30 November 2017
EMA/787932/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): cefuroxime axetil

Procedure No.: PSUSA/00009099/201704

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Send a question via our website www.ema.europa.eu/contact

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/09	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/10	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/11	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/12	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 125 mg comprimate	PT/H/0461/001	8302/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO

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Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/09	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/10	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/11	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 250 mg comprimate	PT/H/0461/002	8303/12	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/01	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/02	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/03	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/04	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/05	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/07	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/08	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/09	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/10	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/11	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/12	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO
Cefuroxima Aurobindo 500 mg comprimate	PT/H/0461/003	8304/06	AUROBINDO PHARMA ROMÂNIA S.R.L.	RO

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Cefuroxime 250mg Tablets	NL/H/0556/002	PL 04416/0627	SANDOZ LTD	UK
CEFUROXIME BIOGARAN 125 mg, comprimé pelliculé	not available	NL14829	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME BIOGARAN 125 mg, comprimé pelliculé	not available	NL14829	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME BIOGARAN 250 mg, comprimé pelliculé	not available	NL14424	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME BIOGARAN 250 mg, comprimé pelliculé	not available	NL14424	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME BIOGARAN 500 mg, comprimé pelliculé	not available	NL21711	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME BIOGARAN 500 mg, comprimé pelliculé	not available	NL21711	LABORATOIRES SAINT-GERMAIN	FR
CEFUROXIME ZENTIVA 250 mg, comprimé pelliculé	not available	NL14426	LABORATOIRES PAUCOURT	FR
CEFUROXIME ZENTIVA 250 mg, comprimé pelliculé	not available	NL14426	LABORATOIRES PAUCOURT	FR
Cefuroxim-ratiopharm® 250 mg Filmtabletten	DE/H/1426/001	70940.00.00	RATIOPHARM GMBH	DE
Cefuroxim-ratiopharm® 500 mg Filmtabletten	DE/H/1426/002	70941.00.00	RATIOPHARM GMBH	DE
Elobact® 125 mg Filmtabletten	UK/H/5462/003	16468.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact® 125 mg Filmtabletten	UK/H/5462/003	16468.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact® 250 mg Filmtabletten	UK/H/5462/004	16468.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact® 250 mg Filmtabletten	UK/H/5462/004	16468.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact® 500 mg Filmtabletten	UK/H/5462/005	16468.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact® 500 mg Filmtabletten	UK/H/5462/005	16468.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Elobact®-Trockensaft 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5462/01	16468.00.01	GLAXOSMITHKLINE GMBH & CO. KG	DE

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Elobact®-Trockensaft 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5462/01	16468.00.01	GLAXOSMITHKLINE GMBH & CO. KG	DE
Oraxim 125 mg/5 ml granulato per sospensione orale	IT/H/0369/004	027002043	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
Oraxim 250 mg compresse rivestite con film	IT/H/0369/001	027002029	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
Oraxim 250 mg granulato per sospensione orale	IT/H/0369/003	027002056	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
Oraxim 250 mg/5 ml granulato per sospensione orale	IT/H/0369/005	027002070	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
Oraxim 500 mg compresse rivestite con film	IT/H/0369/002	027002106	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
Oraxim 500 mg compresse rivestite con film	IT/H/0369/002	027002031	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
TILEXIM 250 mg Compresse rivestite con film	not available	027020027	I.B.N. SAVIO S.R.L.	IT
TILEXIM 500 mg Compresse rivestite con film	not available	027020039	I.B.N. SAVIO S.R.L.	IT
TILEXIM 500 mg Compresse rivestite con film	not available	027020054	I.B.N. SAVIO S.R.L.	IT
Zinadol 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5462/004	1965002	GLAXOSMITHKLINE AEBE	GR
Zinadol 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5462/004	1965002	GLAXOSMITHKLINE AEBE	GR
Zinadol 250 mg/5 ml κοκκία για πόσιμο εναιώρημα	UK/H/5462/002	1965006	GLAXOSMITHKLINE AEBE	GR
Zinadol 250 mg/5 ml κοκκία για πόσιμο εναιώρημα	UK/H/5462/002	1965006	GLAXOSMITHKLINE AEBE	GR
Zinadol 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5462/05	1965003	GLAXOSMITHKLINE AEBE	GR
Zinadol 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5462/05	1965003	GLAXOSMITHKLINE AEBE	GR
ZINNAT 125 mg ENFANTS ET NOURRISSONS, granulés pour suspension buvable en sachet-dose	ES/H/0236/001	NL16770	LABORATOIRE GLAXOSMITHKLINE	FR

