15 October 2020
EMA/649664/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ibuprofen, ibuprofen lysine (not indicated in ductus arteriosus), ibuprofen / caffeine

Procedure no.: PSUSA/00010649/202002

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
SPEDIFEN 400 mg, comprimé pelliculé	not available	3400930021033	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	3400936251663	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	3400936251724	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	3400936251892	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	3400938285154	ZAMBON FRANCE S.A.	FR
Spifen 400 mg, comprimé pelliculé	not available	3400936250833	ZAMBON FRANCE S.A.	FR
Spifen 400 mg, comprimé pelliculé	not available	3400936251083	ZAMBON FRANCE S.A.	FR
Spifen 400 mg, comprimé pelliculé	not available	3400936251144	ZAMBON FRANCE S.A.	FR
Spifen 400 mg, comprimé pelliculé	not available	3400936251205	ZAMBON FRANCE S.A.	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
Spifen 400 mg, comprimé pelliculé	not available	3400936251373	ZAMBON FRANCE S.A.	FR
Spifen 400 mg, comprimé pelliculé	not available	3400936251434	ZAMBON FRANCE S.A.	FR
Spedifen 400 mg granulátum	not available	OGYI-T-7162/01	ZAMBON S.P.A.	HU
Spedifen 400 mg granulátum	not available	OGYI-T-7162/02	ZAMBON S.P.A.	HU
Spedifen 600 mg granulátum	not available	OGYI-T-7162/03	ZAMBON S.P.A.	HU
Spedifen 600 mg granulátum	not available	OGYI-T-7162/04	ZAMBON S.P.A.	HU
Ibalgin krém 50 mg/1 g	not available	29/0051/02-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ibalgin krém 50 mg/1 g	not available	29/0051/02-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ibuprofen 100 mg/5ml oral suspension	not available	PL 00037/0677	ABBOTT LABORATORIES 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386019	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386033	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386021	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386072	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386045	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386058	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386060	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386096	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386084	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386110	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386108	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386134	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386122	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386146	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386159	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386173	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386161	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386197	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386185	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386211	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386209	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386235	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386223	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386247	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386250	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386298	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386286	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386274	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386262	MYLAN S.P.A.	IT
Brufen Analgesico 400 mg compresse rivestite con film	NL/H/2550/002	042386300	MYLAN S.P.A.	IT
Ibuprofen (als lysine) Mylan OTC 200 mg, filmomhulde tabletten	NL/H/2550/001	RVG 111362	MYLAN B.V.	NL
Ibuprofen (als lysine) Mylan OTC 400 mg, filmomhulde tabletten	NL/H/2550/002	RVG 111363	MYLAN B.V.	NL
Ibuprofen LIDERFARM 40 mg/ml Suspension zum Einnehmen	not available	87685.00.00	FARMALIDER, S.A.	DE
ADVILMED ENFANTS ET NOURRISSONS 20 mg/1 ml, suspension buvable en flacon	not available	34009 336 406 2 0	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 432-4	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 431-8	PFIZER SANTE FAMILIALE	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
ADVILCAPS 400 mg, capsule molle	not available	364 427-0	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 631 1 7	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 629 7 4	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 426-4	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 630 5 6	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 626 8 4	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	34009 382 866 2 5	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 433-0	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 632 8 5	PFIZER SANTE FAMILIALE	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
ADVILGEL 5 %, gel	not available	376 851-7	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 430-1	PFIZER SANTE FAMILIALE	FR
ADVILGEL 5 %, gel	not available	376 852-3	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 628 0 6	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 429-3	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	34009 382 865 6 4	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 627 4 5	PFIZER SANTE FAMILIALE	FR
ADVILCAPS 400 mg, capsule molle	not available	364 428-7	PFIZER SANTE FAMILIALE	FR
Dolgit Gel, 50 mg/g, żel	not available	R/6758	DOLORGIET GMBH & CO KG	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
BRUFEN 600 600 mg šumivý granulát	not available	29/0502/08-S	MYLAN IRE HEALTHCARE LIMITED	SK
Ibuprofen 100mg/5ml Oral Suspension	not available	04917/0082	PINEWOOD LABORATORIES LIMITED	UK
Numark Ibuprofen 100mg/5ml Oral Suspension	not available	04917/0082	PINEWOOD LABORATORIES LIMITED	UK
Healthpoint Ibuprofen 100mg/5ml Oral Solution	not available	04917/0082	PINEWOOD LABORATORIES LIMITED	UK
Fenpaed Ibuprofen 100 mg/5ml Oral Suspension	not available	04917/0082	PINEWOOD LABORATORIES LIMITED	UK
Poindol 200 mg granulado para solución oral	not available	76907	FARMALIDER, S.A.	ES
Brufen 20 mg/ml sirup	not available	29/916/92-S/C	MYLAN IRE HEALTHCARE LIMITED	CZ
Ibuprofeno ALGIK 20 mg/ml Oral Suspension	not available	AA865/00801	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	MT
Momentact 400 mg sospensione orale in bustine	not available	035618065	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
Momentact 400 mg sospensione orale in bustine	not available	035618077	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618089	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618091	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618103	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618115	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618127	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618139	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618141	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
SPIDIFEN 600 mg Granulato per soluzione orale aroma cola-limone	not available	026916167	ZAMBON 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
