



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

29 January 2026
EMADOC-1700519818-2975609
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): cefixime

EURD list No. PSUSA/00000594/202506



Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SUPRAX 200 mg filmom obalené tablety	not available	15/0809/95-S	GEDEON RICHTER PLC.	SK
SUPRAX 200 mg filmom obalené tablety	not available	15/0809/95-S	GEDEON RICHTER PLC.	SK
SUPRAX 400 mg dispergovateľné tablety na perorálnu disperziu	not available	15/0858/10-S	GEDEON RICHTER PLC.	SK
SUPRAX 400 mg dispergovateľné tablety na perorálnu disperziu	not available	15/0858/10-S	GEDEON RICHTER PLC.	SK
SUPRAX 400 mg dispergovateľné tablety	not available	15/0858/10-S	GEDEON RICHTER PLC.	SK
CEFTORAL 200 mg/tab δισκία επικαλυμμένα με υμένιο	not available	39138/15-06-2010	VIANEX S.A.	GR
Tricef 200 mg Filmdabletten	AT/H/0132/001	1-19112	MERCK GESELLSCHAFT MBH	AT
Tricef 200 mg Filmdabletten	AT/H/0132/001	1-19112	MERCK GESELLSCHAFT MBH	AT
Tricef 200 mg Filmdabletten	AT/H/0132/001	1-19112	MERCK GESELLSCHAFT MBH	AT
Tricef 400 mg Filmdabletten	AT/H/0132/002	1-22077	MERCK GESELLSCHAFT MBH	AT
Tricef 400 mg Filmdabletten	AT/H/0132/002	1-22077	MERCK GESELLSCHAFT MBH	AT
Tricef 400 mg Filmdabletten	AT/H/0132/002	1-22077	MERCK GESELLSCHAFT MBH	AT
Tricef 200 mg Filmdabletten	AT/H/0132/001	1-19112	MERCK GESELLSCHAFT MBH	AT
Tricef 400 mg Filmdabletten	AT/H/0132/002	1-22077	MERCK GESELLSCHAFT MBH	AT
CEFTORAL, 400mg/TAB, Δισκία επικαλυμμένα με	not available	39515/15-02-2016	VIANEX S.A.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
υμέvio				
Tricef 100 mg/5 ml Trockensaft	not available	1-19111	MERCK GESELLSCHAFT MBH	AT
OROKEN 200 mg, comprimé pelliculé	not available	34009 332 147-2 2	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 311-3 5	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 313-6 4	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 314-2 5	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 315-9 3	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 316-5 4	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 317-1 5	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 556 462-9 7	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 556 463-5 8	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 556 461-2 9	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 331 318-8 3	AMDIPHARM LIMITED	FR
OROKEN 200 mg, comprimé pelliculé	not available	34009 556 460-6 8	AMDIPHARM LIMITED	FR
OROKEN NOURRISSONS 40 mg/5 ml, poudre pour suspension buvable en flacon	not available	34009 335 899-5 0	AMDIPHARM LIMITED	FR
OROKEN ENFANTS 100 mg/5 ml, poudre pour suspension buvable en	not available	34009 335 898-9 9	AMDIPHARM LIMITED	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
flacon				
Suprax 100 mg/5 ml por belsőleges szuszpenzióhoz	not available	OGYI-T-4926/06	GEDEON RICHTER PLC.	HU
Suprax 100 mg/5 ml por belsőleges szuszpenzióhoz	not available	OGYI-T-4926/02	GEDEON RICHTER PLC.	HU
SUPRAX 200 mg filmom obalené tablety	not available	15/0809/95-S	GEDEON RICHTER PLC.	SK
SUPRAX 200 mg filmom obalené tablety	not available	15/0809/95-S	GEDEON RICHTER PLC.	SK
SUPRACEF 400 mg compresse dispersibili	not available	028855070	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
SUPRACEF 400 mg compresse dispersibili	not available	028855068	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
SUPRACEF 100 mg/5 ml granulato per sospensione orale	not available	028855056	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
Suprax 200 mg filmtabletta	not available	OGYI-T-4926/01	GEDEON RICHTER PLC.	HU
SUPRAX 100 mg/5 ml prášok na perorálnu suspenziu	not available	15/0810/95-S	GEDEON RICHTER PLC.	SK
SUPRAX 100 mg/5 ml prášok na perorálnu suspenziu	not available	15/0810/95-S	GEDEON RICHTER PLC.	SK