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ZINNAT 125 mg ENFANTS ET NOURRISSONS, granulés pour suspension buvable en sachet-dose	ES/H/0236/001	NL16770	LABORATOIRE GLAXOSMITHKLINE	FR
Zinnat 125 mg film-coated tablets	UK/H/5462/003	PA 1077/15/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 125 mg film-coated tablets	UK/H/5462/003	PA 1077/15/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 125 mg filmom obalené tablety	UK/H/5462/03	15/0061/88-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 125 mg filmom obalené tablety	UK/H/5462/03	15/0061/88-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ZINNAT 125 mg filmom obložene tablete	not available	HR-H-391373111	GLAXOSMITHKLINE D.O.O.	HR
ZINNAT 125 mg filmom obložene tablete	not available	HR-H-391373111	GLAXOSMITHKLINE D.O.O.	HR
Zinnat 125 mg filmomhulde tabletten	UK/H/5462/03	RVG 13225	GLAXOSMITHKLINE B.V.	NL
Zinnat 125 mg filmomhulde tabletten	UK/H/5462/03	RVG 13225	GLAXOSMITHKLINE B.V.	NL
Zinnat 125 mg filmtabletta	UK/H/5462/03	OGYI-T-1399/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 125 mg filmtabletta	UK/H/5462/03	OGYI-T-1399/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 125 mg granulado para suspensión oral	ES/H/0236/001	59.062	GLAXOSMITHKLINE S.A.	ES
Zinnat 125 mg granulado para suspensión oral	ES/H/0236/001	59.062	GLAXOSMITHKLINE S.A.	ES
Zinnat 125 mg granule pro perorální suspenzi	UK/H/5462/001	15/390/92-A/C	GLAXO GROUP LIMITED	CZ
Zinnat 125 mg granule pro perorální suspenzi	UK/H/5462/001	15/390/92-A/C	GLAXO GROUP LIMITED	CZ
Zinnat 125 mg plėvele dengtos tabletės	UK/H/5462/003	LT/1/94/0478/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg plėvele dengtos tabletės	UK/H/5462/003	LT/1/94/0478/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg plėvele dengtos tabletės	UK/H/5462/003	LT/1/94/0478/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg plėvele dengtos tabletės	UK/H/5462/003	LT/1/94/0478/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg potahované tablety	UK/H/5462/003	15/061/88-A/C	GLAXO GROUP LIMITED	CZ
Zinnat 125 mg potahované tablety	UK/H/5462/003	15/061/88-A/C	GLAXO GROUP LIMITED	CZ

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 125 mg vrecká granulát na perorálnu suspenziu	ES/H/0236/001	15/0244/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 125 mg vrecká granulát na perorálnu suspenziu	ES/H/0236/001	15/0244/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 125 mg, comprimate filmate	UK/H/5462/003	2016/2009/01	GLAXO WELLCOME LTD	RO
Zinnat 125 mg, comprimate filmate	UK/H/5462/003	2016/2009/01	GLAXO WELLCOME LTD	RO
ZINNAT 125 mg, comprimé pelliculé	UK/H/5462/03	NL14830	LABORATOIRE GLAXOSMITHKLINE	FR
ZINNAT 125 mg, comprimé pelliculé	UK/H/5462/03	NL14830	LABORATOIRE GLAXOSMITHKLINE	FR
ZINNAT 125 mg/5 ml ENFANTS ET NOURRISSONS, granulés pour suspension buvable en flacon	UK/H/5462/001	NL16772	LABORATOIRE GLAXOSMITHKLINE	FR
ZINNAT 125 mg/5 ml ENFANTS ET NOURRISSONS, granulés pour suspension buvable en flacon	UK/H/5462/001	NL16772	LABORATOIRE GLAXOSMITHKLINE	FR
Zinnat 125 mg/5 ml granulaat voor orale suspensie	UK/H/5462/01	RVG 14376	GLAXOSMITHKLINE B.V.	NL
Zinnat 125 mg/5 ml granulaat voor orale suspensie	UK/H/5462/01	RVG 14376	GLAXOSMITHKLINE B.V.	NL
Zinnat 125 mg/5 ml granulát na perorálnu suspenziu	UK/H/5462/01	15/0390/92-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 125 mg/5 ml granulát na perorálnu suspenziu	UK/H/5462/01	15/0390/92-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 125 mg/5 ml granulátum belsőleges szuszpenzióhoz	UK/H/5462/01	OGYI-T-1830/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 125 mg/5 ml granulátum belsőleges szuszpenzióhoz	UK/H/5462/01	OGYI-T-1830/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 125 mg/5 ml granule pentru suspensie orală	UK/H/5462/001	5184/2012/01	GLAXO WELLCOME UK LIMITED	RO
Zinnat 125 mg/5 ml granule pentru suspensie orală	UK/H/5462/001	5184/2012/01	GLAXO WELLCOME UK LIMITED	RO
ZINNAT 125 mg/5 ml granule za oralnu suspenziju	not available	UP/I-530-09/12-02/473	GLAXOSMITHKLINE D.O.O.	HR
ZINNAT 125 mg/5 ml granule za oralnu suspenziju	not available	UP/I-530-09/12-02/473	GLAXOSMITHKLINE D.O.O.	HR