SPIDIFEN 600 mg granulato per soluzione orale aroma cola-limone	not available	026916142	ZAMBON ITALIA S.R.L.	IT
Espidifen 400 mg granulado para solución oral sabor menta	not available	60.514	ZAMBON, S.A.U.	ES
Espidifen 600 mg granulado para solución oral sabor albaricoque	not available	68222	ZAMBON, S.A.U.	ES
Brufen Retard 800mg prolonged release tablets	not available	PA 2010/2/7	MYLAN IRE HEALTHCARE LIMITED	IE
BRUFEN® Επικαλυμμένο δισκίο 400 mg	not available	5754	MYLAN IRE HEALTHCARE LIMITED	CY
Ibutop Gel 50 mg/g gel	not available	7878/2015/01	DOLORGIET GMBH & CO KG	RO
proff Schmerzgel, Gel 50mg/g	not available	1 - 21736	DOLORGIET GMBH & CO KG	AT
Ibunin 200 mg Filmtabletten	MT/H/0100/001	1-28281	ACTAVIS GROUP PTC EHF.	AT
ADVIL RAPID 400 mg měkké tobolky	not available	29/596/08-C	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ipren 60 mg suppositorier	not available	46199	MCNEIL SWEDEN AB	SE
Nurofen Orange, 100 mg pehmed närimiskapslid	PL/H/0602/001	925516	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/120	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/121	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/122	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/123	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/124	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/125	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/126	RECKITT BENCKISER KFT	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
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/127	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/128	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/129	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/130	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/131	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/132	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/133	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/134	RECKITT BENCKISER KFT	HU
Nurofen Junior narancsízű 100 mg lágy rágókapszula	PL/H/0602/001	OGYI-T-6793/135	RECKITT BENCKISER KFT	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
Nurofen Junior 100 mg meke kapsule za žvakanje s okusom naranče	PL/H/0602/001	HR-H-987481558	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen Orange 100 mg mīkstās košļājamās kapsulas	PL/H/0602/001	17-0025	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen dla dzieci Junior, 100 mg kapsułki do żucia elastyczne	PL/H/0602/001	23941	RECKITT BENCKISER (POLAND) S.A.	PL
Terbinafină Scholl 1% cremă	DE/H/6185/001	5643/2013/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen for Children Orange Flavoured 100mg Chewable Capsule, Soft	PL/H/0602/001	PL 00063/0723	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Orange 100 mg kramtomosios minkštosios kapsulės	PL/H/0602/001	925516	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Junior pomeranc 100 mg žvýkací mekká tobolka	PL/H/0602/001	07/642/15-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Нурофен за Юноши Портокал 100 mg меки капсули за дъвчене	PL/H/0602/001	20170065	RECKITT BENCKISER (ROMANIA) SRL	BG
Ibuprofeno Worldrugs 40 mg/ml suspensão oral	not available	5677166	LDP TORLAN LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibufem 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	UK
Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	UK
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	UK
Ibuprofen 400mg Caplets	not available	PL 16028/0040	GALPHARM HEALTHCARE LIMITED	UK
Tesco Health Ibuprofen 400mg Tablets	not available	PL 16028/0040	GALPHARM HEALTHCARE LIMITED	UK
Ibutop	not available	18905	DOLORGIET GMBH & CO KG	DK
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/021	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/012	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/020	RATIOPHARM 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
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/016	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/014	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/018	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/013	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/011	RATIOPHARM GMBH	LT
Butifen 400 mg plevele dengtos tabletes	DE/H/2590/001	LT/1/11/2603/015	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/019	RATIOPHARM GMBH	LT
Butifen 400 mg plėvele dengtos tabletės	DE/H/2590/001	LT/1/11/2603/017	RATIOPHARM GMBH	LT
IBU-LYSIN-ratiopharm® 684 mg Filmtabletten 400 mg Ibuprofen	DE/H/2590/001	79779.00.00	RATIOPHARM 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
Ibuxin rapid 400 mg tabletti, kalvopäällysteinen	DE/H/2590/001	28002	RATIOPHARM GMBH	FI
IBU-LYSIN-ratiopharm® 684 mg Filmtabletten 400 mg Ibuprofen	DE/H/2590/001	0687/11080034	RATIOPHARM GMBH	LU
ratioDolor Ibuprofen 400 mg Filmtabletten	DE/H/2590/001	1-30782	TEVA B.V	AT
Brufen Effect 400 mg filmom obložene tablete	not available	HR-H-334193871	MYLAN HRVATSKA D.O.O.	HR
Brufen 400 mg filmom obložene tablete	not available	UP/I-530-09/12-02/348	MYLAN HRVATSKA D.O.O.	HR
Brufen 600 mg filmom obložene tablete	not available	UP/I-530-09/12-02/345	MYLAN HRVATSKA D.O.O.	HR
Ibuprofeno Tarbis Farma 400 mg comprimidos	not available	56.028	TARBIS FARMA, S.L.	ES
Brufen 200 mg comprimidos revestidos por película	not available	8254060	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 200 mg comprimidos revestidos por película	not available	8254078	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen 400 mg comprimidos revestidos por película	not available	5550587	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 400 mg comprimidos revestidos por película	not available	8254052	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 600 mg comprimidos revestidos por película	not available	8660811	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 600 mg comprimidos revestidos por película	not available	5550389	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Ibuprofeno Clover 400 mg capsulas blandas	not available	80975	HC CLOVER PRODUCTOS Y SERVICIOS, S.L.	ES
БРУФЕН 200 mg ефеверсцентни гранули	EE/H/0295/001	20130080	MYLAN EOOD	BG
Brufen, 200 mg kihisevad graanulid	EE/H/0295/001	809113	MYLAN IRE HEALTHCARE LIMITED	EE
Brufen 200 mg Effervescent Granules	EE/H/0295/001	PA 2010/2/3	MYLAN IRE HEALTHCARE LIMITED	IE
FROBEN DOLORE E FEBBRE 200 mg Granulato Effervescente	EE/H/0295/001	041947045	MYLAN IRE HEALTHCARE LIMITED	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
FROBEN DOLORE E FEBBRE 200 mg Granulato Effervescente	EE/H/0295/001	041947058	MYLAN IRE HEALTHCARE LIMITED	IT
FROBEN DOLORE E FEBBRE 200 mg Granulato Effervescente	EE/H/0295/001	041947060	MYLAN IRE HEALTHCARE LIMITED	IT
Brufen 200 mg šnypščiosios granulės	EE/H/0295/001	LT/1/13/3252/001	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 200 mg šnypščiosios granulės	EE/H/0295/001	LT/1/13/3252/002	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 200 mg šnypščiosios granulės	EE/H/0295/001	LT/1/13/3252/003	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 200 mg putojošās granulas	EE/H/0295/001	13-0049	MYLAN IRE HEALTHCARE LIMITED	LV
