



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

24 January 2018
EMA/270645/2015
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: azithromycin (systemic use formulations)

Procedure no.: PSUSA/00010491/201704



Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
3Z, 150 mg, pó para suspensão oral	not available	4453486	JABA RECORDATI, S.A.	PT
3Z, 200 mg, pó para suspensão oral	not available	4453585	JABA RECORDATI, S.A.	PT
3Z, 250 mg, pó para suspensão oral	not available	4453684	JABA RECORDATI, S.A.	PT
3Z, 500 mg, pó para suspensão oral	not available	5367982	JABA RECORDATI, S.A.	PT
3Z, 500 mg, pó para suspensão oral	not available	4453783	JABA RECORDATI, S.A.	PT
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 362 7 0	PFIZER HOLDING FRANCE	FR
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 361 0 2	PFIZER HOLDING FRANCE	FR
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 336 6 8	PFIZER HOLDING FRANCE	FR
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 338 9 7	PFIZER HOLDING FRANCE	FR
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 363 3 1	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
AZADOSE 600 mg, comprimé pelliculé	not available	34009 343 337 2 9	PFIZER HOLDING FRANCE	FR
Azatril 250 mg, film-coated tablets	NL/H/2400/001-002	5215/2012/01-04	ACTAVIS GROUP PTC EHF.	RO
Azibiot 20 mg/ml milteliai geriamajai suspensijai	SK/H/0148/003	LT/1/14/3600/003	KRKA, D.D., NOVO MESTO	LT
Azibiot 20 mg/ml pulbere pentru suspensie orală	SK/H/0148/003	9933/2017/01	KRKA, D.D., NOVO MESTO	RO
Azibiot NEO 20 mg/ml prášok na perorálnu suspenziu	SK/H/0148/003	15/0001/17-S	KRKA, D.D., NOVO MESTO	SK
Azibiot, 20 mg/ml, proszek do sporządzania zawiesiny doustnej	SK/H/0148/003	23963	KRKA, D.D., NOVO MESTO	PL
Azithromycin 250 mg Capsules	not available	PL 42357/0224	AMNEAL PHARMA EUROPE LIMITED	UK
Azithromycin 250mg Tablets	AT/H/0266/001	PL 04416/1235	SANDOZ LTD	UK
Azithromycin 500mg Tablets	DE/H/1903/002	PL 04416/0668	SANDOZ LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Azithromycin Actavis 250 mg filmom obalené tablety	NL/H/2400/001	15/0480/12-S	ACTAVIS GROUP PTC EHF.	SK
Azithromycin Aurovitas 250 mg potahované tablety	PT/H/1475/001	15/481/16-C	AUROVITAS UNIPessoal, LDA.	CZ
Azithromycin Aurovitas 500 mg potahované tablety	PT/H/1475/002	15/482/16-C	AUROVITAS UNIPessoal, LDA.	CZ
Azithromycin Hexal 250 mg - Filmtabletten	AT/H/0266/001	1-25583	HEXAL PHARMA GMBH	AT
Azithromycin Krka 20 mg/ml pulveris iekšķīgi lietojamas suspensijas pagatavošanai	SK/H/0148/003	17-0032	KRKA, D.D., NOVO MESTO	LV
Azithromycin Krka, 20 mg/ml suukaudse suspensiooni pulber	SK/H/0148/003	933217	KRKA, D.D., NOVO MESTO	EE
Azithromycin Sandoz 100 mg/ 5 ml prášok na perorálnu suspenziu	NL/H/0886/001	15/0181/07-S	SANDOZ PHARMACEUTICALS D.D.	SK
Azithromycin Sandoz 500 mg – Filmtabletten	DE/H/1903/002	1-26352	SANDOZ GMBH	AT
Azithromycin Sandoz 500 mg - Granulat zur Herstellung einer	AT/H/0390/003	1-31326	SANDOZ GMBH	AT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Suspension zum Einnehmen auf einem Messlöffel				
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/007	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/010	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/008	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/009	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/011	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/012	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/013	SANDOZ GMBH	LT
Azithromycin Sandoz 500 mg plèvele dengtos tabletès	DE/H/1903/002	LT/1/05/0323/014	SANDOZ GMBH	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
AZITHROMYCINE MONODOSE SANDOZ 250 mg, comprimé pelliculé	not available	387 628-2	SANDOZ	FR
AZITHROMYCINE SANDOZ 250 mg, comprimé pelliculé	not available	379 352-1	SANDOZ	FR