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Zinnat 125 mg/5 ml granules for oral suspension	UK/H/5462/001	PA 1077/15/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 125 mg/5 ml granules for oral suspension	UK/H/5462/001	PA 1077/15/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 125 mg/5 ml granules for oral suspension	UK/H/5462/01	MA 192/01203	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 125 mg/5 ml granules for oral suspension	UK/H/5462/01	MA 192/01203	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/008	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/009	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/007	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/008	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/009	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml granulės geriamajai suspensijai	UK/H/5462/001	LT/1/94/0478/007	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 125 mg/5 ml zrnca za peroralno suspenzijo	UK/H/5462/01	H/92/01704/003	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 125 mg/5 ml zrnca za peroralno suspenzijo	UK/H/5462/01	H/92/01704/003	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 125 mg/5 ml granulato per sospensione orale	UK/H/5462/001	026915049	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 125 mg/5 ml granulato per sospensione orale	UK/H/5462/001	026915049	GLAXOSMITHKLINE S.P.A.	IT
ZINNAT 25 mg/ ml granulas iekšķigi lietojamas suspensijas pagatavošanai	UK/H/5462/01	95-0049	GLAXOSMITHKLINE LATVIA SIA	LV
ZINNAT 25 mg/ ml granulas iekšķigi lietojamas suspensijas pagatavošanai	UK/H/5462/01	95-0049	GLAXOSMITHKLINE LATVIA SIA	LV
Zinnat 250 mg apvalkotās tabletes	UK/H/5462/04	97-0234	GLAXOSMITHKLINE LATVIA SIA	LV
Zinnat 250 mg apvalkotās tabletes	UK/H/5462/04	97-0234	GLAXOSMITHKLINE LATVIA SIA	LV
ZINNAT 250 mg Compresse rivestite con film	UK/H/5462/004	026915025	GLAXOSMITHKLINE S.P.A.	IT

