMITED	NO
Etoricoxib Amneal 90 mg Filmtabletten	NL/H/3461/003	94326.00.00	AMNEAL PHARMA EUROPE LIMITED	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
Etoricoxib Amneal 90 mg tabletti, kalvopäällysteinen	NL/H/3461/003	33113	AMNEAL PHARMA EUROPE LIMITED	FI
Etoricoxib Amneal 90 mg, filmomhulde tabletten	NL/H/3461/003	RVG 116934	AMNEAL PHARMA EUROPE LIMITED	NL
Etoricoxib Apotex 120 mg comprimidos recubiertos con película EFG	not available	81060	APOTEX EUROPE B.V.	ES
Etoricoxib Apotex 30 mg comprimidos recubiertos con película EFG	not available	81052	APOTEX EUROPE B.V.	ES
Etoricoxib Apotex 60 mg comprimidos recubiertos con película EFG	not available	81053	APOTEX EUROPE B.V.	ES
Etoricoxib Apotex 90 mg comprimidos recubiertos con película EFG	not available	81059	APOTEX EUROPE B.V.	ES
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	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
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg comprimate filmate	PT/H/1603/004	9898/2017/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 120 mg, filmomhulde tabletten	PT/H/1603/004	RVG 118782	AUROBINDO PHARMA B.V.	NL
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg comprimate filmate	PT/H/1603/001	9895/2017/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 30 mg, filmomhulde tabletten	PT/H/1603/001	RVG 118779	AUROBINDO PHARMA B.V.	NL
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	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
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg comprimate filmate	PT/H/1603/002	9896/2017/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 60 mg, filmomhulde tabletten	PT/H/1603/002	RVG 118780	AUROBINDO PHARMA B.V.	NL
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	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
Etoricoxib Aurobindo 90 mg comprimate filmate	PT/H/1603/003	9897/2017/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Etoricoxib Aurobindo 90 mg, filmomhulde tabletten	PT/H/1603/003	RVG 118781	AUROBINDO PHARMA B.V.	NL
Etoricoxib Aurovitas 120 mg comprimidos recubiertos con película EFG	PT/H/1614/004	81.890	AUROVITAS SPAIN,S.A.U.	ES
Etoricoxib Aurovitas 120 mg comprimidos revestidos por película	not available	5704846	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 120 mg comprimidos revestidos por película	not available	16/H/0015/004	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 120 mg comprimidos revestidos por película	not available	16/H/0015/004	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 30 mg comprimidos recubiertos con película EFG	PT/H/1614/001	81.887	AUROVITAS SPAIN,S.A.U.	ES
Etoricoxib Aurovitas 30 mg comprimidos revestidos por película	not available	16/H/0015/001	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 30 mg comprimidos revestidos por película	not available	5704465	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 30 mg comprimidos revestidos por película	not available	16/H/0015/001	AUROVITAS UNIPESSOAL, 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
Etoricoxib Aurovitas 60 mg comprimidos recubiertos con película EFG	PT/H/1614/002	81.889	AUROVITAS SPAIN,S.A.U.	ES
Etoricoxib Aurovitas 60 mg comprimidos revestidos por película	not available	16/H/0015/002	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 60 mg comprimidos revestidos por película	not available	5704820	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 60 mg comprimidos revestidos por película	not available	16/H/0015/002	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 90 mg comprimidos recubiertos con película EFG	PT/H/1614/003	81.888	AUROVITAS SPAIN,S.A.U.	ES
Etoricoxib Aurovitas 90 mg comprimidos revestidos por película	not available	16/H/0015/003	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 90 mg comprimidos revestidos por película	not available	5704812	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib Aurovitas 90 mg comprimidos revestidos por película	not available	16/H/0015/003	AUROVITAS UNIPESSOAL, LDA.	PT
Etoricoxib beta 120 mg Filmtabletten	DE/H/4864/004	97658.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Etoricoxib beta 30 mg Filmtabletten	DE/H/4864/001	97655.00.00	BETAPHARM ARZNEIMITTEL 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
Etoricoxib beta 60 mg Filmtabletten	DE/H/4864/002	97656.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Etoricoxib beta 90 mg Filmtabletten	DE/H/4864/003	97657.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Etoricoxib Bluepharma 120 mg comprimidos revestidos por película	not available	5693841	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 120 mg comprimidos revestidos por película	not available	5693858	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 120 mg comprimidos revestidos por película	not available	16/H/0071/004	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 120 mg comprimidos revestidos por película	not available	16/H/0071/004	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 30 mg comprimidos revestidos por película	not available	5693809	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 30 mg comprimidos revestidos por película	not available	5693817	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 30 mg comprimidos revestidos por película	not available	16/H/0071/001	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 30 mg comprimidos revestidos por película	not available	16/H/0071/001	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, 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
Etoricoxib Bluepharma 60 mg comprimidos revestidos por película	not available	5693767	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 60 mg comprimidos revestidos por película	not available	5693775	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 60 mg comprimidos revestidos por película	not available	16/H/0071/002	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 60 mg comprimidos revestidos por película	not available	16/H/0071/002	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 90 mg comprimidos revestidos por película	not available	5693825	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 90 mg comprimidos revestidos por película	not available	5693833	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 90 mg comprimidos revestidos por película	not available	16/H/0071/003	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib Bluepharma 90 mg comprimidos revestidos por película	not available	16/H/0071/003	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Etoricoxib CF 120 mg, filmomhulde tabletten	DE/H/4251/004	RVG 116412	CENTRAFARM B.V.	NL
Etoricoxib CF 30 mg, filmomhulde tabletten	DE/H/4251/001	RVG 116409	CENTRAFARM B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Etoricoxib CF 60 mg, filmomhulde tabletten	DE/H/4251/002	RVG 116410	CENTRAFARM B.V.	NL
Etoricoxib CF 90 mg, filmomhulde tabletten	DE/H/4251/003	RVG 116411	CENTRAFARM B.V.	NL
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239159	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239161	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239173	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239159	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239159	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239161	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239173	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg compresse rivestite con film	ES/H/0264/004	043239159	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644869	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644877	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644901	CHANELLE MEDICAL	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
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644851	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644869	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644877	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644901	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 120 mg comprimidos revestidos por película	ES/H/0264/004	5644851	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 30 mg compresse rivestite con film	ES/H/0264/001	043239019	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 30 mg compresse rivestite con film	ES/H/0264/001	043239019	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 30 mg comprimidos revestidos por película	ES/H/0264/001	5644919	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 30 mg comprimidos revestidos por película	ES/H/0264/001	5644919	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239021	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239033	CHANELLE MEDICAL	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
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239045	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239058	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239060	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239072	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239021	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239033	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239045	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239058	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239060	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg compresse rivestite con film	ES/H/0264/002	043239072	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644802	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644810	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644828	CHANELLE MEDICAL	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
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644778	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644836	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644844	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644802	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644810	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644828	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644778	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644836	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 60 mg comprimidos revestidos por película	ES/H/0264/002	5644844	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239084	CHANELLE MEDICAL	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
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239096	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239110	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239122	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239134	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239108	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239146	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239084	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239096	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239110	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239122	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239134	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239108	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg compresse rivestite con film	ES/H/0264/003	043239146	CHANELLE MEDICAL	IT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	564435	CHANELLE MEDICAL	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
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644943	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644968	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644927	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644950	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644976	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5645007	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	564435	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644943	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644968	CHANELLE MEDICAL	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
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644927	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644950	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5644976	CHANELLE MEDICAL	PT
Etoricoxib Chanelle 90 mg comprimidos revestidos por película	ES/H/0264/003	5645007	CHANELLE MEDICAL	PT
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/02	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/04	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/02	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 120 mg comprimate filmate	ES/H/0264/004	7269/2014/04	CHANELLE MEDICAL	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
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/02	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/04	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/05	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/06	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/02	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/04	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/05	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 60 mg comprimate filmate	ES/H/0264/002	7267/2014/06	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/02	CHANELLE MEDICAL	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
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/04	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/05	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/06	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/07	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/02	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/03	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/04	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/05	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/06	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical 90 mg comprimate filmate	ES/H/0264/003	7268/2014/07	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical30 mg comprimate filmate	ES/H/0264/001	7266/2014/01	CHANELLE MEDICAL	RO
Etoricoxib Chanelle Medical30 mg comprimate filmate	ES/H/0264/001	7266/2014/01	CHANELLE MEDICAL	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
Etoricoxib Ciclum	DE/H/4251/003	5692868	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	5695341	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	5692843	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	5692835	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	5695333	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	5692850	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL 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
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum	DE/H/4251/003	DE/H/4251/003	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	5692876	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL 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
Etoricoxib Ciclum 120 mg comprimidos revestidos por película	DE/H/4251/004	DE/H/4251/004	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 30 mg comprimidos revestidos por película	DE/H/4251/001	5692751	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 30 mg comprimidos revestidos por película	DE/H/4251/001	5692769	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 30 mg comprimidos revestidos por película	DE/H/4251/001	DE/H/4251/001	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 30 mg comprimidos revestidos por película	DE/H/4251/001	DE/H/4251/001	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib Ciclum 60 mg comprimidos revestidos por película	DE/H/4251/002	DE/H/4251/002	CICLUM FARMA UNIPESSOAL LDA.	PT
Etoricoxib cinfa 120 mg comprimidos recubiertos con película EFG	not available	80754	LABORATORIOS CINFA, S.A.	ES
Etoricoxib cinfa 30 mg comprimidos recubiertos con película EFG	not available	80752	LABORATORIOS CINFA, S.A.	ES
Etoricoxib cinfa 60 mg comprimidos recubiertos con película EFG	not available	80751	LABORATORIOS CINFA, 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
Etoricoxib cinfa 90 mg comprimidos recubiertos con película EFG	not available	80753	LABORATORIOS CINFA, S.A.	ES
ETORICOXIB DOC Generici 120 mg compresse rivestite con film	NL/H/3272/003	043688035	DOC GENERICI S.R.L.	IT
Etoricoxib DOC Generici 120 mg film-coated tablets	NL/H/3272/003	RVG 116035	DOC GENERICI S.R.L.	NL
ETORICOXIB DOC Generici 60 mg compresse rivestite con film	NL/H/3272/001	043688011	DOC GENERICI S.R.L.	IT
Etoricoxib DOC Generici 60 mg film-coated tablets	NL/H/3272/001	RVG 116033	DOC GENERICI S.R.L.	NL
ETORICOXIB DOC Generici 90 mg compresse rivestite con film	NL/H/3272/002	043688023	DOC GENERICI S.R.L.	IT
Etoricoxib DOC Generici 90 mg film-coated tablets	NL/H/3272/002	RVG 116034	DOC GENERICI S.R.L.	NL
Etoricoxib EG 120 mg comprimés pelliculés	DE/H/4251/004	BE500497	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 120 mg comprimés pelliculés	DE/H/4251/004	2017040093	EUROGENERICS N.V./S.A.	LU
Etoricoxib EG 120 mg filmomhulde tabletten	DE/H/4251/004	BE500497	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 120 mg Filmtabletten	DE/H/4251/004	BE500497	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 30 mg comprimés pelliculés	DE/H/4251/001	BE500444	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 30 mg comprimés pelliculés	DE/H/4251/001	2017040090	EUROGENERICS N.V./S.A.	LU
Etoricoxib EG 30 mg filmomhulde tabletten	DE/H/4251/001	BE500444	EUROGENERICS N.V./S.A.	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
Etoricoxib EG 30 mg Filmtabletten	DE/H/4251/001	BE500444	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 60 mg comprimés pelliculés	DE/H/4251/002	BE500453	EUROGENERICS SA	BE
Etoricoxib EG 60 mg comprimés pelliculés	DE/H/4251/002	BE500462	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 60 mg comprimés pelliculés	DE/H/4251/002	2017040091	EUROGENERICS N.V./S.A.	LU
Etoricoxib EG 60 mg filmomhulde tabletten	DE/H/4251/002	BE500453	EUROGENERICS SA	BE
Etoricoxib EG 60 mg filmomhulde tabletten	DE/H/4251/002	BE500462	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 60 mg Filmtabletten	DE/H/4251/002	BE500453	EUROGENERICS SA	BE
Etoricoxib EG 60 mg Filmtabletten	DE/H/4251/002	BE500462	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 90 mg comprimés pelliculés	DE/H/4251/003	BE500471	EUROGENERICS SA	BE
Etoricoxib EG 90 mg comprimés pelliculés	DE/H/4251/003	BE500480	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 90 mg comprimés pelliculés	DE/H/4251/003	2017040092	EUROGENERICS N.V./S.A.	LU
Etoricoxib EG 90 mg filmomhulde tabletten	DE/H/4251/003	BE500480	EUROGENERICS N.V./S.A.	BE
Etoricoxib EG 90 mg filmomhulde tabletten	DE/H/4251/003	BE500471	EUROGENERICS SA	BE
Etoricoxib EG 90 mg Filmtabletten	DE/H/4251/003	BE500471	EUROGENERICS SA	BE
Etoricoxib EG 90 mg Filmtabletten	DE/H/4251/003	BE500480	EUROGENERICS N.V./S.A.	BE
