01 December 2022
EMA/919649/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): etoricoxib

Procedure No. PSUSA/00001334/202203

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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
Arcoxia 120 mg Filmtabletten	PT/H/2299/003	1-24676	ORGANONAUSTRIA GMBH	AT
Arcoxia 30 mg Filmtabletten	PT/H/2299/004	1-27325	ORGANONAUSTRIA GMBH	AT
Arcoxia 60 mg Filmtabletten	PT/H/2299/001	1-24674	ORGANONAUSTRIA GMBH	AT
Arcoxia 90 mg Filmtabletten	PT/H/2299/002	1-24675	ORGANONAUSTRIA GMBH	AT
ARCOXIA 120 mg comprimés pelliculés	PT/H/2299/003	BE239556	N.V. ORGANON	BE
ARCOXIA 120 mg comprimés pelliculés	PT/H/2299/003	BE387405	N.V. ORGANON	BE
ARCOXIA 120 mg filmomhulde tabletten	PT/H/2299/003	BE239556	N.V. ORGANON	BE
ARCOXIA 120 mg filmomhulde tabletten	PT/H/2299/003	BE387405	N.V. ORGANON	BE
ARCOXIA 120 mg Filmtabletten	PT/H/2299/003	BE239556	N.V. ORGANON	BE
ARCOXIA 120 mg Filmtabletten	PT/H/2299/003	BE387405	N.V. ORGANON	BE
ARCOXIA 30 mg comprimés pelliculés	PT/H/2299/004	BE303886	N.V. ORGANON	BE
ARCOXIA 30 mg filmomhulde tabletten	PT/H/2299/004	BE303886	N.V. ORGANON	BE
ARCOXIA 30 mg Filmtabletten	PT/H/2299/004	BE303886	N.V. ORGANON	BE
ARCOXIA 60 mg comprimés pelliculés	PT/H/2299/001	BE239531	N.V. ORGANON	BE
ARCOXIA 60 mg comprimés pelliculés	PT/H/2299/001	BE387387	N.V. ORGANON	BE
ARCOXIA 60 mg filmomhulde tabletten	PT/H/2299/001	BE239531	N.V. ORGANON	BE
ARCOXIA 60 mg filmomhulde tabletten	PT/H/2299/001	BE387387	N.V. ORGANON	BE
ARCOXIA 60 mg	PT/H/2299/001	BE239531	N.V. ORGANON	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
Filmtabletten				
ARCOXIA 60 mg Filmtabletten	PT/H/2299/001	BE387387	N.V. ORGANON	BE
ARCOXIA 90 mg comprimés pelliculés	PT/H/2299/002	BE239547	N.V. ORGANON	BE
ARCOXIA 90 mg comprimés pelliculés	PT/H/2299/002	BE387396	N.V. ORGANON	BE
ARCOXIA 90 mg filmomhulde tabletten	PT/H/2299/002	BE239547	N.V. ORGANON	BE
ARCOXIA 90 mg filmomhulde tabletten	PT/H/2299/002	BE387396	N.V. ORGANON	BE
ARCOXIA 90 mg Filmtabletten	PT/H/2299/002	BE239547	N.V. ORGANON	BE
ARCOXIA 90 mg Filmtabletten	PT/H/2299/002	BE387396	N.V. ORGANON	BE
АРКОКСИЯ 120 mg филмирани таблетки	not available	20030176	N.V. ORGANON	BG
АРКОКСИЯ 30 mg филмирани таблетки	not available	20100552	N.V. ORGANON	BG
АРКОКСИЯ 60 mg филмирани таблетки	not available	20030174	N.V. ORGANON	BG
АРКОКСИЯ 90 mg филмирани таблетки	not available	20030175	N.V. ORGANON	BG
ARCOXIA 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/003	19446	N.V. ORGANON	CY
ARCOXIA 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/001	19444	N.V. ORGANON	CY
ARCOXIA 90 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/002	19445	N.V. ORGANON	CY
Arcoxia 30 mg potahované tablety	PT/H/2299/004	29/617/08-C	N.V. ORGANON	CZ
Arcoxia 60 mg potahované tablety	PT/H/2299/001	29/076/03-C	N.V. ORGANON	CZ
Arcoxia 90 mg potahované tablety	PT/H/2299/002	29/077/03-C	N.V. ORGANON	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
ACOXXEL 120 mg Filmtabletten	PT/H/2303/004	77702.00.00	ORGANONHEALTHCARE GMBH	DE
ACOXXEL 30 mg Filmtabletten	PT/H/2303/001	77699.00.00	ORGANONHEALTHCARE GMBH	DE
ACOXXEL 60 mg Filmtabletten	PT/H/2303/002	77700.00.00	ORGANONHEALTHCARE GMBH	DE
ACOXXEL 90 mg Filmtabletten	PT/H/2303/003	77701.00.00	ORGANONHEALTHCARE GMBH	DE
ARCOXIA 120 mg Filmtabletten	PT/H/2299/003	59863.02.00	ORGANONHEALTHCARE GMBH	DE
ARCOXIA 30 mg Filmtabletten	PT/H/2299/004	66968.00.00	ORGANONHEALTHCARE GMBH	DE
ARCOXIA 60 mg Filmtabletten	PT/H/2299/001	59863.00.00	ORGANONHEALTHCARE GMBH	DE
ARCOXIA 90 mg Filmtabletten	PT/H/2299/002	59863.01.00	ORGANONHEALTHCARE GMBH	DE
Arcoxia, filmovertrukne tabletter	PT/H/2299/001	33573	N.V. ORGANON	DK
Arcoxia, filmovertrukne tabletter	PT/H/2299/002	33574	N.V. ORGANON	DK
Arcoxia, filmovertrukne tabletter	PT/H/2299/003	33575	N.V. ORGANON	DK
Arcoxia, filmovertrukne tabletter	PT/H/2299/004	39698	N.V. ORGANON	DK
Arcoxia, 120 mg õhukese polümeerikattega tabletid	PT/H/2299/003	393002	N.V. ORGANON	EE
Arcoxia, 30 mg õhukese polümeerikattega tabletid	PT/H/2299/004	572308	N.V. ORGANON	EE
Arcoxia, 60 mg õhukese polümeerikattega tabletid	PT/H/2299/001	393202	N.V. ORGANON	EE
Arcoxia, 90 mg õhukese polümeerikattega tabletid	PT/H/2299/002	393102	N.V. ORGANON	EE
ACOXXEL 120 mg	PT/H/2303/004	71.590	ORGANONSALUD, S.L.	ES

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comprimidos recubiertos con película				
ACOXXEL 30 mg comprimidos recubiertos con película	PT/H/2303/001	71.587	ORGANONSALUD, S.L.	ES
ACOXXEL 60 mg comprimidos recubiertos con película	PT/H/2303/002	71.588	ORGANONSALUD, S.L.	ES
ACOXXEL 90 mg comprimidos recubiertos con película	PT/H/2303/003	71.589	ORGANONSALUD, S.L.	ES
ARCOXIA 120 mg comprimidos recubiertos con película	PT/H/2299/003	64.930	ORGANONSALUD, S.L.	ES