BRUFEN 200 mg Effervescent Granules	EE/H/0295/001	PL 46302/0003	MYLAN PRODUCTS LIMITED	UK
proff Schmerzcreme	not available	1-20032	DOLORGIET GMBH & CO KG	AT
IBUPROFENE MYLAN ITALIA 400 mg compresse rivestite	not available	042995011	MYLAN 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
BRUFEN 400 mg, comprimés pelliculés	not available	2002016342	MYLAN EPD BVBA/SPRL	LU
BRUFEN FORTE 600 mg, comprimés pelliculés	not available	2002016343	MYLAN EPD BVBA/SPRL	LU
BRUFEN FORTE 600 mg, Filmtabletten	not available	2002016343	MYLAN EPD BVBA/SPRL	LU
BRUFEN 400 mg Filmtabletten	not available	2002016342	MYLAN EPD BVBA/SPRL	LU
BRUFEN 20 mg/ml Sospensione orale	not available	022593228	MYLAN ITALIA S.R.L.	IT
BRUFEN 20 mg/ml Sospensione orale	not available	022593230	MYLAN ITALIA S.R.L.	IT
SPIDIDOL 400 mg compresse rivestite con film	not available	039600061	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg compresse rivestite con film	not available	039600085	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg compresse rivestite con film	not available	039600073	ZAMBON 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
SPEDIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400935251053	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400934750861	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400934750922	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400934751004	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400934751172	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	3400934969126	ZAMBON FRANCE S.A.	FR
Spididol 400 mg compresse rivestite con film	not available	039600010	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg granulato per soluzione orale aroma albicocca	not available	039600022	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg granulato per soluzione orale aroma menta-anice	not available	039600034	ZAMBON 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
SPIDIFEN 600 mg Granulato per soluzione orale aroma albicocca	not available	026916179	ZAMBON ITALIA S.R.L.	IT
SPIDIFEN 600 mg granulato per soluzione orale aroma albicocca	not available	026916130	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg compresse rivestite con film	not available	039600046	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg compresse rivestite con film	not available	039600059	ZAMBON ITALIA S.R.L.	IT
Ibuprofen 800mg PR tablets	not available	PL 00037/0676	ABBOTT LABORATORIES LIMITED	UK
Brufen Syrup	not available	PL 46302/0011	MYLAN PRODUCTS LIMITED	UK
SPIDIFEN 600 mg Granulato per soluzione orale aroma menta-anice	not available	026916155	ZAMBON ITALIA S.R.L.	IT
Spidifen 600 mg granulaat voor drank	not available	BE478853	ZAMBON NV	BE
Spidifen 600 mg granulés pour solution buvable	not available	BE478853	ZAMBON NV	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
Spidifen 400 mg filmom oblozene tablete	not available	UP/I-530-09/12-01/676	ZAMBON S.P.A.	HR
Spidifen 200 mg tablete	not available	UP/I-530-09/12-01/675	ZAMBON S.P.A.	HR
Spidifen 600 mg granulés pour solution buvable	not available	2016040085	ZAMBON NV	LU
SPIDIFEN 400 mg Compresse rivestite con film	not available	026916080	ZAMBON ITALIA S.R.L.	IT
SPEDIFEN 200 mg, comprimé	not available	3400936081109	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	3400936081277	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	3400936081338	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	3400936081567	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	3400936081628	ZAMBON FRANCE S.A.	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
SPIFEN 200 mg, comprimé	not available	3400936081796	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	3400936081857	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	3400936081918	ZAMBON FRANCE S.A.	FR
Spidifen 200 mg comprimès effervescents	not available	BE157726	ZAMBON NV	BE
Spidifen 200 mg bruistabletten	not available	BE157726	ZAMBON NV	BE
Spidifen 200 mg comprimés	not available	2011051165	ZAMBON NV	LU
Spidifen 200 mg comprimés effervescents	not available	2011051164	ZAMBON NV	LU
Spidifen 200 mg granulés pour solution buvable	not available	2011051163	ZAMBON NV	LU
Spidifen 200 mg comprimés	not available	BE386014	ZAMBON NV	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
Spidifen 200 mg tabletten	not available	BE386014	ZAMBON NV	BE
Spidifen 200 mg granulés pour solution buvable	not available	BE157735	ZAMBON NV	BE
Spidifen 200 mg granulaat voor drank	not available	BE157735	ZAMBON NV	BE
Spidifen 400 mg granulés pour solution buvable	not available	BE157805	ZAMBON NV	BE
Spidifen 400 mg granulaat voor drank	not available	BE157805	ZAMBON NV	BE
Spididol analgesico 200 mg compresse	not available	028710034	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL ANALGESICO 200 mg granulato per soluzione orale	not available	028710022	ZAMBON ITALIA S.R.L.	IT
SPIDIFEN 400 mg Granulato per soluzione orale aroma albicocca	not available	026916104	ZAMBON ITALIA S.R.L.	IT
Spedifen 400 mg filmtabletta	not available	OGYI-T-7162/05	ZAMBON S.P.A.	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
Spedifen 400 mg filmtabletta	not available	OGYI-T-7162/06	ZAMBON S.P.A.	HU
Spedifen 400 mg filmtabletta	not available	OGYI-T-7162/07	ZAMBON S.P.A.	HU
Spedifen 400 mg filmtabletta	not available	OGYI-T-7162/08	ZAMBON S.P.A.	HU
Ibuprofeno Zentiva 400 mg comprimido revestido por película	CZ/H/0184/001	5397773	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/02	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/04	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/03	SANOFI ROMANIA SRL	RO
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/002	SANOFI-AVENTIS D.O.O.	SI
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS 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
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/06	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/10	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/05	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/12	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/11	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/07	SANOFI ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/09	SANOFI ROMANIA SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/08	SANOFI ROMANIA SRL	RO
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG

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
Algoflex Rapid 400 mg filmtabletta	CZ/H/0184/001	OGYI-T-21112/02	SANOFI-AVENTIS ZRT	HU
Algoflex Rapid 400 mg filmtabletta	CZ/H/0184/001	OGYI-T-21112/01	SANOFI-AVENTIS ZRT	HU
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
ИБАЛГИН ФАСТ 400 mg филмирани таблетки	CZ/H/0184/001	20090398	SANOFI BULGARIA EOOD	BG
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/01	SANOFI ROMANIA SRL	RO
Ibuprofeno Zentiva 400 mg comprimido revestido por película	CZ/H/0184/001	5397765	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/004	SANOFI-AVENTIS 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