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ZINNAT 250 mg Compresse rivestite con film	UK/H/5462/004	026915025	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 250 mg comprimidos recubiertos con película	UK/H/5462/004	58.305	GLAXOSMITHKLINE S.A.	ES
Zinnat 250 mg comprimidos recubiertos con película	UK/H/5462/004	58.305	GLAXOSMITHKLINE S.A.	ES
Zinnat 250 mg film-coated tablets	UK/H/5462/04	PA 1077/15/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 250 mg film-coated tablets	UK/H/5462/04	PA 1077/15/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 250 mg film-coated tablets	UK/H/5462/04	MA 167/01401	GLAXO WELLCOME UK LIMITED	MT
Zinnat 250 mg film-coated tablets	UK/H/5462/04	MA 167/01401	GLAXO WELLCOME UK LIMITED	MT
Zinnat 250 mg filmom obalené tablety	UK/H/5462/04	15/0243/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 250 mg filmom obalené tablety	UK/H/5462/04	15/0243/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ZINNAT 250 mg filmom obložene tablete	not available	HR-H-074775680	GLAXOSMITHKLINE D.O.O.	HR
ZINNAT 250 mg filmom obložene tablete	not available	HR-H-074775680	GLAXOSMITHKLINE D.O.O.	HR
Zinnat 250 mg filmomhulde tabletten	UK/H/5462/04	RVG 13226	GLAXOSMITHKLINE B.V.	NL
Zinnat 250 mg filmomhulde tabletten	UK/H/5462/04	RVG 13226	GLAXOSMITHKLINE B.V.	NL
Zinnat 250 mg filmsko obložene tablete	UK/H/5462/004	H/92/01704/001	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 250 mg filmsko obložene tablete	UK/H/5462/004	H/92/01704/001	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 250 mg filmtabletta	UK/H/5462/04	OGYI-T-1400/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 250 mg filmtabletta	UK/H/5462/04	OGYI-T-1400/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 250 mg granulado para suspensión oral	ES/H/0236/002	59.063	GLAXOSMITHKLINE S.A.	ES
Zinnat 250 mg granulado para suspensión oral	ES/H/0236/002	59.063	GLAXOSMITHKLINE S.A.	ES
ZINNAT 250 mg Granulato per sospensione orale	ES/H/0236/002	026915052	GLAXOSMITHKLINE S.P.A.	IT
ZINNAT 250 mg Granulato per sospensione orale	ES/H/0236/002	026915052	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 250 mg plėvele dengtos tabletės	UK/H/5462/004	LT/1/94/0478/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 250 mg plėvele dengtos tabletės	UK/H/5462/004	LT/1/94/0478/004	GLAXOSMITHKLINE LIETUVA UAB	LT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 250 mg plėvele dengtos tabletės	UK/H/5462/004	LT/1/94/0478/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 250 mg plėvele dengtos tabletės	UK/H/5462/004	LT/1/94/0478/004	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 250 mg potahované tablety	UK/H/5462/004	15/061/88-B/C	GLAXO GROUP LIMITED	CZ
Zinnat 250 mg potahované tablety	UK/H/5462/004	15/061/88-B/C	GLAXO GROUP LIMITED	CZ
Zinnat 250 mg tabletti, kalvopäällysteinen	UK/H/5462/004	10216	GLAXO OPERATIONS UK LTD	FI
Zinnat 250 mg tabletti, kalvopäällysteinen	UK/H/5462/004	10216	GLAXO OPERATIONS UK LTD	FI
Zinnat 250 mg tabletti, kalvopäällysteinen	UK/H/5462/004	10216	GLAXO OPERATIONS UK LTD	FI
Zinnat 250 mg tabletti, kalvopäällysteinen	UK/H/5462/004	10216	GLAXO OPERATIONS UK LTD	FI
Zinnat 250 mg vrecká granulát na perorálnu suspenziu	ES/H/0236/002	15/0245/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 250 mg vrecká granulát na perorálnu suspenziu	ES/H/0236/002	15/0245/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	16846	GLAXO GROUP LIMITED	CY
Zinnat 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	16846	GLAXO GROUP LIMITED	CY
Zinnat 250 mg, comprimate filmate	UK/H/5462/004	2017/2009/01	GLAXO WELLCOME UK LIMITED	RO
Zinnat 250 mg, comprimate filmate	UK/H/5462/004	2017/2009/01	GLAXO WELLCOME UK LIMITED	RO
ZINNAT 250 mg, comprimé pelliculé	UK/H/5462/04	NL14425	LABORATOIRE GLAXOSMITHKLINE	FR
ZINNAT 250 mg, comprimé pelliculé	UK/H/5462/04	NL14425	LABORATOIRE GLAXOSMITHKLINE	FR
Zinnat 250 mg, comprimés pelliculés	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg, comprimés pelliculés	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg, comprimés pelliculés	UK/H/5462/04	260/02 10 6750	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg, filmomhulde tabletten	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 250 mg, filmomhulde tabletten	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg, filmomhulde tabletten	UK/H/5462/004	260/02 10 6750	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg, filmomhulde tabletten	UK/H/5462/004	260/02 10 6750	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg, Filmtabletten	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg, Filmtabletten	UK/H/5462/004	BE 151121	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg, Filmtabletten	UK/H/5462/004	260/02 10 6750	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg/5 ml granulado para suspensión oral	UK/H/5462/002	62.806	GLAXOSMITHKLINE S.A.	ES
Zinnat 250 mg/5 ml granulado para suspensión oral	UK/H/5462/002	62.806	GLAXOSMITHKLINE S.A.	ES
Zinnat 250 mg/5 ml granulát na perorálnu suspenziu	UK/H/5462/02	15/0246/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 250 mg/5 ml granulát na perorálnu suspenziu	UK/H/5462/02	15/0246/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ZINNAT 250 mg/5 ml Granulato per sospensione orale	UK/H/5462/002	026915076	GLAXOSMITHKLINE S.P.A.	IT
ZINNAT 250 mg/5 ml Granulato per sospensione orale	UK/H/5462/002	026915076	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 250 mg/5 ml granules for oral suspension	UK/H/5462/02	MA 192/01201	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 250 mg/5 ml granules for oral suspension	UK/H/5462/02	MA 192/01201	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 250 mg/5 ml granules for oral suspension.	UK/H/5462/002	PA 1077/15/1	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 250 mg/5 ml granules for oral suspension.	UK/H/5462/002	PA 1077/15/1	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 250 mg/5 ml zrnca za peroralno suspenzijo	UK/H/5462/002	H/92/01704/004	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 250 mg/5 mL κοκκία για πόσιμο εναιώρημα	not available	18086	GLAXO GROUP LIMITED	CY