Etoricoxib Empower 120 mg comprimidos revestidos por película	PT/H/1614/004	5710355	EMPOWER PHARMA	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
Etoricoxib Empower 120 mg comprimidos revestidos por película	PT/H/1614/004	5710355	EMPOWER PHARMA	PT
Etoricoxib Empower 30 mg comprimidos revestidos por película	PT/H/1614/001	5710140	EMPOWER PHARMA	PT
Etoricoxib Empower 30 mg comprimidos revestidos por película	PT/H/1614/001	5710140	EMPOWER PHARMA	PT
Etoricoxib Empower 60 mg comprimidos revestidos por película	PT/H/1614/002	5710231	EMPOWER PHARMA	PT
Etoricoxib Empower 60 mg comprimidos revestidos por película	PT/H/1614/002	5710231	EMPOWER PHARMA	PT
Etoricoxib Empower 90 mg comprimidos revestidos por película	PT/H/1614/003	5710330	EMPOWER PHARMA	PT
Etoricoxib Empower 90 mg comprimidos revestidos por película	PT/H/1614/003	5710330	EMPOWER PHARMA	PT
Etoricoxib Farmoz 120 mg comprimidos revestidos por película	not available	5691852	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 120 mg comprimidos revestidos por película	not available	5691860	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, 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
Etoricoxib Farmoz 120 mg comprimidos revestidos por película	not available	16/H/0023/004	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 120 mg comprimidos revestidos por película	not available	16/H/0023/004	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 30 mg comprimidos revestidos por película	not available	5691712	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 30 mg comprimidos revestidos por película	not available	5691720	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 30 mg comprimidos revestidos por película	not available	16/H/0023/001	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 30 mg comprimidos revestidos por película	not available	16/H/0023/001	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 60 mg comprimidos revestidos por película	not available	5691811	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 60 mg comprimidos revestidos por película	not available	5691829	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 60 mg comprimidos revestidos por película	not available	16/H/0023/002	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, 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
Etoricoxib Farmoz 60 mg comprimidos revestidos por película	not available	16/H/0023/002	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 90 mg comprimidos revestidos por película	not available	5691837	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 90 mg comprimidos revestidos por película	not available	5691845	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 90 mg comprimidos revestidos por película	not available	16/H/0023/003	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Farmoz 90 mg comprimidos revestidos por película	not available	16/H/0023/003	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, 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
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/004	5704622	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 120 mg comprimidos revestidos por película	PT/H/1603/004	5704648	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001	5709571	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001	5704630	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001	5709605	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001/DC	PT/H/1603/001/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001/DC	PT/H/1603/001/DC	GENERIS FARMACÊUTICA, 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
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001/DC	PT/H/1603/001/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001/DC	PT/H/1603/001/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 30 mg comprimidos revestidos por película	PT/H/1603/001/DC	PT/H/1603/001/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002	5709902	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002	5709910	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002	5704663	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002/DC	PT/H/1603/002/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002/DC	PT/H/1603/002/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002/DC	PT/H/1603/002/DC	GENERIS FARMACÊUTICA, 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
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002/DC	PT/H/1603/002/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 60 mg comprimidos revestidos por película	PT/H/1603/002/DC	PT/H/1603/002/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003	5709928	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003	5704671	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003	5709936	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, 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
Etoricoxib Generis 90 mg comprimidos revestidos por película	PT/H/1603/003/DC	PT/H/1603/003/DC	GENERIS FARMACÊUTICA, S.A.	PT
Etoricoxib Glenmark 120 mg Filmtabletten	NL/H/3576/004	96165.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Etoricoxib Glenmark 120 mg, filmomhulde tabletten	NL/H/3576/004	RVG 117989	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Etoricoxib Glenmark 30 mg Filmtabletten	NL/H/3576/001	96162.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Etoricoxib Glenmark 30 mg, filmomhulde tabletten	NL/H/3576/001	RVG 117986	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Etoricoxib Glenmark 60 mg Filmtabletten	NL/H/3576/002	96163.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Etoricoxib Glenmark 60 mg, filmomhulde tabletten	NL/H/3576/002	RVG 117987	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Etoricoxib Glenmark 90 mg Filmtabletten	NL/H/3576/003	96164.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Etoricoxib Glenmark 90 mg, filmomhulde tabletten	NL/H/3576/003	RVG 117988	GLENMARK PHARMACEUTICALS EUROPE LIMITED	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
Etoricoxib HCS 120 mg comprimidos revestidos por película	HU/H/0435/004	5706924	HCS BVBA	PT
Etoricoxib HCS 120 mg comprimidos revestidos por película	HU/H/0435/004	5706932	HCS BVBA	PT
Etoricoxib HCS 120 mg comprimidos revestidos por película	HU/H/0435/004	5706940	HCS BVBA	PT
Etoricoxib HCS 30 mg comprimidos revestidos por película	HU/H/0435/001	5706742	HCS BVBA	PT
Etoricoxib HCS 30 mg comprimidos revestidos por película	HU/H/0435/001	5706759	HCS BVBA	PT
Etoricoxib HCS 60 mg comprimidos revestidos por película	HU/H/0435/002	5706775	HCS BVBA	PT
Etoricoxib HCS 60 mg comprimidos revestidos por película	HU/H/0435/002	5706809	HCS BVBA	PT
Etoricoxib HCS 60 mg comprimidos revestidos por película	HU/H/0435/002	5706817	HCS BVBA	PT
Etoricoxib HCS 90 mg comprimidos revestidos por película	HU/H/0435/003	5706833	HCS BVBA	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
Etoricoxib HCS 90 mg comprimidos revestidos por película	HU/H/0435/003	5706841	HCS BVBA	PT
Etoricoxib HCS 90 mg comprimidos revestidos por película	HU/H/0435/003	5706858	HCS BVBA	PT
Etoricoxib Heumann 120 mg Filmtabletten	DE/H/5092/004	93517.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Etoricoxib Heumann 30 mg Filmtabletten	DE/H/5092/001	93514.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Etoricoxib Heumann 60 mg Filmtabletten	DE/H/5092/002	93515.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Etoricoxib Heumann 90 mg Filmtabletten	DE/H/5092/003	93516.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Etoricoxib Krka 120 mg comprimés pelliculés	HU/H/0435/004	BE501706	KRKA, D.D., NOVO MESTO	BE
Etoricoxib Krka 120 mg comprimidos recubiertos con película EFG	HU/H/0435/004	81717	KRKA, D.D., NOVO MESTO	ES
Etoricoxib Krka 120 mg film-coated tablets	HU/H/0435/004	PA1347/064/004	KRKA, D.D., NOVO MESTO	IE
Etoricoxib Krka 120 mg filmdragerade tabletter	HU/H/0435/004	53602	KRKA, D.D., NOVO MESTO	SE
Etoricoxib Krka 120 mg filmdrasjerte tabletter	HU/H/0435/004	15-10943	KRKA, D.D., NOVO MESTO	NO
Etoricoxib Krka 120 mg filmuhúðaðar töflur	HU/H/0435/004	IS/1/16/099/04	KRKA, D.D., NOVO MESTO	IS
Etoricoxib Krka 120 mg kalvopäällysteiset tabletit	HU/H/0435/004	33739	KRKA, D.D., NOVO MESTO	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
Etoricoxib Krka 30 mg comprimés pelliculés	HU/H/0435/001	BE501671	KRKA, D.D., NOVO MESTO	BE
Etoricoxib Krka 30 mg comprimidos recubiertos con película EFG	HU/H/0435/001	81709	KRKA, D.D., NOVO MESTO	ES
Etoricoxib Krka 30 mg film-coated tablets	HU/H/0435/001	PA1347/064/001	KRKA, D.D., NOVO MESTO	IE
Etoricoxib Krka 30 mg filmdragerade tabletter	HU/H/0435/001	53599	KRKA, D.D., NOVO MESTO	SE
Etoricoxib Krka 30 mg filmdrasjerte tabletter	HU/H/0435/001	15-10940	KRKA, D.D., NOVO MESTO	NO
Etoricoxib Krka 30 mg filmuhúðaðar töflur	HU/H/0435/001	IS/1/16/099/01	KRKA, D.D., NOVO MESTO	IS
Etoricoxib Krka 30 mg kalvopäällysteiset tabletit	HU/H/0435/001	33736	KRKA, D.D., NOVO MESTO	FI
Etoricoxib Krka 60 mg comprimés pelliculés	HU/H/0435/002	BE501680	KRKA, D.D., NOVO MESTO	BE
Etoricoxib Krka 60 mg comprimidos recubiertos con película EFG	HU/H/0435/002	81710	KRKA, D.D., NOVO MESTO	ES
Etoricoxib Krka 60 mg film-coated tablets	HU/H/0435/002	PA1347/064/002	KRKA, D.D., NOVO MESTO	IE
Etoricoxib Krka 60 mg filmdragerade tabletter	HU/H/0435/002	53600	KRKA, D.D., NOVO MESTO	SE
Etoricoxib Krka 60 mg filmdrasjerte tabletter	HU/H/0435/002	15-10941	KRKA, D.D., NOVO MESTO	NO
Etoricoxib Krka 60 mg filmuhúðaðar töflur	HU/H/0435/002	IS/1/16/099/02	KRKA, D.D., NOVO MESTO	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
Etoricoxib Krka 60 mg kalvopäällysteiset tabletit	HU/H/0435/002	33737	KRKA, D.D., NOVO MESTO	FI
Etoricoxib Krka 90 mg comprimés pelliculés	HU/H/0435/003	BE501697	KRKA, D.D., NOVO MESTO	BE
Etoricoxib Krka 90 mg comprimidos recubiertos con película EFG	HU/H/0435/003	81711	KRKA, D.D., NOVO MESTO	ES
Etoricoxib Krka 90 mg film-coated tablets	HU/H/0435/003	PA1347/064/003	KRKA, D.D., NOVO MESTO	IE
Etoricoxib Krka 90 mg filmdragerade tabletter	HU/H/0435/003	53601	KRKA, D.D., NOVO MESTO	SE
Etoricoxib Krka 90 mg filmdrasjerte tabletter	HU/H/0435/003	15-10942	KRKA, D.D., NOVO MESTO	NO
Etoricoxib Krka 90 mg filmuhúðaðar töflur	HU/H/0435/003	IS/1/16/099/03	KRKA, D.D., NOVO MESTO	IS
Etoricoxib Krka 90 mg kalvopäällysteiset tabletit	HU/H/0435/003	33738	KRKA, D.D., NOVO MESTO	FI
Etoricoxib Libra-Pharm 120 mg Filmtabletten	UK/H/3190/004	77702.00.00	MERCK SHARP & DOHME BV	DE
Etoricoxib Libra-Pharm 30 mg Filmtabletten	UK/H/3190/001	77699.00.00	MERCK SHARP & DOHME BV	DE
Etoricoxib Libra-Pharm 60 mg Filmtabletten	UK/H/3190/002	77700.00.00	MERCK SHARP & DOHME BV	DE
Etoricoxib Libra-Pharm 90 mg Filmtabletten	UK/H/3190/003	77701.00.00	MERCK SHARP & DOHME BV	DE
Etoricoxib Micro Labs 120 mg Filmtabletten	ES/H/0264/004	91239.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 120 mg Filmtabletten	ES/H/0264/004	91239.00.00	MICRO LABS 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
Etoricoxib Micro Labs 120 mg Filmtabletten	ES/H/0264/004	91239.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 120 mg Filmtabletten	ES/H/0264/004	91239.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 120 mg Filmtabletten	ES/H/0264/004	91239.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 30 mg Filmtabletten	ES/H/0264/001	91236.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 30 mg Filmtabletten	ES/H/0264/001	91236.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 60 mg Filmtabletten	ES/H/0264/002	91237.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS 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
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Micro Labs 90 mg Filmtabletten	ES/H/0264/003	91238.00.00	MICRO LABS GMBH	DE
Etoricoxib Mylan 120 mg compresse rivestite con film	NL/H/3196/004	043546112	MYLAN S.P.A.	IT
Etoricoxib Mylan 120 mg compresse rivestite con film	NL/H/3196/004	043546124	MYLAN S.P.A.	IT
Etoricoxib Mylan 120 mg comprimidos recubiertos con película EFG	NL/H/3151/004	80376	MYLAN PHARMACEUTICALS S.L.	ES
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662507	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662428	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662465	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662515	MYLAN, 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
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662523	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5662531	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5689633	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg comprimidos revestidos por película	NL/H/3151/004	5689641	MYLAN, LDA	PT
Etoricoxib Mylan 120 mg Film-coated Tablets	NL/H/3151/004	PL 04569/1467	GENERICS [UK] LIMITED	UK
Etoricoxib Mylan 120 mg Filmtabletten	NL/H/3151/004	92028.00.00	GENERICS [UK] LIMITED	DE
Etoricoxib Mylan 120 mg tabletti, kalvopäällysteinen	NL/H/3151/004	32283	MYLAN AB	FI
Etoricoxib Mylan 120 mg, filmomhulde tabletten	NL/H/3151/004	RVG 115517	MYLAN B.V.	NL
Etoricoxib Mylan 120 mg, filmomhulde tabletten	NL/H/3196/004	RVG 115521	MYLAN B.V.	NL
Etoricoxib Mylan 30 mg compresse rivestite con film	NL/H/3196/001	043546011	MYLAN S.P.A.	IT
Etoricoxib Mylan 30 mg compresse rivestite con film	NL/H/3196/001	043546023	MYLAN S.P.A.	IT
Etoricoxib Mylan 30 mg compresse rivestite con film	NL/H/3196/001	043546035	MYLAN 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
Etoricoxib Mylan 30 mg compresse rivestite con film	NL/H/3196/001	043546047	MYLAN S.P.A.	IT
Etoricoxib Mylan 30 mg comprimidos recubiertos con película EFG	NL/H/3151/001	80373	MYLAN PHARMACEUTICALS S.L.	ES
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662234	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662218	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662200	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662226	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662267	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662259	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662242	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5662275	MYLAN, 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
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5689658	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg comprimidos revestidos por película	NL/H/3151/001	5689666	MYLAN, LDA	PT
Etoricoxib Mylan 30 mg Film-coated Tablets	NL/H/3151/001	PL 04569/1464	GENERICS [UK] LIMITED	UK
Etoricoxib Mylan 30 mg Filmtabletten	NL/H/3151/001	92025.00.00	GENERICS [UK] LIMITED	DE
Etoricoxib Mylan 30 mg tabletti, kalvopäällysteinen	NL/H/3151/001	32280	MYLAN AB	FI
Etoricoxib Mylan 30 mg, filmomhulde tabletten	NL/H/3151/001	RVG 115514	MYLAN B.V.	NL
Etoricoxib Mylan 30 mg, filmomhulde tabletten	NL/H/3196/001	RVG 115518	MYLAN B.V.	NL
Etoricoxib Mylan 60 mg compresse rivestite con film	NL/H/3196/002	043546050	MYLAN S.P.A.	IT
Etoricoxib Mylan 60 mg compresse rivestite con film	NL/H/3196/002	043546062	MYLAN S.P.A.	IT
Etoricoxib Mylan 60 mg comprimidos recubiertos con película EFG	NL/H/3151/002	80374	MYLAN PHARMACEUTICALS S.L.	ES
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662309	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662325	MYLAN, 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
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662333	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662317	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662366	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662341	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662374	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5662358	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5689617	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg comprimidos revestidos por película	NL/H/3151/002	5689625	MYLAN, LDA	PT
Etoricoxib Mylan 60 mg Film-coated Tablets	NL/H/3151/002	PL 04569/1465	GENERICS [UK] LIMITED	UK
Etoricoxib Mylan 60 mg Filmtabletten	NL/H/3151/002	92026.00.00	GENERICS [UK] LIMITED	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
Etoricoxib Mylan 60 mg tabletti, kalvopäällysteinen	NL/H/3151/002	32281	MYLAN AB	FI
Etoricoxib Mylan 60 mg, filmomhulde tabletten	NL/H/3151/002	RVG 115515	MYLAN B.V.	NL
Etoricoxib Mylan 60 mg, filmomhulde tabletten	NL/H/3196/002	RVG 115519	MYLAN B.V.	NL
Etoricoxib Mylan 90 mg compresse rivestite con film	NL/H/3196/003	043546074	MYLAN S.P.A.	IT
Etoricoxib Mylan 90 mg compresse rivestite con film	NL/H/3196/003	043546086	MYLAN S.P.A.	IT
Etoricoxib Mylan 90 mg compresse rivestite con film	NL/H/3196/003	043546098	MYLAN S.P.A.	IT
Etoricoxib Mylan 90 mg compresse rivestite con film	NL/H/3196/003	043546100	MYLAN S.P.A.	IT
Etoricoxib Mylan 90 mg comprimidos recubiertos con película EFG	NL/H/3151/003	80375	MYLAN PHARMACEUTICALS S.L.	ES
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662416	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662432	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662408	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662424	MYLAN, 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
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662440	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5662457	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5689575	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg comprimidos revestidos por película	NL/H/3151/003	5689609	MYLAN, LDA	PT
Etoricoxib Mylan 90 mg Film-coated Tablets	NL/H/3151/003	PL 04569/1466	GENERICS [UK] LIMITED	UK
Etoricoxib Mylan 90 mg Filmtabletten	NL/H/3151/003	92027.00.00	GENERICS [UK] LIMITED	DE
Etoricoxib Mylan 90 mg tabletti, kalvopäällysteinen	NL/H/3151/003	32282	MYLAN AB	FI
Etoricoxib Mylan 90 mg, filmomhulde tabletten	NL/H/3151/003	RVG 115516	MYLAN B.V.	NL
Etoricoxib Mylan 90 mg, filmomhulde tabletten	NL/H/3196/003	RVG 115520	MYLAN B.V.	NL
Etoricoxib Normon 120 mg comprimidos revestidos por película	ES/H/0415/004/DC	5711635	LABORATÓRIOS NORMON, S.A.	PT
Etoricoxib Normon 120 mg comprimidos revestidos por película	ES/H/0415/004/DC	5711643	LABORATÓRIOS NORMON, 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
Etoricoxib Normon 120 mg comprimidos revestidos por película	ES/H/0415/004/DC	5711650	LABORATÓRIOS NORMON, S.A.	PT
Etoricoxib Normon 30 mg comprimidos revestidos por película	ES/H/0415/001/DC	5710967	LABORATÓRIOS NORMON, S.A.	ES
Etoricoxib Normon 30 mg comprimidos revestidos por película	ES/H/0415/001/DC	5710975	LABORATÓRIOS NORMON, S.A.	ES
Etoricoxib Normon 30 mg comprimidos revestidos por película	ES/H/0415/001/DC	5710959	LABORATÓRIOS NORMON, S.A.	ES
Etoricoxib Normon 60 mg comprimidos revestidos por película	ES/H/0415/001/DC	5711577	LABORATÓRIOS NORMON, S.A.	ES
Etoricoxib Normon 60 mg comprimidos revestidos por película	ES/H/0415/002/DC	5711551	LABORATÓRIOS NORMON, S.A.	ES