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/006	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/007	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/005	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/003	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/010	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/011	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/008	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/012	SANOFI-AVENTIS D.O.O.	SI
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/009	SANOFI-AVENTIS 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
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
Ibalgin Rapid 400 mg, potahované tablety	CZ/H/0184/001	29/581/09-C	SANOFI-AVENTIS SRO	CZ
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg filmom obalené tablety	CZ/H/0184/001	29/0543/09-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS 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
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN FAST 400 mg, tabletki powlekane	CZ/H/0184/001	16794	SANOFI-AVENTIS SP Z.O.O.	PL
IBALGIN NEO 400 mg filmsko obložene tablete	CZ/H/0184/001	H/13/00749/001	SANOFI-AVENTIS D.O.O.	SI
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND 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
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ibufarmalid, 400 mg 10 ml, zawiesina doustna	not available	21883	FARMALIDER, S.A.	PL
Ibuprofen Farmalider 20 mg/ml oral suspension	FI/H/0799/001	47788	FARMALIDER, S.A.	SE
Ibuprofen Farmalider 40 mg/ml oral suspension	FI/H/0799/002	47789	FARMALIDER, S.A.	SE
Ibuprofen ”Farmalider”, oral suspension	FI/H/0799/001	50518	FARMALIDER, S.A.	DK
Ibuprofen ”Farmalider”, oral suspension	FI/H/0799/002	50519	FARMALIDER, S.A.	DK
Ibuprofen Farmalider 40 mg/ml oraalisuspensio	FI/H/0799/002	30596	FARMALIDER, S.A.	FI
Neobrufen 600 mg granulado efervescente	not available	61610	MYLAN IRE HEALTHCARE LIMITED	ES
Ibudol Pediátrico 200 mg suspensión oral	not available	68.086	KERN PHARMA, 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
Neobrufen retard 800 mg comprimidos de liberación prolongada	not available	61618	MYLAN IRE HEALTHCARE LIMITED	ES
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386019	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386033	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386021	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386072	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386045	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386058	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386060	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386096	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386084	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386110	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386108	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386134	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386122	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386146	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386159	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386173	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386161	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386197	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386185	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386211	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386209	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386235	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386223	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386247	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386250	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386298	MYLAN 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
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386286	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386274	MYLAN S.P.A.	IT
Brufen Analgesico 200 mg compresse rivestite con film	NL/H/2550/001	042386262	MYLAN S.P.A.	IT
Brufen Analgesico 400 mg compresse rivestite con film	NL/H/2550/002	042386300	MYLAN S.P.A.	IT
Ibuprofen (als lysine) Mylan OTC 200 mg, filmomhulde tabletten	NL/H/2550/001	RVG 111362	MYLAN B.V.	NL
Ibuprofen (als lysine) Mylan OTC 400 mg, filmomhulde tabletten	NL/H/2550/002	RVG 111363	MYLAN B.V.	NL
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (φράουλα)	DE/H/2343/002	021713	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Nurofen Forte za djecu 200 mg/5 ml oralna suspenzija s okusom jagode	DE/H/2343/002	HR-H-140153836	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Нурофен за Юноши Ягода 200 mg/5 ml перорална суспензия (Nurofen Junior	DE/H/2343/002	20150019	RECKITT BENCKISER (ROMANIA) SRL	BG

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
Strawberry 200 mg/5 ml oral suspension)				
Nurofen Morango 40 mg/ml suspensão oral	DE/H/2343/002	5630157	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen eperízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/002	OGYI-T-6793/116	RECKITT BENCKISER KFT	HU
Nurofen Junior cu aromă de căpșuni 40 mg/ml suspensie orală	DE/H/2343/002	9339/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de căpșuni 40 mg/ml suspensie orală	DE/H/2343/002	9339/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de căpșuni 40 mg/ml suspensie orală	DE/H/2343/002	9339/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de căpșuni 40 mg/ml suspensie orală	DE/H/2343/002	9339/2016/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de căpșuni 40 mg/ml suspensie orală	DE/H/2343/002	9339/2016/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen voor Kinderen Suikervrij rood 4% suspensie voor oraal gebruik	DE/H/2343/002	BE378831	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
Nurofen pour Enfants Sans sucre rouge 4% suspension buvable	DE/H/2343/002	BE378831	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder Zuckerfrei Rot 4% Suspension zum Einnehmen	DE/H/2343/002	BE378831	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pro děti 4% Jahoda perorální suspenze	DE/H/2343/002	07/969/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen Schmerz- und Fiebersaft Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2343/002	77581.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen eperízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/002	OGYI-T-6793/105	RECKITT BENCKISER KFT	HU
Nurofen eperízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/002	OGYI-T-6793/106	RECKITT BENCKISER KFT	HU
Nurofen for Children Six Plus Strawberry 200mg/5ml Oral Suspension	DE/H/2343/002	PA 979/56/2	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen voor Kinderen Aardbei suspensie, suspensie 200mg/5ml	DE/H/2343/002	RVG 104637	RECKITT BENCKISER HEALTHCARE B.V.	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