Product full name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 250 mg/5 mL κοκκία για πόσιμο εναιώρημα	not available	18086	GLAXO GROUP LIMITED	CY
Zinnat 250 mg/5 ml, granulaat voor orale suspensie	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, granulaat voor orale suspensie	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, granulaat voor orale suspensie	UK/H/5462/002	2002 10 6755	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg/5 ml, granulaat voor orale suspensie	UK/H/5462/002	2002 10 6755	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5462/002	2002 10 6755	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg/5 ml, granulés pour suspension buvable	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, granulés pour suspension buvable	UK/H/5462/002	BE 198195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 250 mg/5 ml, granulés pour suspension buvable	UK/H/5462/002	2002 10 6755	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 250 mg-Filmtabletten	UK/H/5462/004	1-19106	GLAXOSMITHKLINE PHARMA GMBH.	AT
Zinnat 250 mg-Filmtabletten	UK/H/5462/004	1-19106	GLAXOSMITHKLINE PHARMA GMBH.	AT
Zinnat 500 mg apvalkotās tabletes	UK/H/5462/05	97-0235	GLAXOSMITHKLINE LATVIA SIA	LV
Zinnat 500 mg apvalkotās tabletes	UK/H/5462/05	97-0235	GLAXOSMITHKLINE LATVIA SIA	LV
Zinnat 500 mg compresse rivestite con film	UK/H/5462/005	026915037	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 500 mg comprimidos recubiertos con película	UK/H/5462/005	58.309	GLAXOSMITHKLINE S.A.	ES
Zinnat 500 mg comprimidos recubiertos con película	UK/H/5462/005	58.309	GLAXOSMITHKLINE S.A.	ES

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 500 mg film-coated tablets	UK/H/5462/005	PA 1077/15/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Zinnat 500 mg film-coated tablets	UK/H/5462/05	MA192/01202	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 500 mg film-coated tablets	UK/H/5462/05	MA192/01202	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Zinnat 500 mg filmom obalené tablety	UK/H/5462/05	15/0247/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Zinnat 500 mg filmom obalené tablety	UK/H/5462/05	15/0247/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ZINNAT 500 mg filmom obložene tablete	not available	HR-H-565616124	GLAXOSMITHKLINE D.O.O.	HR
ZINNAT 500 mg filmom obložene tablete	not available	HR-H-565616124	GLAXOSMITHKLINE D.O.O.	HR
Zinnat 500 mg filmomhulde tabletten	UK/H/5462/05	RVG 13227	GLAXOSMITHKLINE B.V.	NL
Zinnat 500 mg filmomhulde tabletten	UK/H/5462/05	RVG 13227	GLAXOSMITHKLINE B.V.	NL
Zinnat 500 mg filmsko obložene tablete	UK/H/5462/005	H/92/01704/002	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 500 mg filmsko obložene tablete	UK/H/5462/005	H/92/01704/002	GLAXOSMITHKLINE D.O.O.	SI
Zinnat 500 mg filmtabletta	UK/H/5462/05	OGYI-T-1401/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 500 mg filmtabletta	UK/H/5462/05	OGYI-T-1401/01	GLAXOSMITHKLINE KFT.	HU
Zinnat 500 mg granulado para suspensión oral	ES/H/0236/003	60.656	GLAXOSMITHKLINE S.A.	ES
Zinnat 500 mg granulado para suspensión oral	ES/H/0236/003	60.656	GLAXOSMITHKLINE S.A.	ES
Zinnat 500 mg plėvele dengtos tabletės	UK/H/5462/005	LT/1/94/0478/005	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 500 mg plėvele dengtos tabletės	UK/H/5462/005	LT/1/94/0478/006	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 500 mg plėvele dengtos tabletės	UK/H/5462/005	LT/1/94/0478/005	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 500 mg plėvele dengtos tabletės	UK/H/5462/005	LT/1/94/0478/006	GLAXOSMITHKLINE LIETUVA UAB	LT
Zinnat 500 mg potahované tablety	UK/H/5462/005	15/061/88-C/C	GLAXO GROUP LIMITED	CZ
Zinnat 500 mg potahované tablety	UK/H/5462/005	15/061/88-C/C	GLAXO GROUP LIMITED	CZ
Zinnat 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	16847	GLAXO GROUP LIMITED	CY
Zinnat 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	16847	GLAXO GROUP LIMITED	CY