Etoricoxib Normon 60 mg comprimidos revestidos por película	ES/H/0415/002/DC	5711569	LABORATÓRIOS NORMON, S.A.	PT
Etoricoxib Normon 90 mg comprimidos revestidos por película	ES/H/0415/003	5711601	LABORATÓRIOS NORMON, S.A.	PT
Etoricoxib Normon 90 mg comprimidos revestidos por película	ES/H/0415/003	5711619	LABORATÓRIOS NORMON, 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
Etoricoxib Normon 90 mg comprimidos revestidos por película	ES/H/0415/003	5711627	LABORATÓRIOS NORMON, S.A.	PT
Etoricoxib Orion 120 mg filmdragerade tabletter	SE/H/1554/004	33618	ORION CORPORATION	FI
Etoricoxib Orion 120 mg filmdragerade tabletter	SE/H/1554/004	53358	ORION CORPORATION	SE
Etoricoxib Orion 120 mg filmdrasjerte tabletter	SE/H/1554/004	15-10838	ORION CORPORATION	NO
Etoricoxib Orion 120 mg kalvopäällysteiset tabletit	SE/H/1554/004	33618	ORION CORPORATION	FI
Etoricoxib Orion 30 mg filmdragerade tabletter	SE/H/1554/001	33615	ORION CORPORATION	FI
Etoricoxib Orion 30 mg filmdragerade tabletter	SE/H/1554/001	53355	ORION CORPORATION	SE
Etoricoxib Orion 30 mg filmdrasjerte tabletter	SE/H/1554/001	15-10835	ORION CORPORATION	NO
Etoricoxib Orion 30 mg kalvopäällysteiset tabletit	SE/H/1554/001	33615	ORION CORPORATION	FI
Etoricoxib Orion 60 mg filmdragerade tabletter	SE/H/1554/002	33616	ORION CORPORATION	FI
Etoricoxib Orion 60 mg filmdragerade tabletter	SE/H/1554/002	53356	ORION CORPORATION	SE
Etoricoxib Orion 60 mg filmdrasjerte tabletter	SE/H/1554/002	15-10836	ORION CORPORATION	NO
Etoricoxib Orion 60 mg kalvopäällysteiset tabletit	SE/H/1554/002	33616	ORION CORPORATION	FI
Etoricoxib Orion 90 mg filmdragerade tabletter	SE/H/1554/003	33617	ORION CORPORATION	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
Etoricoxib Orion 90 mg filmdragerade tabletter	SE/H/1554/003	53357	ORION CORPORATION	SE
Etoricoxib Orion 90 mg filmdrasjerte tabletter	SE/H/1554/003	15-10837	ORION CORPORATION	NO
Etoricoxib Orion 90 mg kalvopäällysteiset tabletit	SE/H/1554/003	33617	ORION CORPORATION	FI
Etoricoxib Pensa 120 mg compresse rivestite con film	PT/H/1714/002-004	45220098	PENSA PHARMA S.P.A.	IT
Etoricoxib Pensa 120 mg compresse rivestite con film	PT/H/1714/002-004	45220086	PENSA PHARMA S.P.A.	IT
Etoricoxib Pensa 120 mg compresse rivestite con film	PT/H/1714/002-004	45220074	PENSA PHARMA S.P.A.	IT
Etoricoxib pensa 120 mg comprimidos recubiertos con película EFG	PT/H/1714/004	81.754	PENSA PHARMA, S.A	ES
Etoricoxib pensa 30 mg comprimidos recubiertos con película EFG	PT/H/1714/001	81.755	PENSA PHARMA, S.A	ES
Etoricoxib Pensa 60 mg compresse rivestite con film	PT/H/1714/002-004	45220011	PENSA PHARMA S.P.A.	IT
Etoricoxib Pensa 60 mg compresse rivestite con film	PT/H/1714/002-004	45220023	PENSA PHARMA S.P.A.	IT
Etoricoxib Pensa 60 mg compresse rivestite con film	PT/H/1714/002-004	45220035	PENSA PHARMA S.P.A.	IT
Etoricoxib pensa 60 mg comprimidos recubiertos con película EFG	PT/H/1714/002	81.756	PENSA PHARMA, S.A	ES
Etoricoxib Pensa 90 mg compresse rivestite con film	PT/H/1714/002-004	45220050	PENSA PHARMA 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
Etoricoxib Pensa 90 mg compresse rivestite con film	PT/H/1714/002-004	45220062	PENSA PHARMA S.P.A.	IT
Etoricoxib Pensa 90 mg compresse rivestite con film	PT/H/1714/002-004	45220047	PENSA PHARMA S.P.A.	IT
Etoricoxib pensa 90 mg comprimidos recubiertos con película EFG	PT/H/1714/003	81.757	PENSA PHARMA, S.A	ES
Etoricoxib Pentafarma 120 mg comprimidos revestidos por película	not available	5693700	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 120 mg comprimidos revestidos por película	not available	5693718	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 120 mg comprimidos revestidos por película	not available	16/H/0029/004	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 120 mg comprimidos revestidos por película	not available	16/H/0029/004	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 30 mg comprimidos revestidos por película	not available	5693627	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 30 mg comprimidos revestidos por película	not available	5693635	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 30 mg comprimidos revestidos por película	not available	16/H/0029/001	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL 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
Etoricoxib Pentafarma 30 mg comprimidos revestidos por película	not available	16/H/0029/001	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 60 mg comprimidos revestidos por película	not available	5693643	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 60 mg comprimidos revestidos por película	not available	5693650	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 60 mg comprimidos revestidos por película	not available	16/H/0029/002	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 60 mg comprimidos revestidos por película	not available	16/H/0029/002	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 90 mg comprimidos revestidos por película	not available	5693668	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 90 mg comprimidos revestidos por película	not available	5693676	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 90 mg comprimidos revestidos por película	not available	16/H/0029/003	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib Pentafarma 90 mg comprimidos revestidos por película	not available	16/H/0029/003	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/07	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/08	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/09	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/10	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/11	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/12	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/13	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/14	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/15	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 120 mg comprimate filmate	NL/H/3267/003	8575/2016/17	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/07	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/08	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/09	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/10	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/11	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/12	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/13	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/14	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/15	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 60 mg comprimate filmate	NL/H/3267/001	8573/2016/17	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/07	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/08	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/09	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/10	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/11	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/12	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/13	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/14	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/15	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharmaSwiss 90 mg comprimate filmate	NL/H/3267/002	8574/2016/17	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Etoricoxib PharOS	PT/H/1374/001	PT/H/1374/001	PHAROS - PHARMACEUTICAL ORIENTED SERVICES LIMITED	PT
Etoricoxib PharOS	PT/H/1374/002	PT/H/1374/002	PHAROS - PHARMACEUTICAL ORIENTED SERVICES LIMITED	PT
Etoricoxib PharOS	PT/H/1374/003	PT/H/1374/003	PHAROS - PHARMACEUTICAL ORIENTED SERVICES LIMITED	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
Etoricoxib PharOS	PT/H/1374/004	PT/H/1374/004	PHAROS - PHARMACEUTICAL ORIENTED SERVICES LIMITED	PT
Etoricoxib PUREN 120 mg Filmtabletten	PT/H/1603/004	97036.00.00	PUREN PHARMA GMBH & CO. KG	DE
Etoricoxib PUREN 30 mg Filmtabletten	PT/H/1603/001	97033.00.00	PUREN PHARMA GMBH & CO. KG	DE
Etoricoxib PUREN 60 mg Filmtabletten	PT/H/1603/002	97034.00.00	PUREN PHARMA GMBH & CO. KG	DE
Etoricoxib PUREN 90 mg Filmtabletten	PT/H/1603/003	97035.00.00	PUREN PHARMA GMBH & CO. KG	DE
Etoricoxib ratiopharm 120 mg comprimidos recubiertos con película EFG	DE/H/5032/004	80964	RATIOPHARM ESPANA SA	ES
Etoricoxib ratiopharm 120 mg comprimidos revestidos por película	DE/H/5032/004	5674650	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 120 mg comprimidos revestidos por película	DE/H/5032/004	5674643	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 120 mg filmdragerade tabletter	DE/H/5032/004	32500	RATIOPHARM GMBH	FI
Etoricoxib ratiopharm 120 mg filmuhúðaðar töflur.	DE/H/5032/004	IS/1/16/102/04	RATIOPHARM GMBH	IS
Etoricoxib ratiopharm 120 mg tabletti, kalvopäällysteinen	DE/H/5032/004	32500	RATIOPHARM GMBH	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
Etoricoxib ratiopharm 30 mg comprimidos recubiertos con película EFG	DE/H/5032/001	80961	RATIOPHARM ESPANA SA	ES
Etoricoxib ratiopharm 30 mg comprimidos revestidos por película	DE/H/5032/001	5674601	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 30 mg comprimidos revestidos por película	DE/H/5032/001	5674577	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 30 mg filmdragerade tabletter	DE/H/5032/001	32497	RATIOPHARM GMBH	FI
Etoricoxib ratiopharm 30 mg filmuhúðaðar töflur.	DE/H/5032/001	IS/1/16/102/01	RATIOPHARM GMBH	IS
Etoricoxib ratiopharm 30 mg tabletti, kalvopäällysteinen	DE/H/5032/001	32497	RATIOPHARM GMBH	FI
Etoricoxib ratiopharm 60 mg comprimidos recubiertos con película EFG	DE/H/5032/002	80962	RATIOPHARM ESPANA SA	ES
Etoricoxib ratiopharm 60 mg comprimidos revestidos por película	DE/H/5032/002	5674619	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 60 mg comprimidos revestidos por película	DE/H/5032/002	5674627	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 60 mg comprimidos revestidos por película	DE/H/5032/002	5687157	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 60 mg filmdragerade tabletter	DE/H/5032/002	32498	RATIOPHARM GMBH	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
Etoricoxib ratiopharm 60 mg filmuhúðaðar töflur.	DE/H/5032/002	IS/1/16/102/02	RATIOPHARM GMBH	IS
Etoricoxib ratiopharm 60 mg tabletti, kalvopäällysteinen	DE/H/5032/002	32498	RATIOPHARM GMBH	FI
Etoricoxib ratiopharm 90 mg comprimidos recubiertos con película EFG	DE/H/5032/003	80963	RATIOPHARM ESPANA SA	ES
Etoricoxib ratiopharm 90 mg comprimidos revestidos por película	DE/H/5032/003	5674809	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 90 mg comprimidos revestidos por película	DE/H/5032/003	5674635	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 90 mg comprimidos revestidos por película	DE/H/5032/003	5687165	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib ratiopharm 90 mg filmdragerade tabletter	DE/H/5032/003	32499	RATIOPHARM GMBH	FI
Etoricoxib ratiopharm 90 mg filmuhúðaðar töflur.	DE/H/5032/003	IS/1/16/102/03	RATIOPHARM GMBH	IS
Etoricoxib ratiopharm 90 mg tabletti, kalvopäällysteinen	DE/H/5032/003	32499	RATIOPHARM GMBH	FI
Etoricoxib Rontis 120mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100427/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 120mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100427/14	RONTIS HELLAS SA	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
Etoricoxib Rontis 30mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100431/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 30mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100431/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 60mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100429/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 60mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100429/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 90mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100428/14	RONTIS HELLAS SA	GR
Etoricoxib Rontis 90mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/1375/001-004/DC	100428/14	RONTIS HELLAS SA	GR
ETORICOXIB SANDOZ	DE/H/3909/002	043004112	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004124	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004136	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004148	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004151	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004163	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004175	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004187	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004199	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/002	043004201	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004213	SANDOZ 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
ETORICOXIB SANDOZ	DE/H/3909/003	043004225	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004237	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004249	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004252	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004264	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004276	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004288	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004290	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/003	043004302	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004314	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004326	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004338	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004340	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004353	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004365	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004377	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004389	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004391	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/004	043004403	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004011	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004023	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004035	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004047	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004050	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004062	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004074	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004086	SANDOZ S.P.A.	IT
ETORICOXIB SANDOZ	DE/H/3909/001	043004098	SANDOZ 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
ETORICOXIB SANDOZ	DE/H/3909/001	043004100	SANDOZ S.P.A.	IT
Etoricoxib Sandoz	DE/H/3908/001	NONE	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/002	5659651	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/002	5659669	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/002	5659677	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/003	5659610	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/003	5659628	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/003	5659636	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/003	5659644	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/004	5659578	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/004	5659602	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz	DE/H/3908/002	5659701	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz 120 mg – Filmtabletten	DE/H/3908/004	136589	SANDOZ GMBH	AT
Etoricoxib Sandoz 120 mg comprimés pelliculés	DE/H/3908/004	2015120187	SANDOZ 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
Etoricoxib Sandoz 120 mg comprimidos recubiertos con película EFG	DE/H/3908/004	80263	SANDOZ FARMACÉUTICA, S.A.	ES
Etoricoxib Sandoz 120 mg filmdragerade tabletter	DE/H/3908/004	49819	SANDOZ A/S	SE
Etoricoxib Sandoz 120 mg filmdrasjerte tabletter	DE/H/3908/004	16/11062	SANDOZ A/S	NO
Etoricoxib Sandoz 120 mg filmomhulde tabletten	DE/H/3908/004	BE472142	SANDOZ N.V.	BE
Etoricoxib Sandoz 120 mg filmomhulde tabletten	DE/H/3908/004	BE472186	SANDOZ N.V.	BE
Etoricoxib Sandoz 120 mg tabletti, kalvopäällysteinen	DE/H/3908/004	31733	SANDOZ A/S	FI
Etoricoxib Sandoz 120 mg tabletti, kalvopäällysteinen	DE/H/3908/004	31733	SANDOZ A/S	FI
Etoricoxib Sandoz 120 mg, filmomhulde tabletten	DE/H/3908/004	RVG 114111	SANDOZ B.V.	NL
Etoricoxib Sandoz 30 mg – Filmtabletten	DE/H/3908/001	136586	SANDOZ GMBH	AT
Etoricoxib Sandoz 30 mg comprimés pelliculés	DE/H/3908/001	2015120184	SANDOZ N.V.	LU
Etoricoxib Sandoz 30 mg comprimidos recubiertos con película EFG	DE/H/3908/001	80265	SANDOZ FARMACÉUTICA, S.A.	ES
Etoricoxib Sandoz 30 mg filmdragerade tabletter	DE/H/3908/001	49816	SANDOZ A/S	SE
Etoricoxib Sandoz 30 mg filmdrasjerte tabletter	DE/H/3908/001	16/11059	SANDOZ A/S	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
Etoricoxib Sandoz 30 mg filmomhulde tabletten	DE/H/3908/001	BE472115	SANDOZ N.V.	BE
Etoricoxib Sandoz 30 mg filmomhulde tabletten	DE/H/3908/001	BE472151	SANDOZ N.V.	BE
Etoricoxib Sandoz 30 mg filmomhulde tabletten	DE/H/3908/001	RVG 114108	SANDOZ B.V.	NL
Etoricoxib Sandoz 30 mg tabletti, kalvopäällysteinen	DE/H/3908/001	31730	SANDOZ A/S	FI
Etoricoxib Sandoz 30 mg, filmom obalené tablety	DE/H/3908/001	29/0324/16-S	SANDOZ PHARMACEUTICALS D.D.	SK
Etoricoxib Sandoz 60 mg – Filmtabletten	DE/H/3908/002	136587	SANDOZ GMBH	AT
Etoricoxib Sandoz 60 mg comprimés pelliculés	DE/H/3908/002	2015120185	SANDOZ N.V.	LU
Etoricoxib Sandoz 60 mg comprimidos recubiertos con película EFG	DE/H/3908/002	80262	SANDOZ FARMACÉUTICA, S.A.	ES
Etoricoxib Sandoz 60 mg filmdragerade tabletter	DE/H/3908/002	49817	SANDOZ A/S	SE
Etoricoxib Sandoz 60 mg filmdrasjerte tabletter	DE/H/3908/002	16/11060	SANDOZ A/S	NO
Etoricoxib Sandoz 60 mg filmomhulde tabletten	DE/H/3908/002	BE472160	SANDOZ N.V.	BE
Etoricoxib Sandoz 60 mg filmomhulde tabletten	DE/H/3908/002	BE472124	SANDOZ N.V.	BE
Etoricoxib Sandoz 60 mg tabletti, kalvopäällysteinen	DE/H/3908/002	31731	SANDOZ A/S	FI
Etoricoxib Sandoz 60 mg, filmomhulde tabletten	DE/H/3908/002	RVG 114109	SANDOZ B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Etoricoxib Sandoz 90 mg – Filmtabletten	DE/H/3908/003	136588	SANDOZ GMBH	AT
Etoricoxib Sandoz 90 mg comprimés pelliculés	DE/H/3908/003	2015120186	SANDOZ N.V.	LU
Etoricoxib Sandoz 90 mg comprimidos recubiertos con película EFG	DE/H/3908/003	80264	SANDOZ FARMACÉUTICA, S.A.	ES
Etoricoxib Sandoz 90 mg filmdragerade tabletter	DE/H/3908/003	49818	SANDOZ A/S	SE
Etoricoxib Sandoz 90 mg filmdrasjerte tabletter	DE/H/3908/003	16/11061	SANDOZ A/S	NO
Etoricoxib Sandoz 90 mg filmomhulde tabletten	DE/H/3908/003	BE472177	SANDOZ N.V.	BE
Etoricoxib Sandoz 90 mg filmomhulde tabletten	DE/H/3908/003	BE472133	SANDOZ N.V.	BE
Etoricoxib Sandoz 90 mg tabletti, kalvopäällysteinen	DE/H/3908/003	31732	SANDOZ A/S	FI
Etoricoxib Sandoz 90 mg, filmomhulde tabletten	DE/H/3908/003	RVG 114110	SANDOZ B.V.	NL