AZITHROMYCINE SANDOZ 250 mg, comprimé pelliculé	not available	570 753-7	SANDOZ	FR
AZITHROMYCINE SANDOZ 250 mg, comprimé pelliculé	not available	379 353-8	SANDOZ	FR
AZITHROMYCINE SANDOZ 250 mg, comprimé pelliculé	not available	570 754-3	SANDOZ	FR
Azitromax 2 g depotgranulat till oral suspension	not available	23285	PFIZER AB	SE
Azitromax 250 mg filmdragerade tabletter	not available	12582	PFIZER AB	SE
Azitromax 40 mg/ml pulver till oral suspension	not available	11609	PFIZER AB	SE
Azitromax 40 mg/ml, pulver til mikstur, suspensjon	not available	8025	PFIZER AS	NO

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Azitromax 500 mg filmdragerade tabletter	not available	12583	PFIZER AB	SE
Azitromax 500 mg tabletter	not available	95-245	PFIZER AS	NO
Azitromax 500 mg, pulver till koncentrat till infusionsvätska, lösning	not available	16698	PFIZER AB	SE
Azitromax 600 mg filmdragerade tabletter	not available	13233	PFIZER AB	SE
Azitromicin Krka 20 mg/ml prašek za peroralno suspenzijo	SK/H/0148/003	H/14/00246/006	KRKA, D.D., NOVO MESTO	SI
AZITROMICINA PFIZER 100 mg polvere per sospensione orale	not available	027897065	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 150 mg polvere per sospensione orale	not available	027897077	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 200 mg polvere per sospensione orale	not available	027897089	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 200 mg/5 ml polvere per sospensione orale	not available	027897053	PFIZER 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
AZITROMICINA PFIZER 200 mg/5 ml polvere per sospensione orale	not available	027897026	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 250 mg capsule rigide	not available	027897014	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 300 mg polvere per sospensione orale	not available	027897091	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 400 mg polvere per sospensione orale	not available	027897103	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 500 mg compresse rivestite con film	not available	027897040	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER 500 mg polvere per soluzione per infusione	not available	027897127	PFIZER ITALIA S.R.L.	IT
AZITROMICINA PFIZER Avium 600 mg compresse rivestite con film	not available	027897115	PFIZER ITALIA S.R.L.	IT
Azitromicină Sandoz 500 mg, granule pentru suspensie orală în linguriță unidoză	AT/H/0390/003	4814/2012/01	S.C. SANDOZ S.R.L.	RO
Azitromicina Teva 250 mg compresse rivestite con film	NL/H/0614/001	037555315	TEVA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Azitromicina Teva 250 mg compresse rivestite con film	NL/H/0614/001	037555291	TEVA ITALIA S.R.L.	IT
Azitromicina Teva 250 mg compresse rivestite con film	NL/H/0614/001	037555303	TEVA ITALIA S.R.L.	IT
Azitromicina Teva 250 mg compresse rivestite con film	NL/H/614/001	037555289	TEVA ITALIA S.R.L.	IT
Azitromicina TEVA 250 mg comprimidos recubiertos con película EFG	NL/H/0614/001	67335	TEVA PHARMA S.L.U	ES
Azitromycine Sandoz 100 mg/5 ml, poeder voor orale suspensie	NL/H/0886/001	RVG 32016	SANDOZ B.V.	NL
ORDIPHA 500 mg, comprimé dispersible sécable	not available	395347-9	TONIPHARM	FR
ORDIPHA 500 mg, comprimé dispersible sécable	not available	395347-9	TONIPHARM	FR
Ribotrex 200 mg/5 ml polvere per sospensione orale	not available	028177020	PIERRE FABRE PHARMA S.R.L.	IT
Ribotrex 500 mg compresse rivestite con film	not available	028177032	PIERRE FABRE 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
SUMAMED 100 mg/5 ml milteliai geriamajai suspensijai	not available	LT/1/94/0529/005	TEVA PHARMA B.V.	LT
Sumamed 100 mg/5 ml por sziruphoz	not available	OGYI-T-5272/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Sumamed 100 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-129408559	PLIVA HRVATSKA D.O.O.	HR