ARCOXIA 30 mg comprimidos recubiertos con película	PT/H/2299/004	69.376	ORGANONSALUD, S.L.	ES
ARCOXIA 60 mg comprimidos recubiertos con película	PT/H/2299/001	64.928	ORGANONSALUD, S.L.	ES
ARCOXIA 90 mg comprimidos recubiertos con película	PT/H/2299/002	64.929	ORGANONSALUD, S.L.	ES
EXXIV 120 mg comprimidos recubiertos con película	PT/H/2301/003	64.933	ORGANONSALUD, S.L.	ES
EXXIV 30 mg comprimidos recubiertos con película	PT/H/2301/004	72.076	ORGANONSALUD, S.L.	ES
EXXIV 60 mg comprimidos recubiertos con película	PT/H/2301/001	64.931	ORGANONSALUD, S.L.	ES
EXXIV 90 mg comprimidos recubiertos con película	PT/H/2301/002	64.932	ORGANONSALUD, S.L.	ES
Arcoxia 120 mg filmdragerade tabletter	PT/H/2299/003	17275	N.V. ORGANON	FI
Arcoxia 120 mg tabletti,	PT/H/2299/003	17275	N.V. ORGANON	FI

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kalvopäällysteinen				
Arcoxia 30 mg filmdragerade tabletter	PT/H/2299/004	22403	N.V. ORGANON	FI
Arcoxia 30 mg tabletti, kalvopäällysteinen	PT/H/2299/004	22403	N.V. ORGANON	FI
Arcoxia 60 mg filmdragerade tabletter	PT/H/2299/001	17273	N.V. ORGANON	FI
Arcoxia 60 mg tabletti, kalvopäällysteinen	PT/H/2299/001	17273	N.V. ORGANON	FI
Arcoxia 90 mg filmdragerade tabletter	PT/H/2299/002	17274	N.V. ORGANON	FI
Arcoxia 90 mg tabletti, kalvopäällysteinen	PT/H/2299/002	17274	N.V. ORGANON	FI
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 300 043 0 2	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 387 974 8 0	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 387 976 0 2	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 387 977 7 0	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 387 978 3 1	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 387 980 8 1	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 550 795 0 7	N.V. ORGANON	FR
ARCOXIA 30 mg, comprimé pelliculé	PT/H/2299/004	34009 573 530 9 4	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 275 698 9 0	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 923 4 8	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 924 0 9	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 925 7 7	N.V. ORGANON	FR
ARCOXIA 60 mg,	PT/H/2299/001	34009 387 926 3 8	N.V. ORGANON	FR

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comprimé pelliculé				
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 928 6 7	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 947 0 0	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 948 7 8	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 949 3 9	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 387 950 1 1	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 550 794 9 1	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 501 9 2	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 502 5 3	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 503 1 4	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 510 8 3	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 511 4 4	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 512 0 5	N.V. ORGANON	FR
ARCOXIA 60 mg, comprimé pelliculé	PT/H/2299/001	34009 573 513 7 3	N.V. ORGANON	FR
ARCOXIA® 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/003	71849/26-11-2018	VIANEX S.A.	GR
ARCOXIA® 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/004	71850/08-01-2019	VIANEX S.A.	GR
ARCOXIA® 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/2299/001	71847/26-11-2018	VIANEX S.A.	GR
ARCOXIA® 90 mg επικαλυμμένα με λεπτό	PT/H/2299/002	71848/26-11-2018	VIANEX S.A.	GR

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υμένιο δισκία				
ARCOXIA 120 mg filmom obložene tablete	not available	HR-H-130521686	ORGANON PHARMA DRUŠTVO S OGRANI?ENOM ODGOVORNOŠ?U ZA TRGOVINU I USLUGE	HR
ARCOXIA 30 mg filmom obložene tablete	not available	HR-H-662678488	ORGANON PHARMA DRUŠTVO S OGRANI?ENOM ODGOVORNOŠ?U ZA TRGOVINU I USLUGE	HR
ARCOXIA 60 mg filmom obložene tablete	not available	HR-H-150116833	ORGANON PHARMA DRUŠTVO S OGRANI?ENOM ODGOVORNOŠ?U ZA TRGOVINU I USLUGE	HR
ARCOXIA 90 mg filmom obložene tablete	not available	HR-H-231307450	ORGANON PHARMA DRUŠTVO S OGRANI?ENOM ODGOVORNOŠ?U ZA TRGOVINU I USLUGE	HR
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/01	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/02	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/03	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/04	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/10	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/11	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/12	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/13	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/14	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/15	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg	PT/H/2299/003	OGYI-T-8825/16	ORGANON HUNGARY KFT.	HU