Etoricoxib Sandoz Farmacêutica	not available	5694047	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5694054	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5693940	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5693957	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5694021	SANDOZ FARMACÊUTICA 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
Etoricoxib Sandoz Farmacêutica	not available	5694039	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5693965	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sandoz Farmacêutica	not available	5693973	SANDOZ FARMACÊUTICA LDA.	PT
Etoricoxib Sigillata 120 mg film-coated tablets	SE/H/1576/04	53672	SIGILLATA LIMITED	SE
Etoricoxib Sigillata 30 mg film-coated tablets	SE/H/1576/01	53669	SIGILLATA LIMITED	SE
Etoricoxib Sigillata 60 mg film-coated tablets	SE/H/1576/02	53670	SIGILLATA LIMITED	SE
Etoricoxib Sigillata 90 mg film-coated tablets	SE/H/1576/03	53671	SIGILLATA LIMITED	SE
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/06	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/01	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/07	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/02	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/03	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/04	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 120 mg comprimate filmate	DE/H/4251/004	9357/2016/05	STADA ARZNEIMITTEL AG	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
Etoricoxib STADA 120 mg comprimidos recubiertos con película EFG	DE/H/4251/004	81.220	LABORATORIO STADA, S.L.	ES
Etoricoxib STADA 120 mg filmdragerade tabletter	DE/H/4251/004	51986	STADA ARZNEIMITTEL AG	SE
Etoricoxib STADA 120 mg Filmtabletten	DE/H/4692/004	97254.00.00	STADAPHARM GMBH	DE
Etoricoxib STADA 120 mg kalvopäällysteiset tabletit	DE/H/4251/004	32781	STADA ARZNEIMITTEL AG	FI
Etoricoxib STADA 30 mg comprimate filmate	DE/H/4251/001	9354/2016/02	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 30 mg comprimate filmate	DE/H/4251/001	9354/2016/01	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 30 mg comprimate filmate	DE/H/4251/001	9354/2016/03	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 30 mg comprimate filmate	DE/H/4251/001	9354/2016/04	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 30 mg comprimidos recubiertos con película EFG	DE/H/4251/001	81.217	LABORATORIO STADA, S.L.	ES
Etoricoxib STADA 30 mg filmdragerade tabletter	DE/H/4251/001	51983	STADA ARZNEIMITTEL AG	SE
Etoricoxib STADA 30 mg Filmtabletten	DE/H/4692/001	97251.00.00	STADAPHARM GMBH	DE
Etoricoxib STADA 30 mg kalvopäällysteiset tabletit	DE/H/4251/001	32778	STADA ARZNEIMITTEL AG	FI
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/01	STADA ARZNEIMITTEL AG	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
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/06	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/07	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/08	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/04	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/03	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/10	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/09	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/05	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimate filmate	DE/H/4251/002	9355/2016/02	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 60 mg comprimidos recubiertos con película EFG	DE/H/4251/002	81.218	LABORATORIO STADA, S.L.	ES
Etoricoxib STADA 60 mg filmdragerade tabletter	DE/H/4251/002	51984	STADA ARZNEIMITTEL AG	SE
Etoricoxib STADA 60 mg Filmtabletten	DE/H/4692/002	97252.00.00	STADAPHARM GMBH	DE
Etoricoxib STADA 60 mg kalvopäällysteiset tabletit	DE/H/4251/002	32779	STADA ARZNEIMITTEL AG	FI
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/04	STADA ARZNEIMITTEL AG	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
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/08	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/06	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/02	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/09	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/01	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/10	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/07	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/03	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimate filmate	DE/H/4251/003	9356/2016/05	STADA ARZNEIMITTEL AG	RO
Etoricoxib STADA 90 mg comprimidos recubiertos con película EFG	DE/H/4251/003	81.219	LABORATORIO STADA, S.L.	ES
Etoricoxib STADA 90 mg filmdragerade tabletter	DE/H/4251/003	51985	STADA ARZNEIMITTEL AG	SE
Etoricoxib STADA 90 mg Filmtabletten	DE/H/4692/003	97253.00.00	STADAPHARM GMBH	DE
Etoricoxib STADA 90 mg kalvopäällysteiset tabletit	DE/H/4251/003	32780	STADA ARZNEIMITTEL AG	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
Etoricoxib Synthon 120 mg comprimidos recubiertos con película EFG	NL/H/3269/004	80672	SYNTHON BV	ES
Etoricoxib Synthon 30 mg comprimidos recubiertos con película EFG	NL/H/3269/001	80673	SYNTHON BV	ES
Etoricoxib Synthon 60 mg comprimidos recubiertos con película EFG	NL/H/3269/002	80674	SYNTHON BV	ES
Etoricoxib Synthon 90 mg comprimidos recubiertos con película EFG	NL/H/3269/003	80675	SYNTHON BV	ES
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/01	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/02	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/03	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/04	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/05	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/06	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/07	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/08	TERAPIA S.A.	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
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/09	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/10	TERAPIA S.A.	RO
Etoricoxib Terapia 120 mg comprimate filmate	DE/H/4557/004	9919/2017/11	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/01	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/02	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/03	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/04	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/05	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/06	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/07	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/08	TERAPIA S.A.	RO
Etoricoxib Terapia 30 mg comprimate filmate	DE/H/4557/001	9916/2017/09	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/01	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/02	TERAPIA S.A.	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
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/03	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/04	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/05	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/06	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/07	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/08	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/09	TERAPIA S.A.	RO
Etoricoxib Terapia 60 mg comprimate filmate	DE/H/4557/002	9917/2017/10	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/01	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/02	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/03	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/04	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/05	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/06	TERAPIA S.A.	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
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/07	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/08	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/09	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/10	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/11	TERAPIA S.A.	RO
Etoricoxib Terapia 90 mg comprimate filmate	DE/H/4557/003	9918/2017/12	TERAPIA S.A.	RO
Etoricoxib Teva 120 mg apvalkotās tabletes	DE/H/5031/004	16-0063	TEVA B.V	LV
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684226	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684253	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684339	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684240	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684238	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684341	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684277	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
ETORICOXIB TEVA 120 mg compresse rivestite con film	DE/H/5031/004	043684265	TEVA ITALIA S.R.L.	IT
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/01	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/008	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/004	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/07	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/02	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/05	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/06	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimate filmate	DE/H/5031/004	8805/2016/03	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 120 mg comprimidos recubiertos con película EFG	DE/H/5031/004	80948	TEVA PHARMA S.L.U	ES
Etoricoxib Teva 120 mg comprimidos revestidos por película	DE/H/5031/004	5674007	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 120 mg comprimidos revestidos por película	DE/H/5031/004	5673975	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 120 mg Film-coated Tablets	DE/H/5031/004	PA1986/002/004	TEVA B.V	IE

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Etoricoxib Teva 120 mg Film-coated Tablets	DE/H/5031/004	MA713/00304	TEVA B.V	MT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/030	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/029	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/033	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/028	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/031	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/034	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/027	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/032	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/032	TEVA B.V	LT
Etoricoxib Teva 120 mg plėvele dengtos tabletės	DE/H/5031/004	LT/1/16/3888/035	TEVA B.V	LT
Etoricoxib Teva 120 mg, filmomhulde tabletten	DE/H/5031/004	RVG 115906	TEVA NEDERLAND B.V.	NL
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684291	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684289	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
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684036	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684012	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684048	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 30 mg compresse rivestite con film	DE/H/5031/001	043684024	TEVA ITALIA S.R.L.	IT
Etoricoxib Teva 30 mg comprimate filmate	DE/H/5031/001	8802/2016/01-06	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 30 mg comprimidos recubiertos con película EFG	DE/H/5031/001	80944	TEVA PHARMA S.L.U	ES
Etoricoxib Teva 30 mg comprimidos revestidos por película	DE/H/5031/001	5673942	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 30 mg comprimidos revestidos por película	DE/H/5031/001	5673934	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 30 mg Film-coated Tablets	DE/H/5031/001	PA1986/002/001	TEVA B.V	IE
Etoricoxib Teva 30 mg Film-coated Tablets	DE/H/5031/001	MA713/00301	TEVA B.V	MT
Etoricoxib Teva 30 mg filmdragerade tabletter	DE/H/5031/001	51529	TEVA SWEDEN AB	SE
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/005	TEVA B.V	LT
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/004	TEVA B.V	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
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/003	TEVA B.V	LT
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/002	TEVA B.V	LT
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/001	TEVA B.V	LT
Etoricoxib Teva 30 mg plėvele dengtos tabletės	DE/H/5031/001	LT/1/16/3888/006	TEVA B.V	LT
Etoricoxib Teva 30 mg, filmomhulde tabletten	DE/H/5031/001	RVG 115901	TEVA NEDERLAND B.V.	NL
Etoricoxib Teva 60 mg apvalkotas tabletes	DE/H/5031/002	16-0061	TEVA B.V	LV
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684087	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684099	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684063	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684051	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684075	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684303	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684101	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684113	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
ETORICOXIB TEVA 60 mg compresse rivestite con film	DE/H/5031/002	043684125	TEVA ITALIA S.R.L.	IT
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/001	8803/2016/01	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/001	8803/2016/006	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/001	8803/2016/004	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/002	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/008	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/003	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/009	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/07	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimate filmate	DE/H/5031/002	8803/2016/05	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 60 mg comprimidos recubiertos con película EFG	DE/H/5031/001	80946	TEVA PHARMA S.L.U	ES
Etoricoxib Teva 60 mg comprimidos revestidos por película	DE/H/5031/002	5673959	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 60 mg comprimidos revestidos por película	DE/H/5031/002	5673967	TEVA PHARMA – PRODUTOS FARMACÊUTICOS 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
Etoricoxib Teva 60 mg comprimidos revestidos por película	DE/H/5031/002	5687132	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 60 mg Film-coated Tablets	DE/H/5031/002	PA1986/002/002	TEVA B.V	IE
Etoricoxib Teva 60 mg Film-coated Tablets	DE/H/5031/002	MA713/00302	TEVA B.V	MT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/015	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/013	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/014	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/007	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/012	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/008	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/009	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/011	TEVA B.V	LT
Etoricoxib Teva 60 mg plėvele dengtos tabletės	DE/H/5031/002	LT/1/16/3888/010	TEVA B.V	LT
Etoricoxib Teva 60 mg, filmomhulde tabletten	DE/H/5031/002	RVG 115904	TEVA NEDERLAND B.V.	NL
Etoricoxib Teva 90 mg apvalkotās tabletes	DE/H/5031/003	16-0062	TEVA B.V	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
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684214	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684315	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684176	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684190	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684188	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684152	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684327	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684202	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684164	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684137	TEVA ITALIA S.R.L.	IT
ETORICOXIB TEVA 90 mg compresse rivestite con film	DE/H/5031/003	043684149	TEVA ITALIA S.R.L.	IT
Etoricoxib Teva 90 mg comprimate filmate	DE/H/5031/003	8804/2016/01-11	TEVA PHARMACEUTICALS S.R.L	RO
Etoricoxib Teva 90 mg comprimidos recubiertos con película EFG	DE/H/5031/003	80949	TEVA PHARMA S.L.U	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
Etoricoxib Teva 90 mg comprimidos revestidos por película	DE/H/5031/003	5674353	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 90 mg comprimidos revestidos por película	DE/H/5031/003	5674361	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 90 mg comprimidos revestidos por película	DE/H/5031/003	5687140	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Etoricoxib Teva 90 mg Film-coated Tablets	DE/H/5031/003	PA1986/002/003	TEVA B.V	IE
Etoricoxib Teva 90 mg Film-coated Tablets	DE/H/5031/003	MA713/00303	TEVA B.V	MT
Etoricoxib Teva 90 mg filmdragerade tabletter	DE/H/5031/003	51531	TEVA SWEDEN AB	SE
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/025	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/021	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/020	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/022	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/016	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/026	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/024	TEVA B.V	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
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/018	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/023	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/019	TEVA B.V	LT
Etoricoxib Teva 90 mg plėvele dengtos tabletės	DE/H/5031/003	LT/1/16/3888/017	TEVA B.V	LT
Etoricoxib Teva 90 mg, filmomhulde tabletten	DE/H/5031/003	RVG 115905	TEVA NEDERLAND B.V.	NL
Etoricoxib Teva, 120 mg filmdragerad tablett	DE/H/5031/004	51532	TEVA SWEDEN AB	SE
Etoricoxib Teva, 120 mg õhukese polümeerikattega tabletid	DE/H/5031/004	903916	TEVA B.V	EE
Etoricoxib Teva, 120 mg, tabletki powlekane	SE/H/1574/004	23645	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Etoricoxib Teva, 30 mg õhukese polümeerikattega tabletid	DE/H/5031/001	904116	TEVA B.V	EE
Etoricoxib Teva, 30 mg, tabletki powlekane	SE/H/1574/001	23642	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Etoricoxib Teva, 60 mg filmdragerad tablett	DE/H/5031/002	51530	TEVA SWEDEN AB	SE
Etoricoxib Teva, 60 mg õhukese polümeerikattega tabletid	DE/H/5031/002	904016	TEVA B.V	EE
Etoricoxib Teva, 60 mg, tabletki powlekane	SE/H/1574/002	23643	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Etoricoxib Teva, 90 mg õhukese polümeerikattega tabletid	DE/H/5031/003	904216	TEVA B.V	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