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat 500 mg, comprimate filmate	UK/H/5462/005	2018/2009/01	GLAXO WELLCOME UK LIMITED	RO
Zinnat 500 mg, comprimate filmate	UK/H/5462/005	2018/2009/01	GLAXO WELLCOME UK LIMITED	RO
Zinnat 500 mg, comprimés pelliculés	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, comprimés pelliculés	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, comprimés pelliculés	UK/H/5462/005	260/02 10 6751	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 500 mg, filmomhulde tabletten	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, filmomhulde tabletten	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, filmomhulde tabletten	UK/H/5462/005	260/02 10 6751	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 500 mg, filmomhulde tabletten	UK/H/5462/005	260/02 10 6751	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 500 mg, Filmtabletten	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, Filmtabletten	UK/H/5462/005	BE 154585	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Zinnat 500 mg, Filmtabletten	UK/H/5462/005	260/02 10 6751	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Zinnat 500 mg-Filmtabletten	UK/H/5462/005	1-19105	GLAXOSMITHKLINE PHARMA GMBH.	AT
Zinnat 500 mg-Filmtabletten	UK/H/5462/005	1-19105	GLAXOSMITHKLINE PHARMA GMBH.	AT
Zinnat 500 mg compresse rivestite con film	UK/H/5462/005	026915102	GLAXOSMITHKLINE S.P.A.	IT
Zinnat 500 mg compresse rivestite con film	UK/H/5462/005	026915102	GLAXOSMITHKLINE S.P.A.	IT
Zinnat Suspension 125mg/5ml	UK/H/5462/01	PL10949/0094	GLAXO WELLCOME UK LIMITED	UK
Zinnat Suspension 250mg/5ml	UK/H/5462/02	PL10949/0272	GLAXO WELLCOME UK LIMITED	UK
Zinnat Tablets 125mg	UK/H/5462/003	PL 10949/0095	GLAXO WELLCOME LTD	UK
Zinnat Tablets 250mg	UK/H/5462/004	PL10949/0096	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zinnat Tablets 500mg	UK/H/5462/05	PL 10949/0119	GLAXO WELLCOME LTD	UK
Zinnat, 125 mg, tabletki powlekane	UK/H/5462/03	R/0832	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 125 mg, tabletki powlekane	UK/H/5462/03	R/0832	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 125 mg/5 ml, granulat do sporządzania zawiesiny doustnej	UK/H/5462/01	R/0513	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 125 mg/5 ml, granulat do sporządzania zawiesiny doustnej	UK/H/5462/01	R/0513	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 25 mg/ml suukaudse suspensiooni graanulid	UK/H/5462/01	226698	GLAXO GROUP LIMITED	EE
Zinnat, 25 mg/ml suukaudse suspensiooni graanulid	UK/H/5462/01	226698	GLAXO GROUP LIMITED	EE
Zinnat, 250 mg õhukese polümeerikattega tabletid	UK/H/5462/04	226498	GLAXO GROUP LIMITED	EE
Zinnat, 250 mg õhukese polümeerikattega tabletid	UK/H/5462/04	226498	GLAXO GROUP LIMITED	EE
Zinnat, 250 mg, tabletki powlekane	UK/H/5462/004	R/0833	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 250 mg, tabletki powlekane	UK/H/5462/004	R/0833	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 250 mg/5 ml, granulat do sporządzania zawiesiny doustnej	UK/H/5462/02	4688	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 250 mg/5 ml, granulat do sporządzania zawiesiny doustnej	UK/H/5462/02	4688	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 500 mg õhukese polümeerikattega tabletid	UK/H/5462/05	226598	GLAXO GROUP LIMITED	EE
Zinnat, 500 mg õhukese polümeerikattega tabletid	UK/H/5462/05	226598	GLAXO GROUP LIMITED	EE
Zinnat, 500 mg, tabletki powlekane	UK/H/5462/005	R/0834	GLAXOSMITHKLINE EXPORT LTD	PL
Zinnat, 500 mg, tabletki powlekane	UK/H/5462/005	R/0834	GLAXOSMITHKLINE EXPORT LTD	PL
Zipos 125 mg/5 mL granulado para suspensão oral	not available	4690988	INSTITUTO LUSO-FÁRMACO, LDA	PT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zipos 125 mg/5 mL granulado para suspensão oral	not available	8796904	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 125 mg/5 mL granulado para suspensão oral	not available	4690988	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 125 mg/5 mL granulado para suspensão oral	not available	8796904	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	4691382	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	8767210	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	