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filmtabletta				
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/17	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/18	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/19	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/20	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/21	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg filmtabletta	PT/H/2299/003	OGYI-T-8825/58	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/05	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/22	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/23	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/24	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/25	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/26	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/55	ORGANON HUNGARY KFT.	HU
Arcoxia 30 mg filmtabletta	PT/H/2299/004	OGYI-T-8825/59	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/06	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/07	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/27	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/28	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg	PT/H/2299/001	OGYI-T-8825/29	ORGANON HUNGARY KFT.	HU

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filmtabletta				
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/30	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/31	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/32	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/33	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/34	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/35	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/36	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/37	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/38	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/39	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/40	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/56	ORGANON HUNGARY KFT.	HU
Arcoxia 60 mg filmtabletta	PT/H/2299/001	OGYI-T-8825/60	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/08	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/09	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/41	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/42	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/43	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg	PT/H/2299/002	OGYI-T-8825/44	ORGANON HUNGARY KFT.	HU

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filmtabletta				
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/45	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/46	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/47	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/48	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/49	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/50	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/51	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/52	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/53	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/54	ORGANON HUNGARY KFT.	HU
Arcoxia 90 mg filmtabletta	PT/H/2299/002	OGYI-T-8825/57	ORGANON HUNGARY KFT.	HU
Arcoxia 120 mg film-coated tablets	PT/H/2299/003	PA 0964/009/4	N.V. ORGANON	IE
Arcoxia 30 mg film-coated tablets	PT/H/2299/004	PA 0964/009/1	N.V. ORGANON	IE
Arcoxia 60 mg film-coated tablets	PT/H/2299/001	PA 0964/009/2	N.V. ORGANON	IE
Arcoxia 90 mg film-coated tablets	PT/H/2299/002	PA 0964/009/3	N.V. ORGANON	IE
ARCOXIA 120 mg filmuhúðaðar töflur	PT/H/2299/003	IS/1/02/025/03	N.V. ORGANON	IS
ARCOXIA 30 mg filmuhúðaðar töflur	PT/H/2299/004	IS/1/06/050/01	N.V. ORGANON	IS
ARCOXIA 60 mg filmuhúðaðar töflur	PT/H/2299/001	IS/1/02/025/01	N.V. ORGANON	IS
ARCOXIA 90 mg	PT/H/2299/002	IS/1/02/025/02	N.V. ORGANON	IS

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filmuhúðaðar töflur				
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821406	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821329	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821293	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821305	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821331	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821420	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821368	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821394	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821317	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821418	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821356	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821382	NEOPHARMED GENTILI SPA	IT