Etoricoxib Teva, 90 mg, tabletki powlekane	SE/H/1574/003	23644	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Etoricoxib toLife 120 mg comprimidos revestidos por película	PT/H/1714/004	5704614	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 120 mg comprimidos revestidos por película	PT/H/1714/004	5704606	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 30 mg comprimidos revestidos por película	PT/H/1714/001	5704507	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 30 mg comprimidos revestidos por película	PT/H/1714/001	5704515	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 30 mg comprimidos revestidos por película	PT/H/1714/001	5704473	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 60 mg comprimidos revestidos por película	PT/H/1714/002	5704523	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 60 mg comprimidos revestidos por película	PT/H/1714/002	5704549	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 60 mg comprimidos revestidos por película	PT/H/1714/002	5704531	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 90 mg comprimidos revestidos por película	PT/H/1714/003	5704556	TOLIFE - 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
Etoricoxib toLife 90 mg comprimidos revestidos por película	PT/H/1714/003	5704572	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib toLife 90 mg comprimidos revestidos por película	PT/H/1714/003	5704564	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg apvalkotās tabletes	EE/H/0211/004	16-0083	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747207	ZENTIVA ITALIA SRL	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
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747195	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747169	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747183	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747171	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747207	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747195	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747169	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747183	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compresse rivestite con film	EE/H/0211/004	043747171	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 120 mg compridos revestidos por película	EE/H/0211/004	5692447	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 120 mg compridos revestidos por película	EE/H/0211/004	5694831	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 120 mg compridos revestidos por película	EE/H/0211/004	5692454	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 120 mg compridos revestidos por película	EE/H/0211/004	5692447	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 120 mg compridos revestidos por película	EE/H/0211/004	5694831	SANOFI - PRODUTOS FARMACEUTICOS 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
Etoricoxib Zentiva 120 mg comprimidos revestidos por película	EE/H/0211/004	5692454	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg film-coated tablets	EE/H/0211/004	MA082/08504	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA 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
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg Filmtabletten	EE/H/0211/004	92974.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/018	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/019	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/017	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/016	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/020	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/018	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/019	ZENTIVA, K.S.	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
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/017	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/016	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg plėvele dengtos tabletės	EE/H/0211/004	LT/1/16/3907/020	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 120 mg, õhukese polümeerikattega tabletid	EE/H/0211/004	906916	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747029	ZENTIVA ITALIA SRL	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
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747031	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747017	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747029	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747031	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 30 mg compresse rivestite con film	EE/H/0211/001	043747017	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 30 mg compridos revestidos por película	EE/H/0211/001	5692413	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 30 mg compridos revestidos por película	EE/H/0211/001	5692405	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 30 mg compridos revestidos por película	EE/H/0211/001	5692413	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 30 mg compridos revestidos por película	EE/H/0211/001	5692405	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	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
Etoricoxib Zentiva 30 mg film-coated tablets	EE/H/0211/001	MA082/08501	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg Filmtabletten	EE/H/0211/001	92971.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/003	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/001	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/002	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/003	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/001	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg plėvele dengtos tabletės	EE/H/0211/001	LT/1/16/3907/002	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	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
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 30 mg, õhukese polümeerikattega tabletid	EE/H/0211/001	907216	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	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
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg apvalkotās tabletes	EE/H/0211/002	16-0081	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747043	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747082	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747056	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747094	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747068	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747070	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747043	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747082	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747056	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747094	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747068	ZENTIVA ITALIA SRL	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
Etoricoxib Zentiva 60 mg compresse rivestite con film	EE/H/0211/002	043747070	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5692371	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5694807	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5692363	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5692371	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5694807	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg compridos revestidos por película	EE/H/0211/002	5692363	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	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
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg film-coated tablets	EE/H/0211/002	MA082/08502	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA 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
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg Filmtabletten	EE/H/0211/002	92972.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/008	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/004	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/005	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/006	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/007	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/009	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/008	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/004	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/005	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/006	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/007	ZENTIVA, K.S.	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
Etoricoxib Zentiva 60 mg plėvele dengtos tabletės	EE/H/0211/002	LT/1/16/3907/009	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 60 mg, õhukese polümeerikattega tabletid	EE/H/0211/002	907116	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	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
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg apvalkotās tabletes	EE/H/0211/003	16-0082	ZENTIVA, K.S.	LV
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747120	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747157	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747132	ZENTIVA ITALIA SRL	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
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747106	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747144	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747118	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747120	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747157	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747132	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747106	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747144	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compresse rivestite con film	EE/H/0211/003	043747118	ZENTIVA ITALIA SRL	IT
Etoricoxib Zentiva 90 mg compridos revestidos por película	EE/H/0211/003	5694823	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 90 mg compridos revestidos por película	EE/H/0211/003	5692439	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 90 mg compridos revestidos por película	EE/H/0211/003	5692421	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 90 mg compridos revestidos por película	EE/H/0211/003	5694823	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 90 mg compridos revestidos por película	EE/H/0211/003	5692439	SANOFI - PRODUTOS FARMACEUTICOS 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
Etoricoxib Zentiva 90 mg comprimidos revestidos por película	EE/H/0211/003	5692421	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg film-coated tablets	EE/H/0211/003	MA082/08503	SANOFI MALTA LTD	MT
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA 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
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg Filmtabletten	EE/H/0211/003	92973.00.00	ZENTIVA PHARMA GMBH	DE
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/015	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/013	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/011	ZENTIVA, K.S.	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
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/012	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/014	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/010	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/015	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/013	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/011	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/012	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/014	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg plėvele dengtos tabletės	EE/H/0211/003	LT/1/16/3907/010	ZENTIVA, K.S.	LT
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	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
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zentiva 90 mg, õhukese polümeerikattega tabletid	EE/H/0211/003	907016	ZENTIVA, K.S.	EE
Etoricoxib Zydus 120 mg comprimés pelliculés	DE/H/3469/004	BE445304	ZYDUS FRANCE	BE
Etoricoxib Zydus 120 mg comprimidos recubiertos con película EFG	DE/H/3469/004	79986	ZYDUS FRANCE	ES
Etoricoxib Zydus 120 mg comprimidos revestidos por película	DE/H/3469/004	5633847	ZYDUS FRANCE	PT
Etoricoxib Zydus 120 mg filmomhulde tabletten	DE/H/3469/004	RVG 111531	ZYDUS FRANCE	NL
Etoricoxib Zydus 120 mg filmtabletta	DE/H/3469/004	OGYI-T-225771/04	ZYDUS FRANCE	HU
Etoricoxib Zydus 120 mg Filmtabletten	DE/H/3469/004	86169.00.00	ZYDUS FRANCE	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
Etoricoxib Zydus 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3469/004	3018904	ZYDUS FRANCE	GR
Etoricoxib Zydus 30 mg comprimés pelliculés	DE/H/3469/001	BE445277	ZYDUS FRANCE	BE
Etoricoxib Zydus 30 mg comprimidos recubiertos con película EFG	DE/H/3469/001	79900	ZYDUS FRANCE	ES
Etoricoxib Zydus 30 mg comprimidos revestidos por película	DE/H/3469/001	5633821	ZYDUS FRANCE	PT
Etoricoxib Zydus 30 mg filmomhulde tabletten	DE/H/3469/001	RVG 111528	ZYDUS FRANCE	NL
Etoricoxib Zydus 30 mg filmtabletta	DE/H/3469/001	OGYI-T-225771/01	ZYDUS FRANCE	HU
Etoricoxib Zydus 30 mg Filmtabletten	DE/H/3469/001	86166.00.00	ZYDUS FRANCE	DE
Etoricoxib Zydus 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3469/001	3018901	ZYDUS FRANCE	GR
Etoricoxib Zydus 60 mg comprimés pelliculés	DE/H/3469/002	BE445286	ZYDUS FRANCE	BE
Etoricoxib Zydus 60 mg comprimidos recubiertos con película EFG	DE/H/3469/002	79901	ZYDUS FRANCE	ES
Etoricoxib Zydus 60 mg comprimidos revestidos por película	DE/H/3469/002	5633813	ZYDUS FRANCE	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
Etoricoxib Zydus 60 mg filmomhulde tabletten	DE/H/3469/002	RVG 111529	ZYDUS FRANCE	NL
Etoricoxib Zydus 60 mg filmtabletta	DE/H/3469/002	OGYI-T-225771/02	ZYDUS FRANCE	HU
Etoricoxib Zydus 60 mg Filmtabletten	DE/H/3469/002	86167.00.00	ZYDUS FRANCE	DE
Etoricoxib Zydus 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3469/002	3018902	ZYDUS FRANCE	GR
Etoricoxib Zydus 90 mg comprimés pelliculés	DE/H/3469/003	BE445295	ZYDUS FRANCE	BE
Etoricoxib Zydus 90 mg comprimidos recubiertos con película EFG	DE/H/3469/003	79902	ZYDUS FRANCE	ES
Etoricoxib Zydus 90 mg comprimidos revestidos por película	DE/H/3469/003	5633839	ZYDUS FRANCE	PT
Etoricoxib Zydus 90 mg filmomhulde tabletten	DE/H/3469/003	RVG 111530	ZYDUS FRANCE	NL
Etoricoxib Zydus 90 mg filmtabletta	DE/H/3469/003	OGYI-T-225771/03	ZYDUS FRANCE	HU
Etoricoxib Zydus 90 mg Filmtabletten	DE/H/3469/003	86168.00.00	ZYDUS FRANCE	DE
Etoricoxib Zydus 90 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3469/003	3018903	ZYDUS FRANCE	GR
Etoricoxib Zydus France 120 mg compresse rivestite con film	DE/H/3469/001	042146047	ZYDUS FRANCE	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
ETORICOXIB ZYDUS FRANCE 120 mg comprimé pelliculé	DE/H/3469/004	34009 300 202 4 1	ZYDUS FRANCE	FR
Etoricoxib Zydus France 30 mg compresse rivestite con film	DE/H/3469/001	042146011	ZYDUS FRANCE	IT
ETORICOXIB ZYDUS FRANCE 30 mg comprimé pelliculé	DE/H/3469/001	34009 300 197 9 5	ZYDUS FRANCE	FR
Etoricoxib Zydus France 60 mg compresse rivestite con film	DE/H/3469/001	042146023	ZYDUS FRANCE	IT
ETORICOXIB ZYDUS FRANCE 60 mg comprimé pelliculé	DE/H/3469/002	34009 300 202 2 7	ZYDUS FRANCE	FR
Etoricoxib Zydus France 90 mg compresse rivestite con film	DE/H/3469/001	042146035	ZYDUS FRANCE	IT
ETORICOXIB ZYDUS FRANCE 90 mg comprimé pelliculé	DE/H/3469/003	34009 300 202 3 4	ZYDUS FRANCE	FR
Etoricoxib Zydus, 120 mg, tabletki powlekane	DE/H/3469/004	22052	ZYDUS FRANCE	PL
Etoricoxib Zydus, 30 mg, tabletki powlekane	DE/H/3469/001	22049	ZYDUS FRANCE	PL
Etoricoxib Zydus, 60 mg, tabletki powlekane	DE/H/3469/002	22050	ZYDUS FRANCE	PL
Etoricoxib Zydus, 90, mg tabletki powlekane	DE/H/3469/003	22051	ZYDUS FRANCE	PL
Etoricoxib-ratiopharm 120 mg Filmtabletten	DE/H/5032/004	92420.00.00	RATIOPHARM GMBH	DE
Etoricoxib-ratiopharm 120 mg Filmtabletten	DE/H/5032/004	2016110295	RATIOPHARM GMBH	LU
Etoricoxib-ratiopharm 30 mg Filmtabletten	DE/H/5032/001	92417.00.00	RATIOPHARM 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
Etoricoxib-ratiopharm 30 mg Filmtabletten	DE/H/5032/001	2016110292	RATIOPHARM GMBH	LU
Etoricoxib-ratiopharm 60 mg Filmtabletten	DE/H/5032/002	92418.00.00	RATIOPHARM GMBH	DE
Etoricoxib-ratiopharm 60 mg Filmtabletten	DE/H/5032/002	2016110293	RATIOPHARM GMBH	LU
Etoricoxib-ratiopharm 90 mg Filmtabletten	DE/H/5032/003	92419.00.00	RATIOPHARM GMBH	DE
Etoricoxib-ratiopharm 90 mg Filmtabletten	DE/H/5032/003	2016110294	RATIOPHARM GMBH	LU
Etorikoksib Lek 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/073	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/022	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/021	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/026	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/023	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/027	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/024	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 120 mg filmsko obložene tablete	DE/H/4251/004	H/17/02292/025	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/004	STADA ARZNEIMITTEL AG	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
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/005	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/006	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/002	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/003	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/007	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/009	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/001	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/008	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 60 mg filmsko obložene tablete	DE/H/4251/002	H/17/02292/010	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/015	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/014	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/012	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/011	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/018	STADA ARZNEIMITTEL AG	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
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/016	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/013	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/020	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/019	STADA ARZNEIMITTEL AG	SI
Etorikoksib STADA 90 mg filmsko obložene tablete	DE/H/4251/003	H/17/02292/017	STADA ARZNEIMITTEL AG	SI
Etorilin	NL/H/3271/001	58341	SYNTHON BV	DK
Etorilin	NL/H/3271/004	58344	SYNTHON BV	DK
Etorilin	NL/H/3271/002	58342	SYNTHON BV	DK
Etorilin	NL/H/3271/003	58343	SYNTHON BV	DK
Etoxib 120 mg filmom obložene tablete	HU/H/0435/004	HR-H-399675011	KRKA-FARMA D.O.O.	HR
Etoxib 30 mg filmom obložene tablete	HU/H/0435/001	HR-H-568519667	KRKA-FARMA D.O.O.	HR
Etoxib 60 mg filmom obložene tablete	HU/H/0435/002	HR-H-352347859	KRKA-FARMA D.O.O.	HR
Etoxib 90 mg filmom obložene tablete	HU/H/0435/003	HR-H-589688667	KRKA-FARMA D.O.O.	HR
Etoxib, 120 mg õhukese polümeerikattega tabletid	HU/H/0435/004	921816	KRKA, D.D., NOVO MESTO	EE
Etoxib, 30 mg õhukese polümeerikattega tabletid	HU/H/0435/001	921516	KRKA, D.D., NOVO MESTO	EE
Etoxib, 60 mg õhukese polümeerikattega tabletid	HU/H/0435/002	921616	KRKA, D.D., NOVO MESTO	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
Etoxib, 90 mg õhukese polümeerikattega tabletid	HU/H/0435/003	921716	KRKA, D.D., NOVO MESTO	EE