Nurofen pre deti 4 % jahoda perorálna suspenzia	DE/H/2343/002	07/0004/11-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen pour Enfants Sans sucre rouge 4% suspension buvable	DE/H/2343/002	2010120026	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (φράουλα)	DE/H/2343/002	2075016	RECKITT BENCKISER HELLAS CHEMICALS ABEE	GR
Nurofen dla Dzieci 4% truskawkowy, 40 mg/ml, zawiesina doustna	DE/H/2343/002	17855	RECKITT BENCKISER (POLAND) S.A.	PL
Ibuprofen LIDERFARM 20 mg/ml Suspension zum Einnehmen	not available	87684.00.00	FARMALIDER, S.A.	DE
Neobrufen 400 mg Brausegranulat	SE/H/1184/001	1-31789	MYLAN ÖSTERREICH GMBH	AT
Brufen Granules 400 mg, Brausegranulat	SE/H/1184/001	BE434411	MYLAN EPD BVBA/SPRL	BE
Brufen Granules 400mg, granulés effervescents	SE/H/1184/001	BE434411	MYLAN EPD BVBA/SPRL	BE
Brufen Granules 400 mg, bruisgranulaat	SE/H/1184/001	BE434411	MYLAN EPD BVBA/SPRL	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
Brumare, 400 mg kihisevad graanulid	SE/H/1184/001	805213	MYLAN IRE HEALTHCARE LIMITED	EE
Brufen 400 mg pezsgőgranulátum	SE/H/1184/001	OGYI-T-22073/05	MYLAN EPD KFT.	HU
Brufen 400 mg effervescent Granules	SE/H/1184/001	PA 2010/2/4	MYLAN IRE HEALTHCARE LIMITED	IE
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/002	MYLAN IRE HEALTHCARE LIMITED	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/003	MYLAN IRE HEALTHCARE LIMITED	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/001	MYLAN IRE HEALTHCARE LIMITED	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/005	MYLAN IRE HEALTHCARE LIMITED	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/004	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 400 mg granulado efervescente	SE/H/1184/001	5571567	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen 400 mg brusgranulat	SE/H/1184/001	46807	BGP PRODUCTS AB	SE
BRUFEN INSTANT 400 mg šumivý granulát	SE/H/1184/001	29/0162/13-S	MYLAN IRE HEALTHCARE LIMITED	SK
Brufen Granules 400 mg	SE/H/1184/001	2013110373	MYLAN EPD BVBA/SPRL	LU
Brufen 400 mg Effervescent Granules	SE/H/1184/001	PL 46302/0005	MYLAN PRODUCTS LIMITED	UK
Бруфен 400 mg ефервесцентни гранули	SE/H/1184/001	20130355	MYLAN EOOD	BG
FROBEN DOLORE E INFIAMMAZIONE 400 mg Granulato Effervescente	SE/H/1184/001	043155011	MYLAN IRE HEALTHCARE LIMITED	IT
FROBEN DOLORE E INFIAMMAZIONE 400 mg Granulato Effervescente	SE/H/1184/001	043155035	MYLAN IRE HEALTHCARE LIMITED	IT
FROBEN DOLORE E INFIAMMAZIONE 400 mg Granulato Effervescente	SE/H/1184/001	043155023	MYLAN IRE HEALTHCARE LIMITED	IT
FROBEN DOLORE E INFIAMMAZIONE 400 mg Granulato Effervescente	SE/H/1184/001	043155050	MYLAN IRE HEALTHCARE LIMITED	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
FROBEN DOLORE E INFIAMMAZIONE 400 mg Granulato Effervescente	SE/H/1184/001	043155047	MYLAN IRE HEALTHCARE LIMITED	IT
Brumare 400 mg putojošās granulas	SE/H/1184/001	13-0078	MYLAN IRE HEALTHCARE LIMITED	LV
Brufen Gran 400 mg šumeča zrnca	SE/H/1184/001	H/13/00312/002	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Gran 400 mg šumeča zrnca	SE/H/1184/001	H/13/00312/003	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Gran 400 mg šumeča zrnca	SE/H/1184/001	H/13/00312/001	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Gran 400 mg šumeča zrnca	SE/H/1184/001	H/13/00312/004	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Gran 400 mg šumeča zrnca	SE/H/1184/001	H/13/00312/005	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Granules 400 mg, Brausegranulat	SE/H/1184/001	2013110373	MYLAN EPD BVBA/SPRL	LU
Moment 20 g/100 ml gocce per soluzione orale	not available	025669033	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
Ibuprofen Tablets BP 400 mg	not available	PL 20416/0272	CRESCENT PHARMA LIMITED	UK
Rofen™	not available	PL 20416/0272	CRESCENT PHARMA LIMITED	UK
Dolgit Gel	not available	29/0851/95-S	DOLORGIET GMBH & CO KG	SK
BRUFEN GRANULES 600 mg, granulés effervescents	not available	2002016339	MYLAN EPD BVBA/SPRL	LU
BRUFEN GRANULES 600 mg, Brausegranulat	not available	2002016339	MYLAN EPD BVBA/SPRL	LU
Ibufarmalid, 200 mg/ 10 ml, zawiesina doustna	not available	21882	FARMALIDER, S.A.	PL
Ibuleve/Ibuleve Sports Gel	not available	PL 00173/0413	DIOMED DEVELOPMENTS LIMITED	UK
Nurofen for Children 200 mg/5 ml Strawberry Oral Suspension	UK/H/7028/002/DC	PL 00063/0743	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Childrens Ibuprofen 100mg/5ml Oral Suspension	not available	PL 16028/0075	GALPHARM HEALTHCARE 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
Dolgit krem	not available	29/0751/92-S	DOLORGIET GMBH & CO KG	SK
Ibutop Gel	not available	MA180/00101	DOLORGIET GMBH & CO KG	MT
Easofen Max Strength 400 mg Film-coated Tablets	not available	PA0126/060/002	CLONMEL HEALTHCARE LTD.	IE
Advil, dragees 200 mg	not available	RVG 12471	PFIZER B.V.	NL
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/11	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/09	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/06	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/02	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/05	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/07	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/01	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/08	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/12	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/03	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/10	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra Forte 400 mg capsule moi	not available	7168/2014/04	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/01	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/03	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Advil Ultra 200 mg capsule moi	not available	8319/2015/02	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/04	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil 200 mg drajeuri	not available	7261/2006/01	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil 200 mg drajeuri	not available	7261/2006/02	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil 200 mg drajeuri	not available	7261/2006/05	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil 200 mg drajeuri	not available	7261/2006/03	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil 200 mg drajeuri	not available	7261/2006/04	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
IbuViva 100mg/5ml Geriamoji Suspensija	IE/H/0691/001/MR	LT/1/10/2162/001	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 100mg/5ml Geriamoji Suspensija	IE/H/0691/001/MR	LT/1/10/2162/002	PINEWOOD LABORATORIES LIMITED	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
IbuViva 100mg/5ml Geriamoji Suspensija	IE/H/0691/001/MR	LT/1/10/2162/003	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 100mg/5ml Geriamoji Suspensija	IE/H/0691/001/MR	LT/1/10/2162/004	PINEWOOD LABORATORIES LIMITED	LT
Fenopine 100mg/5ml Paediatric Oral Suspension	IE/H/0691/001/MR	PA0281/088/004	PINEWOOD LABORATORIES LIMITED	IE
Milifen 100 mg/5 ml, zawiesina doustna	IE/H/0691/001/MR	NR17354	PINEWOOD LABORATORIES LIMITED	PL
Brufen Effect 200 mg šumeće granule	not available	UP/I-530-09/13-01/09	MYLAN HRVATSKA D.O.O.	HR