ALGIX 120 mg	UK/H/0533/004	035821343	NEOPHARMED GENTILI SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
ALGIX 120 mg compresse rivestite con film	UK/H/0533/004	035821370	NEOPHARMED GENTILI SPA	IT
ALGIX 30 mg compresse rivestite con film	UK/H/0533/001	035821457	NEOPHARMED GENTILI SPA	IT
ALGIX 30 mg compresse rivestite con film	UK/H/0533/001	035821444	NEOPHARMED GENTILI SPA	IT
ALGIX 30 mg compresse rivestite con film	UK/H/0533/001	035821432	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821469	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821077	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821115	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821014	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821141	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821089	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821139	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821091	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821103	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821040	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821038	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821065	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821053	NEOPHARMED GENTILI SPA	IT
ALGIX 60 mg compresse	UK/H/0533/002	035821127	NEOPHARMED GENTILI SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rivestite con film				
ALGIX 60 mg compresse rivestite con film	UK/H/0533/002	035821026	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821204	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821281	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821279	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821267	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821255	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821242	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821230	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821228	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821192	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821178	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821166	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821154	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821180	NEOPHARMED GENTILI SPA	IT
ALGIX 90 mg compresse rivestite con film	UK/H/0533/003	035821216	NEOPHARMED GENTILI SPA	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820378	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820380	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg	PT/H/2299/003	035820327	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
compresse rivestite con film				
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820341	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820291	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820339	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820303	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820392	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820354	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820366	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820315	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820416	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820428	MSD ITALIA S.R.L.	IT
ARCOXIA 120 mg compresse rivestite con film	PT/H/2299/003	035820404	MSD ITALIA S.R.L.	IT
ARCOXIA 30 mg compresse rivestite con film	PT/H/2299/004	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	PT/H/2299/004	035820430	MSD ITALIA S.R.L.	IT
ARCOXIA 30 mg compresse rivestite con film	PT/H/2299/004	035820568	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820063	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820051	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820048	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820024	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820012	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820036	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820075	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820099	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820101	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820113	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con	PT/H/2299/001	035820125	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
film				
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820137	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820149	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820087	MSD ITALIA S.R.L.	IT