5837083	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	4691481	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	4691382	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	8767210	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	5837083	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg comprimidos revestidos por película	not available	4691481	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg/5 mL granulado para suspensão oral	not available	4691085	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg/5 mL granulado para suspensão oral	not available	3322484	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg/5 mL granulado para suspensão oral	not available	4691085	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 250 mg/5 mL granulado para suspensão oral	not available	3322484	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	4683884	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	8767228	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	5841283	INSTITUTO LUSO-FÁRMACO, LDA	PT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zipos 500 mg comprimidos revestidos por película	not available	4683983	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	4683884	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	8767228	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	5841283	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zipos 500 mg comprimidos revestidos por película	not available	4683983	INSTITUTO LUSO-FÁRMACO, LDA	PT
Zoref 125 mg/5 ml granulado para suspensão oral	UK/H/5462/01	8780700	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 125 mg/5 mL granulado para suspensão oral	UK/H/5462/01	4684080	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 125 mg/5 ml granulado para suspensão oral	UK/H/5462/01	8780700	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 125 mg/5 mL granulado para suspensão oral	UK/H/5462/01	4684080	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	5811187	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	4684486	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	8720417	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	4684585	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	5811187	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	4684486	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	8720417	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg comprimidos revestidos por película	UK/H/5462/04	4684585	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg/5 mL granulado para suspensão oral	UK/H/5462/02	3090685	GLAXO WELLCOME FARMACÊUTICA, LDA	PT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoref 250 mg/5 mL granulado para suspensão oral	UK/H/5462/02	4684189	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg/5 mL granulado para suspensão oral	UK/H/5462/02	3090685	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 250 mg/5 mL granulado para suspensão oral	UK/H/5462/02	4684189	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	5811286	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	4684684	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	8720425	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	4684783	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	5811286	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	4684684	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Zoref 500 mg comprimidos revestidos por película	UK/H/5462/05	8720425	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Зинат 125 mg/5 ml гранули за перорална суспензия	UK/H/5462/01	20020076	GLAXOSMITHKLINE EOOD	BG
Зинат 125 mg/5 ml гранули за перорална суспензия	UK/H/5462/01	20020076	GLAXOSMITHKLINE EOOD	BG
Зинат 250 mg филмирани таблетки	UK/H/5462/04	20020074	GLAXOSMITHKLINE EOOD	BG
Зинат 250 mg филмирани таблетки	UK/H/5462/04	20020074	GLAXOSMITHKLINE EOOD	BG
Зинат 250 mg/5 ml гранули за перорална суспензия	UK/H/5462/02	20060231	GLAXOSMITHKLINE EOOD	BG
Зинат 250 mg/5 ml гранули за перорална суспензия	UK/H/5462/02	20060231	GLAXOSMITHKLINE EOOD	BG
Зинат 500 mg филмирани таблетки	UK/H/5462/05	20020075	GLAXOSMITHKLINE EOOD	BG
Зинат 500 mg филмирани таблетки	UK/H/5462/05	20020075	GLAXOSMITHKLINE EOOD	BG