Brufen Effect 200 mg šumeće granule	not available	UP/I-530-09/12-01/88	MYLAN HRVATSKA D.O.O.	HR
Brufen Effect 400 mg šumeće granule	not available	UP/I-530-09/12-01/545	MYLAN HRVATSKA D.O.O.	HR
Brufen 400 mg šumeće granule	not available	UP/I-530-09/13-01/08	MYLAN HRVATSKA D.O.O.	HR
Brufen 600 mg šumeće granule	not available	UP/I-530-09/12-02/346	MYLAN 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
Actiprofen 200 mg tablets	not available	PA0678/061/002	GLAXOSMITHKLINE CONSUMER HEALTHCARE (IRELAND) LTD	IE
ibuxin rapid 400 mg filmuhúðaðar töflur	IS/H/0353/001	IS/1/12/080/01	RATIOPHARM GMBH	IS
IBUPROFEN TEVA 5% GEL	not available	BE179715	TEVA PHARMA BELGIUM N.V./S.A	BE
Ibuprofen TEVA 5% gel	not available	BE179715	TEVA PHARMA BELGIUM N.V./S.A	BE
Ibuprofen Teva 5 % gel	not available	BE179715	TEVA PHARMA BELGIUM N.V./S.A	BE
Ibutop 5% geel	not available	280299	DOLORGIET GMBH & CO KG	EE
Ibuprofen 200mg/5ml Oral Suspension	not available	04917/0099	PINEWOOD LABORATORIES LIMITED	UK
Flarin 200 mg soft capsules	not available	PL 51724/0001	INFIRST LTD	UK
Perdofemina 400 mg, comprimés pelliculés	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
Perdofemina 400 mg, filmomhulde tabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, comprimés pelliculés	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, filmomhulde tabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, filmomhulde tabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, comprimés pelliculés	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina, 400 mg, Filmtabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina, 400 mg, Filmtabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina, 400 mg, Filmtabletten	not available	BE 228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
ACTIBU FEBBRE E DOLORE 400 mg compresse rivestite con film.	not available	029129069	JOHNSON & JOHNSON 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
Actibu febbre e dolore 200 mg compresse rivestite con film	not available	029129020	JOHNSON & JOHNSON S.P.A.	IT
Actibu febbre e dolore 200 mg compresse rivestite con film	not available	029129032	JOHNSON & JOHNSON S.P.A.	IT
Actibu febbre e dolore 200 mg compresse rivestite con film.	not available	029129018	JOHNSON & JOHNSON S.P.A.	IT
Ibuprofen Twelve Plus Pain Relief 200mg/5ml oral suspension	UK/H/6039/002	PL35533/0034	ASPIRE PHARMA LIMITED	UK
Ibuprofen 200mg Caplets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	UK
Ibuprofen 200mg Tablets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	UK
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	UK
Galprofen Long Lasting Ibuprofen Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Asda Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE 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
Morrisons Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Superdrug Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Wilko Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Boots Ibuprofen Long Lasting Capsules 200mg	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Lloydspharmacy Ibuprofen 200mg Long Lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Teva Ibuprofen 200mg Long lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
The-Cooperative Ibuprofen 200mg Long Lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Tesco Health Long Lasting Pain Relief, 200mg Modified Release Capsules, Hard	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Essential Waitrose Long Lasting Pain Relief 200mg Modified Release Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE 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
Sainsbury's Healthcare Long Lasting Pain Relief 200mg Modified Release Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	UK
Brufen® Επικαλυμμένο με λεπτό υμένιο δισκίο 600 mg	not available	48827/10-09-2009	BGP PRODUCTS S.LTD	GR
Brufen®	not available	48832/10-09-2009	BGP PRODUCTS S.LTD	GR
Brufen®	not available	48830/10-09-2009	BGP PRODUCTS S.LTD	GR
Ibuprofen 600mg Effervescent Granules	not available	PL 00037/0678	ABBOTT LABORATORIES LIMITED	UK
JOINTRAL 100 mg/g gel	not available	040608010	LAB. IT. BIOCHIM. FARM.CO LISAPHARMA S.P.A.	IT
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/04	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/08	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/09	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen rapid 200 mg Weichkapseln	DE/H/1482/001	1-29414	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen rapid 200 mg Weichkapseln	DE/H/1482/001	1-29414	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
НУРОФЕН ЕКСПРЕС 200 mg меки капсули	DE/H/1482/001	20100224	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Rapid 200 mg Capsules	DE/H/1482/001	07/025/11-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Immedia 200 mg Weichkapseln	DE/H/1482/001	71862.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Fastine Liquid Caps 200 mg, capsule, zacht	DE/H/1482/001	RVG 102120	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Express Femina, 200 mg, kapsułki, miękkie	DE/H/1482/001	16931	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Rapid 200 mg Capsules, mäkké kapsuly	DE/H/1482/001	07/0178/11-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/85	RECKITT BENCKISER KFT	HU
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/86	RECKITT BENCKISER KFT	HU
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/87	RECKITT BENCKISER KFT	HU
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/88	RECKITT BENCKISER KFT	HU
Nurofen ® 200mg Liquid Capsules	DE/H/1482/001	20918	RECKITT BENCKISER HELLAS CHEMICALS ABEE	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
Nurofen ® 200mg Μαλακές Κάψουλες	DE/H/1482/001	2075015	RECKITT BENCKISER HELLAS CHEMICALS ABEE	GR
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/89	RECKITT BENCKISER KFT	HU
Nurofen Rapid 200 mg lágy kapszula	DE/H/1482/001	OGYI-T-6793/90	RECKITT BENCKISER KFT	HU
Nurofen Express 200 mg capsule moi	DE/H/1482/001	7253/2014/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Ibuprofeno Farmalid 400 mg solución para perfusión	not available	82808	NUTRA ESSENTIAL OTC, S.L.	ES
ALGOFEN 200 mg compresse rivestite	not available	023766037	LABORATORIO FARMACEUTICO S.I.T. S.R.L.	IT
ALGOFEN 200 mg compresse rivestite	not available	023766025	LABORATORIO FARMACEUTICO S.I.T. S.R.L.	IT
Ibugel 5% w/w Gel	not available	0278/020/001	DERMAL LABORATORIES LIMITED	IE
Ibuprofen FARMALIDER 20 mg/ml peroralna suspenzija	DE/H/2597/001	5363-I-1601/12	FARMALIDER, S.A.	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
Ibuprofen FARMALIDER 20 mg/ml peroralna suspenzija	DE/H/2597/001	5363-I-1602/12	FARMALIDER, S.A.	SI
Ibuprofen Farmalider 20 mg/ml peroralna suspenzija	DE/H/2597/001	5363-I-1603/12	FARMALIDER, S.A.	SI
Ibuprofen FARMALIDER 20 mg/ml, zawiesina doustna	DE/H/2597/001/DC	21257	FARMALIDER, S.A.	PL
Ibuprofeno Farmalider 20 mg/ml suspensão oral	DE/H/2597/001	5442108	FARMALIDER, S.A.	PT
Ibuprofeno Farmalider 20 mg/ml suspensão oral	DE/H/2597/001	5442116	FARMALIDER, S.A.	PT