ARCOXIA 60 mg compresse rivestite con film	PT/H/2299/001	035820570	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820164	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820152	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820176	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820188	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820253	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820289	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820238	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820190	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg	PT/H/2299/002	035820226	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
compresse rivestite con film				
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820277	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820214	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820265	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820240	MSD ITALIA S.R.L.	IT
ARCOXIA 90 mg compresse rivestite con film	PT/H/2299/002	035820202	MSD ITALIA S.R.L.	IT
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

Product 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	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
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	UK/H/0534/002	035822030	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
film				
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
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	UK/H/0534/003	035822156	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
compresse rivestite con film				
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
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

Product 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	035822283	ABIOGEN PHARMA S.P.A.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890298	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890425	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890300	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890312	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890324	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890336	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890348	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890351	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890363	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890375	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890387	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con	PT/H/2302/003	035890399	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
film				
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890401	ADDENDA PHARMA S.R.L.	IT
TAUXIB 120 mg compresse rivestite con film	PT/H/2302/003	035890413	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	PT/H/2302/004	035890449	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	PT/H/2302/004	035890437	ADDENDA PHARMA S.R.L.	IT
TAUXIB 30 mg compresse rivestite con film	PT/H/2302/004	035890452	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890019	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890021	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890033	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890045	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890058	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890060	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890072	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg	PT/H/2302/001	035890084	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
compresse rivestite con film				
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890096	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890108	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890110	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890122	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890134	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	UK/H/0535/001	035890146	ADDENDA PHARMA S.R.L.	IT
TAUXIB 60 mg compresse rivestite con film	PT/H/2302/001	035890464	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890159	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890161	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890173	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890185	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890197	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	PT/H/2302/002	035890209	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890211	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890223	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890235	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890247	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890250	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890262	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890274	ADDENDA PHARMA S.R.L.	IT