Suprax 200mg Film-coated Tablets	not available	PA1142/042/001	AMDIPHARM LIMITED	IE
Suprax 200mg Film-coated Tablets.	not available	PA1142/042/001	AMDIPHARM LIMITED	IE
Suprax 200mg Film-coated Tablets.	not available	PA1142/042/001	AMDIPHARM LIMITED	IE
Suprax 200mg Film-	not available	PA1142/042/001	AMDIPHARM LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
coated Tablets.				
Denvar 200 mg cápsulas duras	not available	58.819	MERCK, S.L.	ES
Denvar 400 mg cápsulas duras	not available	60.469	MERCK, S.L.	ES
Denvar 100 mg/5 ml granulado para suspensión oral	not available	59.733	MERCK, S.L.	ES
Denvar 200 mg cápsulas duras	not available	58.819	MERCK, S.L.	ES
Denvar 200 mg cápsulas duras	not available	58.819	MERCK, S.L.	ES
Denvar 400 mg cápsulas duras	not available	60.469	MERCK, S.L.	ES
Denvar 100 mg/5 ml granulado para suspensión oral	not available	59.733	MERCK, S.L.	ES
Suprax 100 mg/5 ml por bolsóleges szuszpenzióhoz	not available	OGYI-T-4926/06	GEDEON RICHTER PLC.	HU
Suprax 100 mg/5 ml por bolsóleges szuszpenzióhoz	not available	OGYI-T-4926/02	GEDEON RICHTER PLC.	HU
CEFIXORAL 400 mg compresse dispersibili	not available	027134079	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
CEFIXORAL 400 mg compresse rivestite	not available	027134030	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
CEFIXORAL 100 mg/5 ml granulato per sospensione orale	not available	027134105	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
CEFIXORAL 400 mg compresse dispersibili	not available	027134081	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE -	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
			S.R.L.	
CEFTORAL, 100 mg/5ml, κόνις για πόσιμο εναιώρημα	not available	17283/15-06-2010	VIANEX S.A.	GR
UNIXIME 100 mg/5 ml granulato per sospensione orale	not available	027152091	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
UNIXIME 400 mg compresse dispersibili	not available	027152065	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
UNIXIME 400 mg compresse dispersibili	not available	027152077	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
UNIXIME 400 mg compresse rivestite.	not available	027152038	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
Suprax Tablets 200 mg	not available	PL 12762/0684	MERCURY PHARMACEUTICALS LTD.	XI
Suprax Tablets 200 mg	not available	PL 12762/0684	MERCURY PHARMACEUTICALS LTD.	XI
Suprax Tablets 200 mg	not available	PL 12762/0684	MERCURY PHARMACEUTICALS LTD.	XI
Suprax Tablets 200 mg	not available	PL 12762/0684	MERCURY PHARMACEUTICALS LTD.	XI
Tricef 20 mg/mL powder for oral suspension.	PT/H/2835/002	MA1550/00102	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	MT
Tricef 400 mg film-coated tablets.	PT/H/2835/001	MA1550/00101	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	MT
Tricef 400 mg film-coated tablets.	PT/H/2835/001	MA1550/00101	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	MT
Tricef 400 mg film-coated tablets.	PT/H/2835/001	MA1550/00101	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	MT
Tricef 400 mg film-coated tablets.	PT/H/2835/001	MA1550/00101	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tricef 20 mg/ml pó para suspensão oral	PT/H/2835/002	9725200	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	PT
Tricef 400 mg comprimidos revestidos por película	PT/H/2835/001	4549796	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	PT
Tricef 400 mg comprimidos revestidos por película	PT/H/2835/001	4549895	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	PT
Tricef 400 mg comprimidos revestidos por película	PT/H/2835/001	9725119	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	PT
Tricef 400 mg comprimidos revestidos por película	PT/H/2835/001	4549994	BIALPORT - PRODUTOS FARMACÊUTICOS, S.A.	PT