11 March 2021
EMA/165987/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: naproxen

Procedure no.: PSUSA/00002125/202008

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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
Actromadol 660 mg Comprimidos de liberación modificada	not available	83.348	BAYER HISPANIA SL	ES
Aktren Naproxen Filmtablette mit 220 mg Naproxen-Natrium	not available	38459.00.00	BAYER VITAL GMBH	DE
ALEVE – Filmtabletten	not available	1-22747	BAYER AUSTRIA GMBH	AT
ALEVE 220 mg comprimés pelliculés	not available	BE198213	BAYER SA NV	BE
ALEVE 220 mg comprimés pelliculés	not available	2000090015	BAYER SA NV	LU
ALEVE 220 mg filmomhulde tabletten	not available	BE198213	BAYER SA NV	BE
ALEVE 220 mg, comprimé pelliculé	not available	34009 274 295 8 3	BAYER HEALTHCARE	FR
ALEVE 220 mg, comprimé pelliculé	not available	34009 379 191 8 0	BAYER HEALTHCARE	FR
ALEVE 220 mg, comprimé pelliculé	not available	34009 492 050 7 3	BAYER HEALTHCARE	FR
ALEVE 220 mg, comprimé pelliculé	not available	34009 379 192 4 1	BAYER HEALTHCARE	FR
ALEVE 220 mg, comprimé pelliculé	not available	34009 380 481 6 2	BAYER HEALTHCARE	FR
ALEVE 220 mg, Filmtabletten	not available	BE 198213	BAYER SA NV	BE
ALEVE 220 mg, Filmtabletten	not available	2000090015	BAYER SA NV	LU
Aleve Classic bij pijn en koorts 220, omhulde tabletten	not available	RVG 19630	BAYER BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Aleve Extra 660 mg retard tabletta	not available	OGYI-T-8021/06	BAYER HUNGARIA KFT	HU
Aleve Extra 660 mg retard tabletta	not available	OGYI-T-8021/08	BAYER HUNGARIA KFT	HU
Aleve Extra 660 mg retard tabletta	not available	OGYI-T-8021/07	BAYER HUNGARIA KFT	HU
Aleve Feminax bij menstruatiepijn 275	not available	RVG 09006	BAYER BV	NL
Aleve filmtabletta	not available	OGYI-T-8021/03	BAYER HUNGARIA KFT	HU
Aleve filmtabletta	not available	OGYI-T-8021/04	BAYER HUNGARIA KFT	HU
Aleve filmtabletta	not available	OGYI-T-8021/05	BAYER HUNGARIA KFT	HU
Aleve filmtabletta	not available	OGYI-T-8021/01	BAYER HUNGARIA KFT	HU
Aleve filmtabletta	not available	OGYI-T-8021/02	BAYER HUNGARIA KFT	HU
Aleve Intense 550 mg, omhulde tablet.	not available	RVG 14484	BAYER BV	NL
Aleve Select 275 mg, omhulde tabletten	not available	RVG 09007	BAYER BV	NL
Aleve, 220 mg, tabletki powlekane	not available	7756	BAYER SP.Z.O.O	PL
Aleve® Filmtablette mit 220 mg Naproxen-Natrium	not available	38457.00.00	BAYER VITAL 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
ALEVETABS 220 mg, comprimé pelliculé	not available	34009 274 544 8 6	BAYER HEALTHCARE	FR
ALEVETABS 220 mg, comprimé pelliculé	not available	34009 274 286 9 2	BAYER HEALTHCARE	FR
ALEVETABS 220 mg, comprimé pelliculé	not available	34009 350 622 0 8	BAYER HEALTHCARE	FR
ALEVETABS 220 mg, comprimé pelliculé	not available	34009 350 621 4 7	BAYER HEALTHCARE	FR
ANAPROX® 550mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	128352/25-11-2019	MINERVA PHARMACEUTICAL S.A	GR
Antalgin 550 mg comprimidos recubiertos con película.	not available	57.152	ATNAHS PHARMA NETHERLANDS B.V.	ES
Apranax 275 mg comprimés pelliculés	not available	BE116837	ATNAHS PHARMA NETHERLANDS B.V.	BE
Apranax 275 mg comprimés pelliculés	not available	BE116837	ATNAHS PHARMA NETHERLANDS B.V.	LU
Apranax 275 mg filmomhulde tabletten	not available	BE116837	ATNAHS PHARMA NETHERLANDS B.V.	BE
Apranax 275 mg filmtabletta	not available	OGYI-T- 3742/01	BAUSCH HEALTH IRELAND LIMITED	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
APRANAX 275 mg, comprimé pelliculé	not available	34009 325 343 4 0	ATNAHS PHARMA UK LIMITED	FR
APRANAX 275 mg, comprimé pelliculé	not available	34009 325 002 2 2	ATNAHS PHARMA UK LIMITED	FR
APRANAX 275 mg, comprimé pelliculé	not available	34009 325 003 9 0	ATNAHS PHARMA UK LIMITED	FR