TAUXIB 90 mg compresse rivestite con film	PT/H/2302/002	035890286	ADDENDA PHARMA S.R.L.	IT
ARCOXIA 120 mg plėvele dengtos tabletės	PT/H/2299/003	LT/1/08/1419/008	N.V. ORGANON	LT
ARCOXIA 120 mg plėvele dengtos tabletės	PT/H/2299/003	LT/1/08/1419/009	N.V. ORGANON	LT
ARCOXIA 120 mg plėvele dengtos tabletės	PT/H/2299/003	LT/1/08/1419/010	N.V. ORGANON	LT
ARCOXIA 120 mg plėvele dengtos tabletės	PT/H/2299/003	LT/1/08/1419/011	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/001	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele	PT/H/2299/004	LT/1/08/1419/012	N.V. ORGANON	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
dengtos tabletės				
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/013	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/014	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/015	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/016	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/017	N.V. ORGANON	LT
ARCOXIA 30 mg plėvele dengtos tabletės	PT/H/2299/004	LT/1/08/1419/018	N.V. ORGANON	LT
ARCOXIA 60 mg plėvele dengtos tabletės	PT/H/2299/001	LT/1/08/1419/002	N.V. ORGANON	LT
ARCOXIA 60 mg plėvele dengtos tabletės	PT/H/2299/001	LT/1/08/1419/003	N.V. ORGANON	LT
ARCOXIA 60 mg plėvele dengtos tabletės	PT/H/2299/001	LT/1/08/1419/004	N.V. ORGANON	LT
ARCOXIA 60 mg plėvele dengtos tabletės	PT/H/2299/001	LT/1/08/1419/019	N.V. ORGANON	LT
ARCOXIA 90 mg plėvele dengtos tabletės	PT/H/2299/002	LT/1/08/1419/005	N.V. ORGANON	LT
ARCOXIA 90 mg plėvele dengtos tabletės	PT/H/2299/002	LT/1/08/1419/006	N.V. ORGANON	LT
ARCOXIA 90 mg plėvele dengtos tabletės	PT/H/2299/002	LT/1/08/1419/007	N.V. ORGANON	LT
ARCOXIA 120 mg comprimés pelliculés	PT/H/2299/003	2008019636	N.V. ORGANON	LU
ARCOXIA 30 mg comprimés pelliculés	PT/H/2299/004	2008020011	N.V. ORGANON	LU
ARCOXIA 60 mg comprimés pelliculés	PT/H/2299/001	2008019634	N.V. ORGANON	LU
ARCOXIA 90 mg comprimés pelliculés	PT/H/2299/002	2008019635	N.V. ORGANON	LU
ARCOXIA 120 mg apvalkotās tabletes	PT/H/2299/003	03-0005	N.V. ORGANON	LV
ARCOXIA 30 mg	PT/H/2299/004	07-0353	N.V. ORGANON	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
apvalkotās tabletes				
ARCOXIA 60 mg apvalkotās tabletes	PT/H/2299/001	03-0003	N.V. ORGANON	LV
ARCOXIA 90 mg apvalkotās tabletes	PT/H/2299/002	03-0004	N.V. ORGANON	LV
ARCOXIA 120 mg film-coated tablets	PT/H/2299/003	MA031/02103	N.V. ORGANON	MT
ARCOXIA 60 mg film-coated tablets	PT/H/2299/001	MA031/02102	N.V. ORGANON	MT
ARCOXIA 90 mg film-coated tablets	PT/H/2299/002	MA031/02101	N.V. ORGANON	MT
ARCOXIA 120 mg filmomhulde tabletten	PT/H/2299/003	RVG 27707	N.V. ORGANON	NL
ARCOXIA 30 mg filmomhulde tabletten	PT/H/2299/004	RVG 34279	N.V. ORGANON	NL
ARCOXIA 60 mg filmomhulde tabletten	PT/H/2299/001	RVG 27705	N.V. ORGANON	NL
ARCOXIA 90 mg filmomhulde tabletten	PT/H/2299/002	RVG 27706	N.V. ORGANON	NL
ARCOXIA 120 mg filmdrasjerte tabletter	PT/H/2299/003	02-1052	N.V. ORGANON	NO
ARCOXIA 30 mg filmdrasjerte tabletter	PT/H/2299/004	06-4274	N.V. ORGANON	NO
ARCOXIA 60 mg filmdrasjerte tabletter	PT/H/2299/001	02-1050	N.V. ORGANON	NO
ARCOXIA 90 mg filmdrasjerte tabletter	PT/H/2299/002	02-1051	N.V. ORGANON	NO
ARCOXIA, 120 mg, tabletki powlekane	PT/H/2299/003	11468	ORGANON POLSKA SP. Z O.O.	PL
ARCOXIA, 30 mg, tabletki powlekane	PT/H/2299/004	22580	ORGANON POLSKA SP. Z O.O.	PL
ARCOXIA, 60 mg, tabletki powlekane	PT/H/2299/001	11466	ORGANON POLSKA SP. Z O.O.	PL
ARCOXIA, 90 mg, tabletki powlekane	PT/H/2299/002	11467	ORGANON POLSKA SP. Z O.O.	PL
Acoxxel 120 mg comprimidos revestidos por película	PT/H/2303/004	PT/H/2303/004/E01	ORGANONPORTUGAL, SOCIEDADE 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
Acoxxel 30 mg comprimidos revestidos por película	PT/H/2303/001	PT/H/2303/001/E01	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Acoxxel 60 mg comprimidos revestidos por película	PT/H/2303/002	PT/H/2303/002/E01	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Acoxxel 90 mg comprimidos revestidos por película	PT/H/2303/003	PT/H/2303/003/E01	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106084	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106183	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106282	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106381	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106480	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106589	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106688	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106787	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4106886	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos	PT/H/2299/003	4106985	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película			LDA.	