Sumamed 100 mg/5 ml pulveris iekšķīgi lietojamas suspensijas pagatavošanai	not available	94-0230	TEVA PHARMA B.V.	LV
Sumamed 100 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-5272/07	TEVA GYÓGYSZERGYÁR ZRT	HU
Sumamed 1200, 200 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-406709782	PLIVA HRVATSKA D.O.O.	HR
Sumamed 125 mg comprimate filmate	not available	8765/2016/01	PLIVA LJUBLJANA D.O.O.;	RO
Sumamed 125 mg filmom obalené tablety	not available	15/0351/92-C/S	TEVA PHARMACEUTICALS SLOVAKIA S.R.O	SK
Sumamed 125 mg filmom obložene tablete	not available	HR-H-844869851	PLIVA HRVATSKA D.O.O.	HR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SUMAMED 125 mg filmsko obložene tablete	not available	H/92/01472/001	PLIVA LJUBLJANA D.O.O.;	SI
SUMAMED 125 mg potahované tablety	not available	15-351-92-A-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Sumamed 1500, 200 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-096496463	PLIVA CROATIA LTD	HR
Sumamed 250 mg capsule	not available	730/2008/01	PLIVA LJUBLJANA D.O.O.;	RO
Sumamed 250 mg cietās kapsulas	not available	94-0231	TEVA PHARMA B.V.	LV
SUMAMED 250 mg kapsule	not available	HR-H-304657205	TEVA B.V	HR
Sumamed 250 mg kemény kapszula	not available	OGYI-T-5272/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Sumamed 250 mg kietosios kapsulės	not available	LT/1/94/0529/002	TEVA PHARMA B.V.	LT
SUMAMED 250 mg trde kapsule	not available	H/92/01472/002	PLIVA LJUBLJANA D.O.O.;	SI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SUMAMED 40 mg/ml prašek za peroralno suspenzijo	not available	H/92/01472/006	PLIVA LJUBLJANA D.O.O.;	SI
Sumamed 500 mg apvalkotas tabletes	not available	00-0863	TEVA PHARMA B.V.	LV
Sumamed 500 mg comprimate filmate	not available	8766/2016/01	PLIVA LJUBLJANA D.O.O.;	RO
Sumamed 500 mg filmom obalené tablety	not available	15/0351/92-C/S	TEVA PHARMACEUTICALS SLOVAKIA S.R.O	SK
Sumamed 500 mg filmom obložene tablete	not available	HR-H-425760378	PLIVA CROATIA LTD	HR
SUMAMED 500 mg filmsko obložene tablete	not available	H/92/01472/003	PLIVA LJUBLJANA D.O.O.;	SI
Sumamed 500 mg filmtabletta	not available	OGYI-T-5272/05	TEVA GYÓGYSZERGYÁR ZRT	HU
SUMAMED 500 mg infuze prášek pro infuzní roztok	not available	15/223/03-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Sumamed 500 mg plėvele dengtos tabletės	not available	LT/1/94/0529/004	TEVA PHARMA B.V.	LT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sumamed 500 mg plėvele dengtos tabletės	not available	LT/1/94/0529/003	TEVA PHARMA B.V.	LT
SUMAMED 500 mg potahované tablety	not available	15/351/92-B/C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Sumamed 500 mg prašak za koncentrat za otopinu za infuziju	not available	HR-H-261024118	PLIVA HRVATSKA D.O.O.	HR
SUMAMED 500 mg prašek za raztopino za infundiranje	not available	H/92/01472/007	PLIVA LJUBLJANA D.O.O.;	SI
Sumamed forte 200 mg/5 ml milteliai geriamajai suspensijai	not available	LT/1/94/0529/007	TEVA PHARMA B.V.	LT
Sumamed forte 200 mg/5 ml milteliai geriamajai suspensijai	not available	LT/1/94/0529/006	TEVA PHARMA B.V.	LT
Sumamed forte 200 mg/5 ml milteliai geriamajai suspensijai	not available	LT/1/94/0529/008	TEVA PHARMA B.V.	LT
Sumamed forte 200 mg/5 ml por sziruphoz	not available	OGYI-T-5272/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Sumamed forte 200 mg/5 ml por sziruphoz	not available	OGYI-T-5272/08	TEVA GYÓGYSZERGYÁR ZRT	HU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sumamed forte 200 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-096865641	PLIVA HRVATSKA D.O.O.	HR
Sumamed forte 200 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-096865641	PLIVA HRVATSKA D.O.O.	HR
Sumamed Forte 200 mg/5 ml pulbere pentru suspensie orală	not available	8767/2016/01-02-03	PLIVA LJUBLJANA D.O.O.;	RO