APRANAX 275 mg, comprimé pelliculé	not available	34009 325 004 5 1	ATNAHS PHARMA UK LIMITED	FR
APRANAX 275 mg, comprimé pelliculé	not available	34009 325 005 1 2	ATNAHS PHARMA UK LIMITED	FR
Apranax 550 mg comprimés pelliculés	not available	BE150997	ATNAHS PHARMA NETHERLANDS B.V.	BE
Apranax 550 mg comprimés pelliculés	not available	BE150997	ATNAHS PHARMA NETHERLANDS B.V.	LU
Apranax 550 mg filmomhulde tabletten	not available	BE150997	ATNAHS PHARMA NETHERLANDS B.V.	BE
Apranax 550 mg filmtabletta	not available	OGYI-T- 4030/01	BAUSCH HEALTH IRELAND LIMITED	HU
APRANAX 550 mg, comprimé pelliculé sécable	not available	34009 326 613 5 0	ATNAHS PHARMA UK LIMITED	FR
APRANAX 750 mg, comprimé	not available	34009 335 364 4 2	ATNAHS PHARMA UK 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
Dolormin für Frauen bei Menstruationsbeschwerden mit Naproxen	not available	35873.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin für Frauen bei Menstruationsbeschwerden mit Naproxen	not available	35873.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin für Frauen bei Menstruationsbeschwerden mit Naproxen	not available	35873.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin GS mit Naproxen	not available	35876.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin GS mit Naproxen	not available	35876.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin GS mit Naproxen 250 mg Tabletten	not available	35876.00.00	JOHNSON & JOHNSON GMBH	DE
Eox	IT/H/0110/001	33253	MYLAN DENMARK APS	DK
Galpharm Period Pain Relief 250mg Gastro-Resistant Tablets	not available	PL 16028/0144	GALPHARM HEALTHCARE LIMITED	UK
Laser 750 mg compresse a rilascio modificato	not available	023886082	EURO-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
Lasonil antinfiammatorio e antireumatico 220 mg compresse rivestite con film	not available	032790040	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 220 mg compresse rivestite con film	not available	032790038	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 220 mg compresse rivestite con film	not available	032790014	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 220 mg compresse rivestite con film	not available	032790026	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 660 mg compresse a rilascio modificato	not available	032790053	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 660 mg compresse a rilascio modificato	not available	032790065	BAYER SPA	IT
Lasonil antinfiammatorio e antireumatico 660 mg compresse a rilascio modificato	not available	032790077	BAYER 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
LUNDIRAN 250 mg cápsulas	not available	54.246	INDUSTRIA QUÍMICA Y FARMACEÚTICA VIR, S.A.	ES
Miranax 550 mg tabletti, kalvopäällysteinen	not available	10830	ATNAHS PHARMA UK LIMITED	FI
Miranax® 550 mg – Filmtabletten	not available	1-19930	GRÜNENTHAL GES. M.B.H.	AT
MOMENDOL 10% gel	IT/H/0110/002	025829197	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
MOMENDOL 10% gel	IT/H/0110/002	025829209	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
MOMENDOL 10% w/w GEL	IT/H/0110/002	34588/14-5-2012	ANGELINI PHARMA HELLAS S.A.	GR
MOMENDOL 10% w/w GEL	IT/H/0110/002	34588/14-5-2012	ANGELINI PHARMA HELLAS S.A.	GR
Momendol 100 mg/g (10%) gel	IT/H/0110/002	5359559	ANGELINI FARMACÊUTICA, LDA	PT
Momendol 100 mg/g (10%) gel	IT/H/0110/002	5359567	ANGELINI FARMACÊUTICA, LDA	PT
Momendol 220 mg capsule molli	not available	025829211	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. 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
Momendol 220 mg capsule molli	not available	025829223	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momendol 220 mg capsule molli	not available	025829235	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momendol 220 mg capsule molli	not available	025829247	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momendol 220 mg capsule molli	not available	025829250	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