Ibuprofeno Farmalider 20 mg/ml suspensão oral	DE/H/2597/001	5442124	FARMALIDER, S.A.	PT
Ibudol roll-on 50 mg/g gel envase con aplicador de bola	not available	72.989	KERN PHARMA, S.L.	ES
Ibuprofeno Algik 20 mg/ml Suspensão Oral	not available	5106109	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Ibuprofeno Algik 20 mg/ml suspensão oral	not available	5106075	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	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
Perdophen 200 mg, filmomhulde tabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg, comprimés pelliculés	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN, 200 mg Filmtabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg, filmomhulde tabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN, 200 mg Filmtabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg, comprimés pelliculés	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN, 200 mg Filmtabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg, comprimés pelliculés	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg, filmomhulde tabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
Brufen 200 mg granulado efervescente	not available	5082680	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 200 mg granulado efervescente	not available	5082789	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 200 mg granulado efervescente	not available	5474317	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 200 mg granulado efervescente	not available	5474325	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 600 mg granulado efervescente	not available	5550488	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Ibuprofeno Farmalider 400 mg concentrado para solución para perfusión	not available	83570	FARMALIDER, S.A.	ES
BRUFEN 400 400 mg filmom obalené tablety	not available	29/0110/06-S	MYLAN IRE HEALTHCARE LIMITED	SK
AAH Pharmaceutical's Ibuprofen pain relief gel maximum strength 10% w/w	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Boots Ibuprofen Max Strength 10% Gel	not available	PL 10972/0082	MERCURY PHARMA GROUP 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
Fenbid Forte 10% Gel	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Ibuprofen 10% gel	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Numark's Ibuprofen Pain Relief Gel Maximum Strength 10% w/w	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Lloydspharmacy's Maximum strength Ibuprofen 10% gel	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Morrison's Ibuprofen Pain Relief Gel Maximum Strength 10% w/w	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Phorpain gel maximum strength	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Tesco's Ibuprofen pain relief gel maximum strength 10% w/w	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
Thornton &Ross's Ibuprofen 10% gel	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	UK
БРУФЕН РЕТАРД 800 mg таблетки с удължено освобождаване	SE/H/0912/005	20110338	MYLAN EOOD	BG

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
Brufen, 20 mg/ml suukaudne suspensioon	SE/H/0912/004	750711	MYLAN IRE HEALTHCARE LIMITED	EE
Brufen, 800 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0912/005	750811	MYLAN IRE HEALTHCARE LIMITED	EE
Brufen 800 mg retard tabletta	SE/H/0912/005	OGYI-T-22073/02	MYLAN EPD KFT.	HU
Brufen 800 mg retard tabletta	SE/H/0912/005	OGYI-T-22073/01	MYLAN EPD KFT.	HU
Brufen 20 mg/ml geriamoji suspensija	SE/H/0912/004	LT/1/10/1885/010	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 20 mg/ml geriamoji suspensija	SE/H/0912/004	LT/1/10/1885/009	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 800 mg pailginto atpalaidavimo tabletės	SE/H/0912/005	LT/1/10/1885/008	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 800 mg pailginto atpalaidavimo tabletės	SE/H/0912/005	LT/1/10/1885/007	MYLAN IRE HEALTHCARE LIMITED	LT
Brufen 20 mg/ml suspensija iekšķīgai lietošanai	SE/H/0912/004	11-0250	MYLAN IRE HEALTHCARE LIMITED	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen 800 mg ilgstošās darbības tabletes	SE/H/0912/005	11-0251	MYLAN IRE HEALTHCARE LIMITED	LV
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/01	BGP PRODUCTS AB	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/02	BGP PRODUCTS AB	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/04	BGP PRODUCTS AB	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/06	BGP PRODUCTS AB	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/03	BGP PRODUCTS AB	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/05	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/01	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/02	BGP PRODUCTS AB	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/03	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/04	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/05	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/06	BGP PRODUCTS AB	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/07	BGP PRODUCTS AB	RO
Brufen Retard 800 mg comprimate cu eliberare prelungită	SE/H/0912/005	7461/2015/01	BGP PRODUCTS AB	RO
Brufen Retard 800 mg comprimate cu eliberare prelungită	SE/H/0912/005	7461/2015/02	BGP PRODUCTS AB	RO
Brufen 200 mg filmsko obložene tablete	SE/H/0912/001	H/10/01998/001	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/016	MYLAN IRE HEALTHCARE LIMITED	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
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/018	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/019	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/020	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/017	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/021	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/022	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/023	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 200 mg filmsko obložene tablete	SE/H/0912/001	H/10/01998/002	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 200 mg filmsko obložene tablete	SE/H/0912/001	H/10/01998/003	MYLAN IRE HEALTHCARE LIMITED	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
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/004	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/005	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/006	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/007	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/008	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/009	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/010	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/011	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/012	MYLAN IRE HEALTHCARE LIMITED	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
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/013	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen retard 800 mg tablete s podaljšanim sproščanjem	SE/H/0912/005	H/10/01998/014	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen retard 800 mg tablete s podaljšanim sproščanjem	SE/H/0912/005	H/10/01998/015	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen Retard 800 mg prolonged-release tablets	SE/H/0912/005	MA1187/03202	MYLAN IRE HEALTHCARE LIMITED	MT