Sumamed forte 200 mg/5 ml pulveris iekškigi lietojamas suspensijas pagatavošanai	not available	00-0864	TEVA PHARMA B.V.	LV
Sumamed forte prášok na sirup	not available	15/0352/92-C/S	TEVA PHARMACEUTICALS SLOVAKIA S.R.O	SK
SUMAMED forte sirup prášek pro perorální suspenzi	not available	15/352/92-B/C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
SUMAMED FORTE, 200 mg/5 ml suukaudse suspensiooni pulber	not available	357201	TEVA PHARMA B.V.	EE
SUMAMED forte, 200 mg/5 ml, proszek do sporządzania zawiesiny doustnej	not available	7422	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sumamed prášok na sirup	not available	15/0352/92-C/S	TEVA PHARMACEUTICALS SLOVAKIA S.R.O	SK
Sumamed S 500 mg filmom obložene tablete	not available	HR-H-732776477	PLIVA HRVATSKA D.O.O.	HR
SUMAMED S 500 mg filmsko obložene tablete	not available	H/92/01472/004	PLIVA LJUBLJANA D.O.O.;	SI
Sumamed S 500 mg filmtableta	not available	OGYI-T-5272/06	TEVA GYÓGYSZERGYÁR ZRT	HU
SUMAMED sirup prášek pro perorální suspenzi	not available	15/352/92-A/C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
SUMAMED STD potahované tablety	not available	15/249/02-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Sumamed tvrdé kapsuly	not available	15/0226/90-C/S	TEVA PHARMACEUTICALS SLOVAKIA S.R.O	SK
SUMAMED tvrdé tobolky	not available	15/226/90-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
SUMAMED za otroke 20 mg/ml prašek za peroralno suspenzijo	not available	H/92/01472/005	PLIVA LJUBLJANA D.O.O.;	SI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SUMAMED, 100 mg/5 ml suukaudse suspensiooni pulber	not available	357101	TEVA PHARMA B.V.	EE
SUMAMED, 100 mg/5 ml, proszek do sporządzania zawiesiny doustnej	not available	7421	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Sumamed, 125 mg, tabletki powlekane	not available	8462	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
SUMAMED, 250 mg kõvakapslid	not available	292599	TEVA PHARMA B.V.	EE
Sumamed, 250 mg, kapsułki twarde	not available	7423	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
SUMAMED, 500 mg õhukese polümeerikattega tabletid	not available	292499	TEVA PHARMA B.V.	EE
SUMAMED, 500 mg, proszek do sporządzania roztworu do infuzji	not available	9016	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Sumamed, 500 mg, tabletki powlekane	not available	7424	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
TROZOCINA 2 g granulato per sospensione orale a rilascio prolungato	not available	027948102	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE 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
TROZOCINA 200 mg/5 ml polvere per sospensione orale	not available	027948052	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
TROZOCINA 200 mg/5 ml polvere per sospensione orale	not available	027948076	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
TROZOCINA 250 mg capsule rigide	not available	027948049	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
TROZOCINA 500 mg compresse rivestite con film	not available	027948064	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
TROZOCINA 500 mg polvere per soluzione per infusione	not available	027948090	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
TROZOCINA Avium 600 mg compresse rivestite con film	not available	027948088	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
Ultreon® 600 mg Filmtabletten	not available	37344.00.00	PFIZER PHARMA PFE GMBH	DE
Zetamax, 2 g, granulati o przedłużonym uwalnianiu do sporządzania zawiesiny doustnej	PT/H/0146/001	15609	PFIZER EUROPE MA EEIG	PL