MOMENDOL 220 mg compresse rivestite con film.	IT/H/0110/001	025829185	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
MOMENDOL 220 mg compresse rivestite con film.	IT/H/0110/001	025829084	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momendol 220 mg Filmtabletten	IT/H/0110/001	1-24912	ANGELINI PHARMA ÖSTERREICH GMBH	AT
Momendol 220 mg Filmtabletten	IT/H/0110/001	53767.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	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
MOMENDOL 220 mg granulato per soluzione orale.	not available	025829122	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momendol 220 mg επικαλυμμένα με λεπτό υμένιο δισκία	IT/H/0110/001	70434/11/31-8-2012	ANGELINI PHARMA HELLAS S.A.	GR
Momendol 220 mg επικαλυμμένα με λεπτό υμένιο δισκία	IT/H/0110/001	70434/11/31-8-2012	ANGELINI PHARMA HELLAS S.A.	GR
Momendol comprimidos revestidos por película	IT/H/0110/001	4031381	ANGELINI FARMACÊUTICA, LDA	PT
Momendol comprimidos revestidos por película	IT/H/0110/001	5814082	ANGELINI FARMACÊUTICA, LDA	PT
NAPREBEN 10% gel	not available	027669050	MERQURIO PHARMA S.R.L.	IT
NAPREBEN 550 mg capsule rigide	not available	027669011	MERQURIO PHARMA S.R.L.	IT
Napren-E 250 mg enterotabletter	not available	6998	TAKEDA AS	NO
Napren-E 375 mg enterotabletter	not available	95-1154	TAKEDA AS	NO
Napren-E 500 mg enterotabletter	not available	6999	TAKEDA AS	NO
NAPRIUS 10% GEL	not available	024667141	AESCULAPIUS FARMACEUTICI S.R.L.	IT
NAPRIUS 500 mg Compresse	not available	024667139	AESCULAPIUS FARMACEUTICI S.R.L.	IT
NAPRIUS 500 mg Granulato per sospensione orale	not available	024667154	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Naprometin 500 mg tabletti	not available	8765	ATNAHS PHARMA UK LIMITED	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
NAPROSSENE ANGELINI 220 mg compresse rivestite con film	not available	043800010	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
NAPROSSENE ANGELINI 220 mg compresse rivestite con film	not available	043800022	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Naprostad® 250 mg Filmtabletten	not available	904.01.01	STADA ARZNEIMITTEL AG	DE
Naprostad® 500 mg Filmtabletten	not available	13523.00.00	STADA ARZNEIMITTEL AG	DE
NAPROSYN 10 %, γέλη	not available	69908/16/24-5-2017	MINERVA PHARMACEUTICAL S.A	GR
Naprosyn 10% gel	not available	023177102	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Naprosyn 250 mg compresse gastroresistenti	not available	023177203	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Naprosyn 250 mg comprimidos	not available	4533899	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 250 mg comprimidos	not available	9382424	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 250 mg comprimidos	not available	4533998	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 250 mg comprimidos	not available	9382440	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 250 mg granulato per sospensione orale	not available	023177090	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA 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
NAPROSYN 250 mg, γαστροανθεκτικό δισκίο	not available	23554/12-3-14	MINERVA PHARMACEUTICAL S.A	GR
NAPROSYN 250 mg, δισκία	not available	60295/13/22-5-2014	MINERVA PHARMACEUTICAL S.A	GR
NAPROSYN 250 mg, υπόθετα	not available	79750/09/7-4-10	MINERVA PHARMACEUTICAL S.A	GR
Naprosyn 250mg Tablets	not available	PL 43252/0004	ATNAHS PHARMA UK LIMITED	UK
Naprosyn 500 mg compresse gastroresistenti	not available	023177215	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Naprosyn 500 mg comprimidos	not available	56.267	ATNAHS PHARMA UK LIMITED	ES