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4107082	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4107181	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4107280	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4107389	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4242780	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 120 mg comprimidos revestidos por película	PT/H/2299/003	4242889	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 30 mg comprimidos revestidos por película	PT/H/2299/004	5063037	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 30 mg comprimidos revestidos por película	PT/H/2299/004	5118625	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 30 mg comprimidos revestidos por película	PT/H/2299/004	5715032	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4107488	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4107587	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4107686	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg	PT/H/2299/001	4107785	ORGANONPORTUGAL,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película			SOCIEDADE UNIPESSOAL LDA.	
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4107884	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4107983	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108080	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108189	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108288	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108387	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108486	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108585	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108684	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4108783	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4242384	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 60 mg comprimidos revestidos por película	PT/H/2299/001	4242483	ORGANONPORTUGAL, SOCIEDADE 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
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4108882	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4108981	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109088	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109187	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109286	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109385	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109484	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109583	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109682	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109781	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109880	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4109989	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos	PT/H/2299/002	4110086	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película			LDA.	
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4110185	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4242582	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Arcoxia 90 mg comprimidos revestidos por película	PT/H/2299/002	4242681	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Auxib 120 mg comprimidos revestidos por película	PT/H/2303/004	PT/H/2300/003/E02	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Auxib 30 mg comprimidos revestidos por película	PT/H/2303/001	PT/H/2300/004/E01	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Auxib 60 mg comprimidos revestidos por película	PT/H/2303/002	PT/H/2300/001/E02	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Auxib 90 mg comprimidos revestidos por película	PT/H/2303/003	PT/H/2300/002/E02	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114385	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114484	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114583	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114682	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114781	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg	PT/H/2302/003	4114880	ORGANONPORTUGAL,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película			SOCIEDADE UNIPESSOAL LDA.	
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114989	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115085	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115184	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115283	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115382	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115481	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115580	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115689	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4241584	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4241683	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114385	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114484	ORGANONPORTUGAL, SOCIEDADE 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
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114583	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114682	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114781	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114880	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4114989	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115085	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115184	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115283	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115382	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115481	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115580	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4115689	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 120 mg comprimidos revestidos	PT/H/2302/003	4241584	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película			LDA.	
Turox 120 mg comprimidos revestidos por película	PT/H/2302/003	4241683	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5063052	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5118617	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5715040	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5063052	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5118617	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 30 mg comprimidos revestidos por película	PT/H/2302/004	5715040	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115788	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115887	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115986	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116083	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116182	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg	PT/H/2302/001	4116281	ORGANONPORTUGAL,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película			SOCIEDADE UNIPESSOAL LDA.	
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116380	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116489	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116588	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116687	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116786	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116885	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116984	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4117081	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4241188	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4241287	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115788	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115887	ORGANONPORTUGAL, SOCIEDADE 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
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4115986	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116083	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116182	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116281	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116380	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116489	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116588	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116687	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116786	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116885	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4116984	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4117081	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 60 mg comprimidos revestidos	PT/H/2302/001	4241188	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película			LDA.	
Turox 60 mg comprimidos revestidos por película	PT/H/2302/001	4241287	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117180	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117289	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117388	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117487	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117586	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117685	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117784	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117883	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117982	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118089	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118188	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg	PT/H/2302/002	4118287	ORGANONPORTUGAL,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película			SOCIEDADE UNIPESSOAL LDA.	