ZITHROMAX	not available	19592	PFIZER HELLAS, A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zithromax 2 g granulado de libertação prolongada para suspensão oral	PT/H/0146/001	5017827	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 2 g granulado de libertação prolongada para suspensão oral	PT/H/0146/001	5017827	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 200 mg in 5 ml suspension	not available	MA505/03901	PFIZER HELLAS, A.E.	MT
Zithromax 200 mg in 5 ml suspension	not available	PL 00057/0336	PFIZER LIMITED	UK
Zithromax 200 mg/5 ml - Trockensaft	not available	1-20313	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
ZITHROMAX 200 mg/5 ml κόνις για πόσιμο εναιώρημα	not available	41167/10/31-05-2011	PFIZER HELLAS, A.E.	GR
ZITHROMAX 200 mg/5ml κόνις για πόσιμο εναιώρημα	not available	16336	PFIZER HELLAS, A.E.	CY
ZITHROMAX 250 mg CAPSULES	not available	MA505/03902	PFIZER HELLAS, A.E.	MT
ZITHROMAX 250 mg CAPSULES	not available	PL 00057/0335	PFIZER LIMITED	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZITHROMAX 250 mg filmdragerade tabletter azitromycin	not available	12016	PFIZER OY	FI
Zithromax 250 mg tabletti, kalvopäällysteinen	not available	12016	PFIZER OY	FI
ZITHROMAX 250 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	44407/5-11-2009	PFIZER HELLAS, A.E.	GR
ZITHROMAX 250 mg καψάκια σκληρά	not available	44405/5-11-2009	PFIZER HELLAS, A.E.	GR
ZITHROMAX 250 mg, comprimé pelliculé	not available	34009 351 774 9 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZITHROMAX 250 mg, comprimé pelliculé	not available	34009 351 773 2 2	PFIZER HOLDING FRANCE (S.C.A.)	FR
Zithromax 250, tabletten 250 mg	not available	RVG 19432	PFIZER B.V.	NL
ZITHROMAX 40 mg/ml ENFANTS, poudre pour suspension buvable	not available	34009 356 564 2 1	PFIZER HOLDING FRANCE	FR
ZITHROMAX 40 mg/ml ENFANTS, poudre pour suspension buvable	not available	34009 356 565 9 9	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZITHROMAX 40 mg/ml ENFANTS, poudre pour suspension buvable	not available	34009 356 561 3 1	PFIZER HOLDING FRANCE	FR
ZITHROMAX 40 mg/ml ENFANTS, poudre pour suspension buvable	not available	34009 356 563 6 0	PFIZER HOLDING FRANCE	FR
Zithromax 40 mg/ml jauhe oraalisuspensiota varten	not available	11615	PFIZER OY	FI
Zithromax 40 mg/ml pó para suspensão oral	not available	3969581	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 40 mg/ml pó para suspensão oral	not available	4576286	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 40 mg/ml pó para suspensão oral	not available	2248284	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 40 mg/ml pó para suspensão oral	not available	2248383	LABORATÓRIOS PFIZER, LDA.	PT
ZITHROMAX 40 mg/ml pulver till oral suspension azitromycin	not available	11615	PFIZER OY	FI
Zithromax 500 mg - Filmtabletten	not available	1-21939	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zithromax 500 mg comprimido revestido por película	not available	4576385	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax 500 mg comprimido revestido por película	not available	2521284	LABORATÓRIOS PFIZER, LDA.	PT
ZITHROMAX 500 mg filmdragerade tabletter azitromycin	not available	12017	PFIZER OY	FI
Zithromax 500 mg tabletti, kalvopäällysteinen	not available	12017	PFIZER OY	FI
ZITHROMAX 500 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	44409/5-11-2009	PFIZER HELLAS, A.E.	GR
Zithromax 500, tabletten 500 mg	not available	RVG 19433	PFIZER B.V.	NL
Zithromax Capsules 250 mg	not available	PA 19/47/1	PFIZER LIMITED	IE
Zithromax i.v. 500 mg – Pulver zur Herstellung einer Infusionslösung	not available	1-24636	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Zithromax IV 500 mg infuusiokuiva-aine, liuosta varten	not available	16152	PFIZER OY	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zithromax IV 500 mg pó para solução para perfusão	not available	3496982	LABORATÓRIOS PFIZER, LDA.	PT