Naprosyn 500 mg comprimidos revestidos	not available	4161899	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 500 mg comprimidos revestidos	not available	9382432	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 500 mg comprimidos revestidos	not available	4534095	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 500 mg comprimidos revestidos	not available	9382457	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn 500 mg granulato per sospensione orale	not available	023177138	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Naprosyn 500 mg supposte	not available	023177088	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
NAPROSYN 500 mg, γαστροανθεκτικό δισκίο	not available	21622/12-3-14	MINERVA PHARMACEUTICAL S.A	GR
NAPROSYN 500 mg, δισκία	not available	45244/22-5-2014	MINERVA PHARMACEUTICAL S.A	GR
NAPROSYN 500 mg, υπόθετα	not available	21344/7-4-10	MINERVA PHARMACEUTICAL 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
Naprosyn 500mg Tablets	not available	PL 43252/0005	ATNAHS PHARMA UK LIMITED	UK
Naprosyn 750 mg compresse a rilascio modificato	not available	023177189	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Naprosyn EC 250 mg Gastro-Resistant Tablets	not available	PL 43252/0006	ATNAHS PHARMA UK LIMITED	UK
Naprosyn EC 250mg Gastro-resistant Tablets.	not available	PA 22657/0002/001	ATNAHS PHARMA NETHERLANDS B.V.	IE
Naprosyn EC 375mg Gastro-Resistant Tablets	not available	PL 43252/0007	ATNAHS PHARMA UK LIMITED	UK
Naprosyn EC 500 mg comprimidos gastrorresistentes	not available	4561585	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn EC 500 mg comprimidos gastrorresistentes	not available	8789826	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn EC 500 mg comprimidos gastrorresistentes	not available	4561684	ATNAHS PHARMA NETHERLANDS B.V.	PT
Naprosyn EC 500 mg comprimidos gastrorresistentes	not available	8789834	ATNAHS PHARMA NETHERLANDS B.V.	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
Naprosyn EC 500mg Gastro-Resistant Tablets	not available	PL 43252/0008	ATNAHS PHARMA UK LIMITED	UK
Naprosyn EC 500mg Gastro-resistant Tablets.	not available	PA 22657/0002/002	ATNAHS PHARMA NETHERLANDS B.V.	IE
Naprosyn Entero 250 mg enterotabletter	not available	11956	ATNAHS PHARMA UK LIMITED	SE
Naprosyn Entero 500 mg enterotabletter	not available	11946	ATNAHS PHARMA UK LIMITED	SE
NAPROSYNE 1000 mg, comprimé	not available	34009 334 499-3 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 316 844 4 2	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 316 846 7 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 316 843 8 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 321 767 4 8	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 318 482 2 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 321 766 8 7	LABORATOIRES GRÜNENTHAL S.A.S.	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
NAPROSYNE 250 mg, comprimé	not available	34009 316 845 0 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 250 mg, comprimé	not available	34009 318 481 6 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 500 mg, comprimé	not available	34009 350 241 7 6	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 500 mg, comprimé	not available	34009 323 810 4 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 500 mg, comprimé	not available	34009 331 389 2 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 500 mg, comprimé	not available	34009 323 809 6 1	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE 500 mg, comprimé	not available	34009 323 811 0 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Naprosyne Enteric Coated 250 mg comprimés gastro-résistants	not available	BE155565	ATNAHS PHARMA NETHERLANDS B.V.	BE