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118386	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118485	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4241386	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4241485	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117180	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117289	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117388	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117487	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117586	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117685	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117784	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117883	ORGANONPORTUGAL, SOCIEDADE 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
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4117982	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118089	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118188	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118287	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118386	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4118485	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4241386	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
Turox 90 mg comprimidos revestidos por película	PT/H/2302/002	4241485	ORGANONPORTUGAL, SOCIEDADE UNIPESSOAL LDA.	PT
ARCOXIA 120 mg comprimate filmate	not available	854/2008/01	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 120 mg comprimate filmate	not available	854/2008/02	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 120 mg comprimate filmate	not available	854/2008/03	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 120 mg comprimate filmate	not available	854/2008/04	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/01	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/02	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/03	ORGANON BIOSCIENCES 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
ARCOXIA 30 mg comprimate filmate	not available	7877/2015/04	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/01	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/02	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/03	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 60 mg comprimate filmate	not available	852/2008/04	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/01	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/02	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/03	ORGANON BIOSCIENCES S.R.L.	RO
ARCOXIA 90 mg comprimate filmate	not available	853/2008/04	ORGANON BIOSCIENCES S.R.L.	RO
Arcoxia 120 mg filmdragerade tabletter	PT/H/2299/003	18367	MERCK SHARP & DOHME BV	SE
ARCOXIA 30 mg filmdragerade tabletter	PT/H/2299/004	24178	MERCK SHARP & DOHME BV	SE
Arcoxia 60 mg filmdragerade tabletter	PT/H/2299/001	18365	MERCK SHARP & DOHME BV	SE
Arcoxia 90 mg filmdragerade tabletter	PT/H/2299/002	18366	MERCK SHARP & DOHME BV	SE
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/042	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/043	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/044	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/045	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/046	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/047	N.V. ORGANON	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	PT/H/2299/003	H/02/00207/048	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/049	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/050	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/051	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/052	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/053	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/054	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/055	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/056	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/057	N.V. ORGANON	SI
ARCOXIA 120 mg filmsko obložene tablete	PT/H/2299/003	H/02/00207/058	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/001	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/002	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/003	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/004	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/005	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/006	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/007	N.V. ORGANON	SI
ARCOXIA 30 mg filmsko obložene tablete	PT/H/2299/004	H/02/00207/059	N.V. ORGANON	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	PT/H/2299/001	H/02/00207/008	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/009	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/010	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/011	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/012	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/013	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/014	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/015	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/016	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/017	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/018	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/019	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/020	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/021	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/022	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/023	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/024	N.V. ORGANON	SI
ARCOXIA 60 mg filmsko obložene tablete	PT/H/2299/001	H/02/00207/060	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/025	N.V. ORGANON	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	PT/H/2299/002	H/02/00207/026	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/027	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/028	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/029	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/030	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/031	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/032	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/033	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/034	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/035	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/036	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/037	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/038	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/039	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/040	N.V. ORGANON	SI
ARCOXIA 90 mg filmsko obložene tablete	PT/H/2299/002	H/02/00207/041	N.V. ORGANON	SI
ARCOXIA 120 mg filmom obalené tablety	PT/H/2299/003	29/0005/03-S	N.V. ORGANON	SK
ARCOXIA 30 mg filmom obalené tablety	PT/H/2299/004	29/0125/08-S	N.V. ORGANON	SK
ARCOXIA 60 mg filmom obalené tablety	PT/H/2299/001	29/0003/03-S	N.V. ORGANON	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 filmom obalené tablety	PT/H/2299/002	29/0004/03-S	N.V. ORGANON	SK
ARCOXIA® 120 mg film-coated tablets	PT/H/2299/003	PL 00025/0644	MERCK SHARP & DOHME LTD.	XI
ARCOXIA® 30 mg film-coated tablets	PT/H/2299/004	PL 00025/0641	MERCK SHARP & DOHME LTD.	XI
ARCOXIA® 60 mg film-coated tablets	PT/H/2299/001	PL 00025/0642	MERCK SHARP & DOHME LTD.	XI
ARCOXIA® 90 mg film-coated tablets	PT/H/2299/002	PL 00025/0643	MERCK SHARP & DOHME LTD.	XI