Zithromax IV 500 mg pó para solução para perfusão	not available	5462569	LABORATÓRIOS PFIZER, LDA.	PT
ZITHROMAX IV 500 mg pulver till infusionsvätska, lösning azitromycin	not available	16152	PFIZER OY	FI
ZITHROMAX MONODOSE 250 mg, comprimé pelliculé	not available	34009 351 777 8 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZITHROMAX MONODOSE 250 mg, comprimé pelliculé	not available	34009 351 778 4 1	PFIZER HOLDING FRANCE (S.C.A.)	FR
Zithromax Powder for Oral Suspension 200 mg/5 ml	not available	PA 19/47/2	PFIZER LIMITED	IE
ZITHROMAX, 500mg/φιαλίδιο, κόνις για διάλυμα προς έγχυση	not available	44411/5-11-2009	PFIZER HELLAS, A.E.	GR
Zithromax, suspensie (poeder voor) 200 mg/5 ml	not available	RVG 14999	PFIZER B.V.	NL
Zithromax® 250 mg Filmtabletten	not available	37174.00.00	PFIZER PHARMA PFE GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zithromax® 500 mg Filmtabletten	not available	37174.01.00	PFIZER PHARMA PFE GMBH	DE
Zithromax® i.v. 500 mg Pulver zur Herstellung einer Infusionslösung	not available	37174.00.01	PFIZER PHARMA PFE GMBH	DE
Zithromax® Trockensaft 200 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	not available	25156.00.01	PFIZER PHARMA PFE GMBH	DE
Zitinn 250 mg filmtabletta	NL/H/2400/001	OGYI-T-22373/01-04	ACTAVIS GROUP PTC EHF.	HU
Zitrocin 100 mg/5 ml por sziruphoz	not available	OGYI-T-9027/03	TEVA MAGYARORSZÁG ZRT	HU
Zitrocin 250 mg kemény kapszula	not available	OGYI-T-9027/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Zitrocin 500 mg filmtabletta	not available	OGYI-T-9027/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Zitrocin forte 200 mg/5 ml por sziruphoz	not available	OGYI-T-9027/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Zitrocin forte 200 mg/5 ml por sziruphoz	not available	OGYI-T-9027/07	TEVA GYÓGYSZERGYÁR ZRT	HU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zitrocin S 500 mg filmtabletta	not available	OGYI-T-9027/06	TEVA GYÓGYSZERGYÁR ZRT	HU
ZITROMAX 100 mg polvere per sospensione orale	not available	027860081	PFIZER ITALIA S.R.L.	IT
ZITROMAX 1000 mg polvo para suspensión oral en sobre	not available	60.065	PFIZER, S.L.	ES
ZITROMAX 150 mg polvere per sospensione orale	not available	027860093	PFIZER ITALIA S.R.L.	IT
ZITROMAX 2 g granulato per sospensione orale a rilascio prolungato	not available	027860168	PFIZER ITALIA S.R.L.	IT
ZITROMAX 200 mg polvere per sospensione orale	not available	027860105	PFIZER ITALIA S.R.L.	IT
ZITROMAX 200 mg/5 ml Poeder voor orale suspensie	not available	BE165961	PFIZER S.A. (BELGIUM)	BE
ZITROMAX 200 mg/5 ml polvere per sospensione orale	not available	027860055	PFIZER ITALIA S.R.L.	IT
ZITROMAX 200 mg/5 ml polvere per sospensione orale	not available	027860079	PFIZER 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
ZITROMAX 200 mg/5 ml polvere per sospensione orale	not available	027860067	PFIZER ITALIA S.R.L.	IT
ZITROMAX 200 mg/5 ml polvere per sospensione orale	not available	027860028	PFIZER ITALIA S.R.L.	IT
ZITROMAX 200 mg/5 ml polvo para suspensión oral en frasco	not available	59.615	PFIZER, S.L.	ES
ZITROMAX 200 mg/5 ml Poudre pour suspension buvable	not available	2011111331	PFIZER S.A. (BELGIUM)	LU
Zitromax 200 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	not available	BE165961	PFIZER S.A. (BELGIUM)	BE
Zitromax 200 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen Azithromycin	not available	2011111331	PFIZER S.A. (BELGIUM)	LU
ZITROMAX 200 mg/5ml Poudre pour suspension buvable	not available	BE165961	PFIZER S.A. (BELGIUM)	BE
ZITROMAX 250 mg cápsulas duras	not available	59.616	PFIZER, S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZITROMAX 250 mg capsule rigide	not available	027860016	PFIZER ITALIA S.R.L.	IT
ZITROMAX 250 mg Comprimés pelliculés	not available	BE193261	PFIZER S.A. (BELGIUM)	BE
ZITROMAX 250 mg Comprimés pelliculés	not available	1998080024	PFIZER S.A. (BELGIUM)	LU