Naprosyne Enteric Coated 250 mg comprimés gastro-résistants	not available	BE155565	ATNAHS PHARMA NETHERLANDS B.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Naprosyne Enteric Coated 250 mg maagsapresistente tabletten	not available	BE155565	ATNAHS PHARMA NETHERLANDS B.V.	BE
Naprosyne Enteric Coated 500 mg comprimés gastro-résistants	not available	BE155583	ATNAHS PHARMA NETHERLANDS B.V.	BE
Naprosyne Enteric Coated 500 mg comprimés gastro-résistants	not available	BE155583	ATNAHS PHARMA NETHERLANDS B.V.	LU
Naprosyne Enteric Coated 500 mg maagsapresistente tabletten	not available	BE155583	ATNAHS PHARMA NETHERLANDS B.V.	BE
NAPROSYNE® 500 mg, suppositoire	not available	34009 320 505 6 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE® 500 mg, suppositoire	not available	34009 320 501 0 9	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE® 500 mg, suppositoire	not available	34009 320 502 7 7	LABORATOIRES GRÜNENTHAL S.A.S.	FR
NAPROSYNE® 500 mg, suppositoire	not available	34009 320 503 3 8	LABORATOIRES GRÜNENTHAL S.A.S.	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
Naproxen 250mg Tablets	not available	PL 36722/0052	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Naproxen 500mg Tablets	not available	PL 44041/0026	NOUMED LIFE SCIENCES	UK
Naproxen 500mg Tablets	not available	PL 36722/0053	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Naproxen AB 250 mg comprimés	NL/H/3472/001	BE531591	AUROBINDO PHARMA B.V.	BE
Naproxen AB 250 mg comprimés	NL/H/3472/001	BE531617	AUROBINDO PHARMA B.V.	BE
Naproxen AB 250 mg tabletten	NL/H/3472/001	BE531591	AUROBINDO PHARMA B.V.	BE
Naproxen AB 250 mg tabletten	NL/H/3472/001	BE531617	AUROBINDO PHARMA B.V.	BE
Naproxen AB 250 mg Tabletten	NL/H/3472/001	BE531591	AUROBINDO PHARMA B.V.	BE
Naproxen AB 250 mg Tabletten	NL/H/3472/001	BE531617	AUROBINDO PHARMA B.V.	BE
Naproxen AB 500 mg comprimés	NL/H/3472/002	BE531600	AUROBINDO PHARMA B.V.	BE
Naproxen AB 500 mg comprimés	NL/H/3472/002	BE531626	AUROBINDO PHARMA B.V.	BE
Naproxen AB 500 mg tabletten	NL/H/3472/002	BE531600	AUROBINDO PHARMA B.V.	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
Naproxen AB 500 mg tabletten	NL/H/3472/002	BE531626	AUROBINDO PHARMA B.V.	BE
Naproxen AB 500 mg Tabletten	NL/H/3472/002	BE531600	AUROBINDO PHARMA B.V.	BE
Naproxen AB 500 mg Tabletten	NL/H/3472/002	BE531626	AUROBINDO PHARMA B.V.	BE
Naproxen Orion 25 mg/ml oral suspension	FI/H/879/01/DC	PL 27925/0087	ORION CORPORATION	UK
Naproxen Orion 250 mg tabletit	NL/H/3472/001	33891	AUROBINDO PHARMA (MALTA) LIMITED	FI
Naproxen Saft InfectoPharm® 250 mg/5 ml Suspension zum Einnehmen	not available	17.522	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	AT
Naproxen Sandoz 250 mg tabletten	not available	BE274687	SANDOZ N.V.	BE
Naproxen Sandoz 500 mg tabletten	not available	BE273883	SANDOZ N.V.	BE
Naproxen Tablets BP 500mg.	not available	PL 21880/0111	MEDREICH PLC	UK
Naproxeno Aurobindo 250 mg comprimidos	NL/H/3472/001	80.805	LABORATORIOS AUROBINDO S.L.U.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Period Pain Reliever 250mg Gastro-resistant Tablets	not available	PL 20075/0619	ACCORD HEALTHCARE LIMITED	UK
Proxen 500 mg – Filmtabletten	not available	17231	GRÜNENTHAL GES. M.B.H.	AT
Synflex 275 mg capsule rigide	not available	024722011	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Synflex 550 mg compresse rivestite	not available	024722047	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Synflex 550 mg compresse rivestite	not available	024722112	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Synflex 550 mg granulato per sospensione orale	not available	024722086	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Момендол 220 mg филмирани таблетки	not available	20050220	ANGELINI PHARMA BULGARIA EOOD	BG
Момендол 220 mg филмирани таблетки	not available	20050220	ANGELINI PHARMA BULGARIA EOOD	BG