Evetore 120 mg Filmtabletten	NL/H/3271/004	2016060161	SYNTHON BV	LU
Evetore 120 mg tabletten	NL/H/3271/004	RVG 116030	SYNTHON BV	NL
Evetore 30 mg Filmtabletten	NL/H/3271/001	2016060158	SYNTHON BV	LU
Evetore 30 mg tabletten	NL/H/3271/001	RVG 116027	SYNTHON BV	NL
Evetore 60 mg Filmtabletten	NL/H/3271/002	2016060159	SYNTHON BV	LU
Evetore 60 mg tabletten	NL/H/3271/002	RVG 116028	SYNTHON BV	NL
Evetore 90 mg Filmtabletten	NL/H/3271/003	2016060160	SYNTHON BV	LU
Evetore 90 mg tabletten	NL/H/3271/003	RVG 116029	SYNTHON BV	NL
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822295	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822307	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822319	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822321	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822333	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822345	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822358	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822360	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822372	ABIOGEN PHARMA 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
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822384	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822396	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822408	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822410	ABIOGEN PHARMA S.P.A.	IT
EXINEF 120 mg compresse rivestite con film	UK/H/0534/004	035822422	ABIOGEN PHARMA S.P.A.	IT
EXINEF 30 mg compresse rivestite con film	UK/H/0534/001	035822446	ABIOGEN PHARMA S.P.A.	IT
EXINEF 30 mg compresse rivestite con film	UK/H/0534/001	035822434	ABIOGEN PHARMA S.P.A.	IT
EXINEF 30 mg compresse rivestite con film	UK/H/0534/001	035822459	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822016	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822028	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822030	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822042	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822055	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822067	ABIOGEN PHARMA 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
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822079	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822081	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822093	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822105	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822117	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822129	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822131	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822143	ABIOGEN PHARMA S.P.A.	IT
EXINEF 60 mg compresse rivestite con film	UK/H/0534/002	035822461	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822156	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822168	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822170	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822182	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822194	ABIOGEN PHARMA 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
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822206	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822218	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822220	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822232	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822244	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822257	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822269	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822271	ABIOGEN PHARMA S.P.A.	IT
EXINEF 90 mg compresse rivestite con film	UK/H/0534/003	035822283	ABIOGEN PHARMA S.P.A.	IT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/003	4110284	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110383	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110482	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110581	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110680	BIAL - PORTELA & Cª , SA	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
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110789	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110888	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4110987	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111084	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4242285	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111183	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111282	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111589	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4242186	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111381	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos revestidos por película	UK/H/0534/004	4111480	BIAL - PORTELA & Cª , SA	PT
EXXIV 120 mg comprimidos recubiertos con película	UK/H/0534/004	64.933	LABORATORIOS ABELLÓ, S.A.	ES
EXXIV 30 mg comprimidos revestidos por película	UK/H/0534/004	5121678	BIAL - PORTELA & Cª , SA	PT
EXXIV 30 mg comprimidos revestidos por película	UK/H/0534/001	5063045	BIAL - PORTELA & Cª , SA	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
EXXIV 30 mg comprimidos recubiertos con película	UK/H/0534/001	72.076	LABORATORIOS ABELLÓ, S.A.	ES
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/001	4111688	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4111787	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4111886	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4111985	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112082	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112181	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112280	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112389	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112488	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4241881	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112587	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112686	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112983	BIAL - PORTELA & Cª , SA	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
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4241782	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112785	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos revestidos por película	UK/H/0534/002	4112884	BIAL - PORTELA & Cª , SA	PT
EXXIV 60 mg comprimidos recubiertos con película	UK/H/0534/002	64.931	LABORATORIOS ABELLÓ, S.A.	ES
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/002	4113080	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4118584	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113189	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113288	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113387	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113486	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113585	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113684	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113783	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4242087	BIAL - PORTELA & Cª , SA	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
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113882	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4113981	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4114286	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4241980	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4114088	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos revestidos por película	UK/H/0534/003	4114187	BIAL - PORTELA & Cª , SA	PT
EXXIV 90 mg comprimidos recubiertos con película	UK/H/0534/003	64.932	LABORATORIOS ABELLÓ, S.A.	ES
EXXIV®120 mg film-coated tablets	UK/H/0534/004	PL 0025/0430	MERCK SHARP & DOHME LTD.	UK
EXXIV®30 mg film-coated tablets	UK/H/0534/001	PL 0025/0480	MERCK SHARP & DOHME LTD.	UK
EXXIV®60 mg film-coated tablets	UK/H/0534/002	PL 0025/0428	MERCK SHARP & DOHME LTD.	UK
EXXIV®90 mg film-coated tablets	UK/H/0534/003	PL 0025/0429	MERCK SHARP & DOHME LTD.	UK
Еторикоксиб Актавис 120 mg филмирани таблетки	SE/H/1574/004	20170079	ACTAVIS GROUP PTC EHF.	BG
Еторикоксиб Актавис 30 mg филмирани таблетки	SE/H/1574/002	20170077	ACTAVIS GROUP PTC EHF.	BG
Еторикоксиб Актавис 30 mg филмирани таблетки	SE/H/1574/001	20170076	ACTAVIS GROUP PTC EHF.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Еторикоксиб Актавис 90 mg филмирани таблетки	SE/H/1574/003	20170078	ACTAVIS GROUP PTC EHF.	BG
Halliztolva 120 mg film-coated tablets	SE/H/1633/04	55108	SIGILLATA LIMITED	SE
Halliztolva 30 mg film-coated tablets	SE/H/1633/01	55105	SIGILLATA LIMITED	SE
Halliztolva 60 mg film-coated tablets	SE/H/1633/02	55106	SIGILLATA LIMITED	SE
Halliztolva 90 mg film-coated tablets	SE/H/1633/03	55107	SIGILLATA LIMITED	SE
IMESOL 120 mg comprimate filmate	RO/H/0157/004	8568/2016/01	TERAPIA S.A.	RO
IMESOL 120 mg comprimate filmate	RO/H/0157/004	8568/2016/02	TERAPIA S.A.	RO
IMESOL 30 mg comprimate filmate	RO/H/0157/001	8565/2016/01	TERAPIA S.A.	RO
IMESOL 30 mg comprimate filmate	RO/H/0157/001	8565/2016/02	TERAPIA S.A.	RO
IMESOL 60 mg comprimate filmate	RO/H/0157/002	8566/2016/01	TERAPIA S.A.	RO
IMESOL 60 mg comprimate filmate	RO/H/0157/002	8566/2016/02	TERAPIA S.A.	RO
IMESOL 90 mg comprimate filmate	RO/H/0157/003	8567/2016/01	TERAPIA S.A.	RO
IMESOL 90 mg comprimate filmate	RO/H/0157/003	8567/2016/02	TERAPIA S.A.	RO
Itoroxx 30 mg filmtabletta	HU/H/0444/001	OGYI-T-23101/01	KRKA, D.D., NOVO MESTO	HU
Itoroxx 30 mg filmtabletta	HU/H/0444/001	OGYI-T-23101/02	KRKA, D.D., NOVO MESTO	HU
Itoroxx 30 mg filmtabletta	HU/H/0444/001	OGYI-T-23101/03	KRKA, D.D., NOVO MESTO	HU
Itoroxx 30 mg filmtabletta	HU/H/0444/001	OGYI-T-23101/04	KRKA, D.D., NOVO MESTO	HU
Itoroxx 30 mg filmtabletta	HU/H/0444/001	OGYI-T-23101/05	KRKA, D.D., NOVO MESTO	HU
Itoroxx 60 mg filmtabletta	HU/H/0444/002	OGYI-T-23101/06	KRKA, D.D., NOVO MESTO	HU
Itoroxx 60 mg filmtabletta	HU/H/0444/002	OGYI-T-23101/07	KRKA, D.D., NOVO MESTO	HU
Itoroxx 60 mg filmtabletta	HU/H/0444/002	OGYI-T-23101/08	KRKA, D.D., NOVO MESTO	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
Itoroxx 60 mg filmtabletta	HU/H/0444/002	OGYI-T-23101/09	KRKA, D.D., NOVO MESTO	HU
Itoroxx 60 mg filmtabletta	HU/H/0444/002	OGYI-T-23101/10	KRKA, D.D., NOVO MESTO	HU
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/01	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/02	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/03	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/04	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/05	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/06	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/07	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/08	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/09	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/10	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/11	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/12	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/13	S.C. SANDOZ S.R.L.	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
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/14	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/15	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/16	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/17	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/18	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/19	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/20	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/21	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/22	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/23	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/24	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/25	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/26	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/27	S.C. SANDOZ S.R.L.	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
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/28	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/29	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/30	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/31	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg comprimate filmate	DE/H/3908/004	7797/2015/32	S.C. SANDOZ S.R.L.	RO
KOSTAROX 120 mg filmom obložene tablete	DE/H/3908/004	HR-H-677376885	SANDOZ D.O.O.	HR
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/097	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/098	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/099	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/100	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/101	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/102	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/103	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/104	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/105	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/106	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/107	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/108	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/109	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/110	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/111	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/112	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/113	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/114	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/115	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/116	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/117	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/118	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/119	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/120	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/121	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/122	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/123	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/124	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/125	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/126	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/127	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 120 mg filmsko obložene tablete	DE/H/3908/004	H/15/02067/128	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/01	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/02	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/04	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/05	S.C. SANDOZ S.R.L.	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
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/06	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/07	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/08	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/09	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/10	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/11	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/12	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/13	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/14	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/15	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/16	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/17	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/18	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/19	S.C. SANDOZ S.R.L.	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
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/20	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/21	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/22	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/23	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/24	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/25	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/26	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/27	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/28	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/29	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/30	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/31	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/32	S.C. SANDOZ S.R.L.	RO
KOSTAROX 30 mg comprimate filmate	DE/H/3908/001	7794/2015/03	S.C. SANDOZ S.R.L.	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
KOSTAROX 30 mg filmom obložene tablete	DE/H/3908/001	HR-H-144691129	SANDOZ D.O.O.	HR
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/028	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/030	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/029	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/025	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/027	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/031	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/026	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 30 mg filmsko obložene tablete	DE/H/3908/001	H/15/02067/032	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/01	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/02	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/03	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/04	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/05	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/06	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/07	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/08	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/10	S.C. SANDOZ S.R.L.	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
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/11	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/12	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/13	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/14	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/15	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/16	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/17	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/18	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/19	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/20	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/21	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/22	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/23	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/24	S.C. SANDOZ S.R.L.	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
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/25	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/26	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/27	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/28	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/29	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/30	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/31	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/32	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg comprimate filmate	DE/H/3908/002	7795/2015/09	S.C. SANDOZ S.R.L.	RO