ZITROMAX 250 mg Filmomhulde tabletten	not available	BE193261	PFIZER S.A. (BELGIUM)	BE
Zitromax 250 mg Filmtabletten	not available	BE193261	PFIZER S.A. (BELGIUM)	BE
Zitromax 250 mg Filmtabletten Azithromycin	not available	1998080024	PFIZER S.A. (BELGIUM)	LU
Zitromax 250 mg filmuhúðaðar töflur	not available	970281	PFIZER APS	IS
ZITROMAX 250 mg polvo para suspensión oral en sobre	not available	59.620	PFIZER, S.L.	ES
ZITROMAX 300 mg polvere per sospensione orale	not available	027860117	PFIZER 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
Zitromax 40 mg/ml mixtúrduft, dreifa	not available	920013	PFIZER APS	IS
ZITROMAX 400 mg polvere per sospensione orale	not available	027860129	PFIZER ITALIA S.R.L.	IT
ZITROMAX 500 mg compresse rivestite con film	not available	027860042	PFIZER ITALIA S.R.L.	IT
ZITROMAX 500 mg Comprimés pelliculés	not available	BE193243	PFIZER S.A. (BELGIUM)	BE
ZITROMAX 500 mg Comprimés pelliculés	not available	2011111333	PFIZER S.A. (BELGIUM)	LU
ZITROMAX 500 mg comprimidos recubiertos con película	not available	61.272	PFIZER, S.L.	ES
ZITROMAX 500 mg Filmomhulde tabletten	not available	BE193243	PFIZER S.A. (BELGIUM)	BE
Zitromax 500 mg Filmtabletten	not available	BE193243	PFIZER S.A. (BELGIUM)	BE
Zitromax 500 mg Filmtabletten Azithromycin	not available	2011111333	PFIZER S.A. (BELGIUM)	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zitromax 500 mg filmuhúðaðar töflur	not available	970282	PFIZER APS	IS
Zitromax 500 mg innrennslisstofn, lausn	not available	IS/1/02/106/01	PFIZER APS	IS
ZITROMAX 500 mg polvere per soluzione per infusione	not available	027860156	PFIZER ITALIA S.R.L.	IT
ZITROMAX 500 mg polvo para solución para perfusión	not available	64.834	PFIZER, S.L.	ES
ZITROMAX 500 mg polvo para suspensión oral en sobre	not available	60.066	PFIZER, S.L.	ES
ZITROMAX 600 mg Comprimés pelliculés	not available	BE193252	PFIZER S.A. (BELGIUM)	BE
ZITROMAX 600 mg Comprimés pelliculés	not available	2011111334	PFIZER S.A. (BELGIUM)	LU
ZITROMAX 600 mg Filmomhulde tabletten	not available	BE193252	PFIZER S.A. (BELGIUM)	BE
Zitromax 600 mg Filmtabletten	not available	BE193252	PFIZER S.A. (BELGIUM)	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
Zitromax 600 mg Filmtabletten Azithromycin	not available	2011111334	PFIZER S.A. (BELGIUM)	LU
ZITROMAX Avium 600 mg comprese rivestite con film	not available	027860143	PFIZER ITALIA S.R.L.	IT
Zitromax UNO forðakyrni fyrir mixtúru, dreifu	not available	IS/1/05/134/01	PFIZER APS	IS
Zitromax UNO, depotgranulat til oral suspension	not available	37612	PFIZER APS	DK
Zitromax, filmovertrukne tabletter	not available	18479	PFIZER APS	DK
Zitromax, filmovertrukne tabletter	not available	19118	PFIZER APS	DK
Zitromax, filmovertrukne tabletter	not available	19117	PFIZER APS	DK
Zitromax, Pulver til koncentrat til infusionsvæske, opløsning	not available	32123	PFIZER APS	DK
Zitromax, pulver til oral suspension	not available	14083	PFIZER APS	DK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zmax 2 g retard granulátum belsőleges szuszpenzióhoz	PT/H/0146/001	OGYI-T-20770/01	PFIZER KFT.	HU
Азибиот 20 mg/ml прах за перорална суспензия	SK/H/0148/003	20170121	KRKA, D.D., NOVO MESTO	BG
Зимакс 2 g гранули с удължено освобождаване за перорална суспензия	not available	20060293	PFIZER EUROPE MA EEIG	BG
Сумамед 250 mg капсули, твърди	not available	20000866	ECOPHARM GROUP	BG
СУМАМЕД 500 mg прах за инфузионен разтвор	not available	20010799	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Сумамед 500 mg филмирани таблетки	not available	20010175	ECOPHARM GROUP	BG
Сумамед сироп 100mg/5ml прах за перорална суспензия	not available	20000867	ECOPHARM GROUP	BG
Сумамед форте сироп 200mg/5ml прах за перорална суспензия	not available	20010176	ECOPHARM GROUP	BG