KOSTAROX 60 mg filmom obložene tablete	DE/H/3908/002	HR-H-034074000	SANDOZ D.O.O.	HR
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/033	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/034	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/035	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/036	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/042	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/037	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/043	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/039	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/041	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/040	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/038	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/044	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/045	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/046	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/048	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/049	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/050	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/051	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/052	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/053	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/054	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/055	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/056	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/057	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/058	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/059	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/060	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/061	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/062	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/064	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/063	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 60 mg filmsko obložene tablete	DE/H/3908/002	H/15/02067/047	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/01	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/02	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/03	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/04	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/05	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/06	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/07	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/08	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/09	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/10	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/11	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/12	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/13	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/14	S.C. SANDOZ S.R.L.	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
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/15	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/16	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/17	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/18	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/19	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/20	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/21	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/22	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/23	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/24	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/25	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/26	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/27	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/28	S.C. SANDOZ S.R.L.	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
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/29	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/31	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/30	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg comprimate filmate	DE/H/3908/003	7796/2015/32	S.C. SANDOZ S.R.L.	RO
KOSTAROX 90 mg filmom obložene tablete	DE/H/3908/003	HR-H-031947675	SANDOZ D.O.O.	HR
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/065	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/066	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/067	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/068	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/069	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/070	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/071	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/072	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/073	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/074	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/075	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/076	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/077	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/078	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/079	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/080	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/081	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/082	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/083	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/084	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/085	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/086	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/087	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/088	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/089	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/090	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/091	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/092	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/093	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/094	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/095	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX 90 mg filmsko obložene tablete	DE/H/3908/003	H/15/02067/096	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
KOSTAROX, 30 mg, tabletki powlekane	DE/H/3908/001	23499	SANDOZ GMBH	PL
KOSTAROX, 60 mg, tabletki powlekane	DE/H/3908/002	23500	SANDOZ GMBH	PL
KOSTAROX, 90 mg, tabletki powlekane	DE/H/3908/003	23501	SANDOZ GMBH	PL
KOSTAROX,120 mg, tabletki powlekane	DE/H/3908/004	23502	SANDOZ GMBH	PL
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/25	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T -23054/26	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/27	SANDOZ 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
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/31	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/29	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/28	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T -23054/32	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 120 mg filmtabletta	DE/H/3908/004	OGYI-T-23054/30	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T -23054/07	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T-23054/08	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T-23054/04	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T -23054/05	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T-23054/01	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T-23054/02	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T -23054/03	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 30 mg filmtabletta	DE/H/3908/001	OGYI-T-23054/06	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T -23054/09	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T -23054/10	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T -23054/11	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T-23054/12	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T-23054/14	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T-23054/15	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T-23054/16	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 60 mg filmtabletta	DE/H/3908/002	OGYI-T-23054/13	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T -23054/17	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T -23054/18	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T -23054/19	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T -23054/21	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T-23054/22	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T-23054/23	SANDOZ HUNGÁRIA KFT	HU
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T-23054/24	SANDOZ 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
LINZAVO 90 mg filmtabletta	DE/H/3908/003	OGYI-T -23054/20	SANDOZ HUNGÁRIA KFT	HU
Narox 120 mg Film-coated tablets	not available	22268	DELORBIS PHARMACEUTICALS LTD	CY
Narox 30 mg Film-coated tablets	not available	22265	DELORBIS PHARMACEUTICALS LTD	CY
Narox 60 mg Film-coated tablets	not available	22266	DELORBIS PHARMACEUTICALS LTD	CY
Narox 90 mg Film-coated tablets	not available	22267	DELORBIS PHARMACEUTICALS LTD	CY
Oxidraxib 120 mg apvalkotās tabletes	NL/H/3267/003	16-0020	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LV
Oxidraxib 120 mg filmomhulde tabletten	NL/H/3267/003	RVG 116008	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	NL
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/035	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/036	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/037	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/038	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/039	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/040	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/041	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/042	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/043	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/044	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/045	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/046	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/047	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/048	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/049	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/050	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg filmsko obložene tablete	NL/H/3267/003	H/16/02096/051	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/035	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/036	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/037	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/038	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/039	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/040	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/041	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/042	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/043	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/044	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/045	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/046	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/047	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/048	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/049	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/050	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 120 mg plėvele dengtos tabletės	NL/H/3267/003	LT/1/16/3889/051	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
OXIDRAXIB 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/3267/003	72609/14/31.08.16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	GR
Oxidraxib 60 mg apvalkotās tabletes	NL/H/3267/001	16-0018	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Oxidraxib 60 mg filmomhulde tabletten	NL/H/3267/001	RVG 116006	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	NL
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/003	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/004	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/005	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/006	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/007	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/008	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/009	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/010	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/011	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/012	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/013	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/014	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/015	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/016	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg filmsko obložene tablete	NL/H/3267/001	H/16/02096/017	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/003	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/004	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/005	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/006	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/007	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/008	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/009	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/010	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/011	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/012	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/013	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/014	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/015	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/016	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 60 mg plėvele dengtos tabletės	NL/H/3267/001	LT/1/16/3889/017	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
OXIDRAXIB 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/3267/001	72607/14/31.08.16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	GR
Oxidraxib 90 mg apvalkotas tabletes	NL/H/3267/002	16-0019	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LV
Oxidraxib 90 mg filmomhulde tabletten	NL/H/3267/002	RVG 116007	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	NL
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/018	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/019	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/020	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/021	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/022	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/023	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/024	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/025	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/026	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/027	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/028	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/029	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/030	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/031	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/032	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/033	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg filmsko obložene tablete	NL/H/3267/002	H/16/02096/034	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/019	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/020	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/021	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/022	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/023	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/024	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/025	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/026	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/027	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/028	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/029	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/030	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/031	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/032	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/033	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Oxidraxib 90 mg plevele dengtos tabletes	NL/H/3267/002	LT/1/16/3889/034	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Oxidraxib 90 mg plėvele dengtos tabletės	NL/H/3267/002	LT/1/16/3889/018	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
OXIDRAXIB 90 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/3267/002	72608/14/31.08.16	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	GR
Oxidraxib, 120 mg õhukese polümeerikattega tabletid	NL/H/3267/003	901616	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
Oxidraxib, 60 mg õhukese polümeerikattega tabletid	NL/H/3267/001	901516	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
Oxidraxib, 90 mg õhukese polümeerikattega tabletid	NL/H/3267/002	901716	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/01	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/02	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/03	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/04	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/05	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/06	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/07	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/08	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
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/09	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/10	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg comprimate filmate	HU/H/0435/004	9484/2016/11	KRKA, D.D., NOVO MESTO	RO
Roticox 120 mg filmom obalené tablety	HU/H/0435/004	29/0459/16-S	KRKA, D.D., NOVO MESTO	SK
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/033	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/034	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/035	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/036	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/037	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/038	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/039	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/040	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/041	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/042	KRKA, D.D., NOVO MESTO	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
Roticox 120 mg filmsko obložene tablete	HU/H/0435/004	H/17/02295/043	KRKA, D.D., NOVO MESTO	SI
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/33	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/34	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/35	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/36	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/37	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/38	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/39	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/40	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/41	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/42	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg filmtabletta	HU/H/0435/004	OGYI-T-23097/43	KRKA, D.D., NOVO MESTO	HU
Roticox 120 mg potahované tablet	HU/H/0435/004	29/476/16-C	KRKA, D.D., NOVO MESTO	CZ
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/01	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/02	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/03	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/04	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/05	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/06	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/07	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/08	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg comprimate filmate	HU/H/0435/001	9481/2016/09	KRKA, D.D., NOVO MESTO	RO
Roticox 30 mg filmom obalené tablety	HU/H/0435/001	29/0456/16-S	KRKA, D.D., NOVO MESTO	SK
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/001	KRKA, D.D., NOVO MESTO	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
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/002	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/003	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/004	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/005	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/006	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/007	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/008	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmsko obložene tablete	HU/H/0435/001	H/17/02295/009	KRKA, D.D., NOVO MESTO	SI
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/01	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/02	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/03	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/04	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/05	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/06	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/07	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/08	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg filmtabletta	HU/H/0435/001	OGYI-T-23097/09	KRKA, D.D., NOVO MESTO	HU
Roticox 30 mg potahované tablet	HU/H/0435/001	29/473/16-C	KRKA, D.D., NOVO MESTO	CZ
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/01	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/02	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
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/03	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/04	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/05	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/06	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/07	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/08	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/09	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/10	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg comprimate filmate	HU/H/0435/002	9482/2016/11	KRKA, D.D., NOVO MESTO	RO
Roticox 60 mg filmom obalené tablety	HU/H/0435/002	29/0457/16-S	KRKA, D.D., NOVO MESTO	SK
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/010	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/011	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/012	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/013	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/014	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/015	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/016	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/017	KRKA, D.D., NOVO MESTO	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
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/018	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/019	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmsko obložene tablete	HU/H/0435/002	H/17/02295/020	KRKA, D.D., NOVO MESTO	SI
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/10	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/11	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/12	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/13	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/14	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/15	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/16	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/17	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/18	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/19	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg filmtabletta	HU/H/0435/002	OGYI-T-23097/20	KRKA, D.D., NOVO MESTO	HU
Roticox 60 mg potahované tablet	HU/H/0435/002	29/474/16-C	KRKA, D.D., NOVO MESTO	CZ
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/01	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/02	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/03	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/04	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/05	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/06	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/07	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/08	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/09	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/10	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
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/11	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg comprimate filmate	HU/H/0435/003	9483/2016/12	KRKA, D.D., NOVO MESTO	RO
Roticox 90 mg filmom obalené tablety	HU/H/0435/003	29/0458/16-S	KRKA, D.D., NOVO MESTO	SK
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/021	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/022	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/023	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/024	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/025	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/026	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/027	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/028	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/029	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/030	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/031	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmsko obložene tablete	HU/H/0435/003	H/17/02295/032	KRKA, D.D., NOVO MESTO	SI
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/21	KRKA, D.D., NOVO MESTO	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
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/22	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/23	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/24	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/25	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/26	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/27	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/28	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/29	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/30	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/31	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg filmtabletta	HU/H/0435/003	OGYI-T-23097/32	KRKA, D.D., NOVO MESTO	HU
Roticox 90 mg potahované tablet	HU/H/0435/003	29/475/16-C	KRKA, D.D., NOVO MESTO	CZ
Roticox, 120 mg, tabletki powlekane	HU/H/0435/004	23768	KRKA, D.D., NOVO MESTO	PL
Roticox, 30 mg, tabletki powlekane	HU/H/0435/001	23765	KRKA, D.D., NOVO MESTO	PL
Roticox, 60 mg, tabletki powlekane	HU/H/0435/002	23766	KRKA, D.D., NOVO MESTO	PL
Roticox, 90 mg, tabletki powlekane	HU/H/0435/003	23767	KRKA, D.D., NOVO MESTO	PL
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890298	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890425	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890300	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890312	ADDENDA PHARMA 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
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890324	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890336	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/01-3	035890348	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890351	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890363	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890375	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890387	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890399	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890401	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	UK/H/0535/001-003	035890413	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	UK/H/0535/001-003	035890449	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	UK/H/0535/001-003	035890437	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	UK/H/0535/001-003	035890452	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890019	ADDENDA PHARMA 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
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890021	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890033	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890045	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890058	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890060	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890072	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890084	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890096	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890108	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890110	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890122	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890134	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	035890146	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001-003	35890464	ADDENDA PHARMA 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
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890159	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890161	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890173	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890185	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890197	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890209	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890211	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890223	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890235	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890247	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/001-003	035890250	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/01-3	035890262	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/01-3	035890274	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	UK/H/0535/01-3	035890286	ADDENDA PHARMA 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
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115481	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114880	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4241584	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115580	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115382	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115283	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115689	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114484	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115085	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114989	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114682	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114385	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4115184	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114583	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
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4114781	MERCK SHARP & DOHME, LDA.	PT
Turox 120 mg comprimidos revestidos por película	UK/H/0535/003	4241683	MERCK SHARP & DOHME, LDA.	PT
Turox 30 mg comprimidos revestidos por película	UK/H/0535/004	5063052	MERCK SHARP & DOHME, LDA.	PT
Turox 30 mg comprimidos revestidos por película	UK/H/0535/004	5118617	MERCK SHARP & DOHME, LDA.	PT
Turox 30 mg comprimidos revestidos por película	UK/H/0535/004	5715040	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4117081	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116489	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116083	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116984	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4241287	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116786	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116588	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116687	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116281	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
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4115788	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4115986	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116182	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116380	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4115887	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4241188	MERCK SHARP & DOHME, LDA.	PT
Turox 60 mg comprimidos revestidos por película	UK/H/0535/001	4116885	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117388	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117289	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117180	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4118188	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117982	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4241485	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117685	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
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117784	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4241386	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4118386	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117883	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4118287	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4118485	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117586	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4118089	MERCK SHARP & DOHME, LDA.	PT
Turox 90 mg comprimidos revestidos por película	UK/H/0535/002	4117487	MERCK SHARP & DOHME, LDA.	PT
TUROX® 120 mg film-coated tablets	UK/H/0535/003	PL 0025/0433	MERCK SHARP & DOHME LTD.	UK
TUROX® 30 mg film-coated tablets	UK/H/0535/004	PL 0025/0481	MERCK SHARP & DOHME LTD.	UK
TUROX® 60 mg film-coated tablets	UK/H/0535/001	PL 0025/0431	MERCK SHARP & DOHME LTD.	UK
TUROX® 90 mg film-coated tablets	UK/H/0535/002	PL 0025/0432	MERCK SHARP & DOHME LTD.	UK
АРКОКСИЯ 120 mg филмирани таблетки	not available	20030176	MERCK SHARP & DOHME BULGARIA EOOD	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
АРКОКСИЯ 30 mg филмирани таблетки	not available	20100552	MERCK SHARP & DOHME BULGARIA EOOD	BG
АРКОКСИЯ 60 mg филмирани таблетки	not available	20030174	MERCK SHARP & DOHME BULGARIA EOOD	BG
АРКОКСИЯ 90 mg филмирани таблетки	not available	20030175	MERCK SHARP & DOHME BULGARIA EOOD	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 120 mg филмирани таблетки еторикоксиб	EE/H/0211/004	20160197	ZENTIVA, K.S.	BG
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG
Долоксиб 30 mg филмирани таблетки еторикоксиб	EE/H/0211/001	20160194	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 60 mg филмирани таблетки еторикоксиб	EE/H/0211/002	20160195	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Долоксиб 90 mg филмирани таблетки еторикоксиб	EE/H/0211/003	20160196	ZENTIVA, K.S.	BG
Еторикоксиб Тева 120 mg филмирани таблетки	DE/H/5031/004	20160114	TEVA PHARMACEUTICAL BULGARIA EOOD	BG
Еторикоксиб Тева 120 mg филмирани таблетки	DE/H/5031/004	20160114	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Еторикоксиб Тева 30 mg филмирани таблетки	DE/H/5031/001	20160111	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Еторикоксиб Тева 30 mg филмирани таблетки	DE/H/5031/001	20160111	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Еторикоксиб Тева 60 mg филмирани таблетки	DE/H/5031/002	20160112	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Еторикоксиб Тева 90 mg филмирани таблетки	DE/H/5031/003	20160113	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
КОКСИЕНТ 120 mg филмирани таблетки	IS/H/0244/004	II-35187	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 120 mg филмирани таблетки	IS/H/0244/004	II-35187	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 120 mg филмирани таблетки	IS/H/0244/004	II-35187	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 120 mg филмирани таблетки	IS/H/0244/004	II-35187	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 120 mg филмирани таблетки	IS/H/0244/004	II-35187	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 60 mg филмирани таблетки	IS/H/0244/002	II-35185	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 60 mg филмирани таблетки	IS/H/0244/002	II-35185	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
КОКСИЕНТ 60 mg филмирани таблетки	IS/H/0244/002	II-35185	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 60 mg филмирани таблетки	IS/H/0244/002	II-35185	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 60 mg филмирани таблетки	IS/H/0244/002	II-35185	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 90 mg филмирани таблетки	IS/H/0244/003	II-35186	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 90 mg филмирани таблетки	IS/H/0244/003	II-35186	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 90 mg филмирани таблетки	IS/H/0244/003	II-35186	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 90 mg филмирани таблетки	IS/H/0244/003	II-35186	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОКСИЕНТ 90 mg филмирани таблетки	IS/H/0244/003	II-35186	ALVOGEN MALTA OPERATIONS (ROW) LTD	BG
КОСТАРОКС 120 MG ФИЛМИРАНИ ТАБЛЕТКИ	DE/H/3908/004	20160272	SANDOZ PHARMACEUTICALS D.D.	BG
КОСТАРОКС 30 MG ФИЛМИРАНИ ТАБЛЕТКИ	DE/H/3908/001	20160269	SANDOZ PHARMACEUTICALS D.D.	BG
КОСТАРОКС 60 MG ФИЛМИРАНИ ТАБЛЕТКИ	DE/H/3908/002	20160270	SANDOZ PHARMACEUTICALS D.D.	BG
КОСТАРОКС 90 MG ФИЛМИРАНИ ТАБЛЕТКИ	DE/H/3908/003	20160271	SANDOZ PHARMACEUTICALS D.D.	BG
Ротикокс 120 mg филмирани таблетки	HU/H/0435/004	20170069	KRKA, D.D., NOVO MESTO	BG
Ротикокс 30 mg филмирани таблетки	HU/H/0435/001	20170066	KRKA, D.D., NOVO MESTO	BG

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Ротикокс 60 mg филмирани таблетки	HU/H/0435/002	20170067	KRKA, D.D., NOVO MESTO	BG
Ротикокс 90 mg филмирани таблетки	HU/H/0435/003	20170068	KRKA, D.D., NOVO MESTO	BG