Brufen 400 mg film-coated tablets	SE/H/0912/002	MA1187/03201	MYLAN IRE HEALTHCARE LIMITED	MT
Ibuprofen Teva, 200 mg, tabletki powlekane	PL/H/0592/001	20772	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Ibuprofen Teva MAX, 400 mg tabletki powlekane	PL/H/0592/002	20773	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Ibuprofen B. Braun 200 mg infusioonilahus	ES/H/0392/002/DC	1001020	B.BRAUN MELSUNGEN AG	EE
Ibuprofen B. Braun 200 mg infuusioneste, liuos	ES/H/0392/002	36701	B.BRAUN MELSUNGEN AG	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
Ibuprofen B. Braun 200 mg infusionsvätska, lösning	ES/H/0392/002	36701	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 200 mg šķīdums infūzijām	ES/H/0392/002/DC	20-0016	B.BRAUN MELSUNGEN AG	LV
Ibuprofen B. Braun 200 mg infusjonsvæske, oppløsning	ES/H/0392/002	18-12653	B.BRAUN MELSUNGEN AG	NO
Ibuprofen B. Braun 200 mg infúzny roztok	ES/H/0392/002	07/0041/20-S	B.BRAUN MELSUNGEN AG	SK
Ibuprofen 200 mg solution for infusion	ES/H/0392/002	PL 03551/0155	B.BRAUN MELSUNGEN AG	UK
Ibuprofen B. Braun 200 mg infusionsvätska, lösning	ES/H/0392/002	58662	B.BRAUN MELSUNGEN AG	SE
Ibuprofen B. Braun 200 mg oplossing voor infusie	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg, solution pour perfusion	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	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
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg, solution pour perfusion	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg oplossing voor infusie	ES/H/0392/002	BE557600	B.BRAUN MELSUNGEN AG	BE
Brufen Suspensão 20 mg/ml suspensão oral	not available	8523209	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Ibuspray	not available	PL 00173/0150	DIOMED DEVELOPMENTS LIMITED	UK
Espididol 400 mg comprimidos recubiertos	not available	65843	ZAMBON, S.A.U.	ES
ADVIL RAPID 400 mg, mäkké kapsuly	not available	29/0330/09-S	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	SK
ADVIL 5 %, gel	not available	357 599-4	PFIZER SANTE FAMILIALE SAS	FR
ADVIL 5 %, gel	not available	360 944-0	PFIZER SANTE FAMILIALE SAS	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
ADVIL 5 %, gel	not available	357 598-8	PFIZER SANTE FAMILIALE SAS	FR
Brufen SR 800 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-02/347	MYLAN HRVATSKA D.O.O.	HR
Moment 200 mg capsule molli	not available	025669197	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg capsule molli	not available	025669209	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Dolgit Gel	not available	29/0851/95-S	DOLORGIET GMBH & CO KG	SK
Ibuprofen 100mg/5ml Oral Suspension	not available	29831/0695	WOCKHARDT UK LTD	UK
Nurofen dla dzieci Forte pomarańczowy, 40 mg/ml, zawiesina doustna	DE/H/2206/001	22448	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/024	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex cu aroma de portocale 40 mg/ml suspensie orala	DE/H/2206/001	9340/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nuroflex cu aroma de portocale 40 mg/ml suspensie orala	DE/H/2206/001	9340/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de portocale 40 mg/ml suspensie orala	DE/H/2206/001	9340/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de portocale 40 mg/ml suspensie orala	DE/H/2206/001	9340/2016/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de portocale 40 mg/ml suspensie orala	DE/H/2206/001	9340/2016/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Нурофен за Деца Форте Портокал 200 mg/5 ml перорална суспензия (Nurofen for Children Forte Orange 200 mg/5 ml oral suspension)	DE/H/2206/001	20100670	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Forte Orange, 40 mg/ml, suukaudne suspensioon	DE/H/2206/001	707310	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2206/001	76553.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen for Children Orange 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2206/001	10-0476	RECKITT BENCKISER (POLAND) S.A.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/013	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/014	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/015	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/016	RECKITT BENCKISER (POLAND) S.A.	LT
Junifen 40 mg/ml suspensión oral sabor naranja	DE/H/2206/001	73.047	RECKITT BENCKISER HEALTHCARE S.A.	ES
Ipren 20 mg/ml, oral suspension	not available	19260	MCNEIL SWEDEN AB	SE
Ipren 20 mg/ml, oral suspension	not available	19260	MCNEIL SWEDEN AB	SE
Ipren 20 mg/ml, oral suspension	not available	19260	MCNEIL SWEDEN AB	SE
IBUTOP gel	not available	NR 1996070435	OMEGA PHARMA BELGIUM NV	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
IBUTOP GEL	not available	NR 1996070435	OMEGA PHARMA BELGIUM NV	LU
Ozonol 50 mg/g gel	not available	2322188	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322485	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322584	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322287	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322386	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322782	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322089	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2321982	PERRIGO PORTUGAL LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ozonol 50 mg/g gel	not available	2322683	PERRIGO PORTUGAL LDA	PT
Ozonol 50 mg/g gel	not available	2322881	PERRIGO PORTUGAL LDA	PT
IBUTOP 5% GEL	not available	BE159056	OMEGA PHARMA BELGIUM NV	BE
IBUTOP 5% gel	not available	BE159056	OMEGA PHARMA BELGIUM NV	BE
IBUTOP 5% gel	not available	BE159056	OMEGA PHARMA BELGIUM NV	BE
Solvium 50 mg/g gel	not available	3439692	PERRIGO PORTUGAL LDA	PT
Solvium 50 mg/g gel	not available	3439593	PERRIGO PORTUGAL LDA	PT
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 079 5 0	MYLAN MEDICAL SAS	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 257 0 1	MYLAN MEDICAL SAS	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
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 259 3 0	MYLAN MEDICAL SAS	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 258 7 9	MYLAN MEDICAL SAS	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 260 1 2	MYLAN MEDICAL SAS	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 080 3 2	MYLAN MEDICAL SAS	FR
MIG pediatric 20 mg/ml suspensie orală	DE/H/2463/001	7529/2015/02	BERLIN-CHEMIE AG	RO
MIG pediatric 20 mg/ml suspensie orală	DE/H/2463/001	7529/2015/01	BERLIN-CHEMIE AG	RO
МИГ за деца 20 mg/ml перорална суспензия	DE/H/2463/001	20110424	BERLIN-CHEMIE AG	BG
Ibustar 20 mg/ml belsőleges szuszpenzió gyermekek részére	DE/H/2463/001	OGYI-T-20626/04	BERLIN-CHEMIE AG	HU
Ibustar 20 mg/ml belsőleges szuszpenzió gyermekek részére	DE/H/2463/001	OGYI-T-20626/05	BERLIN-CHEMIE AG	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
Ibustar bērniem 100 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2463/001	11-0290	BERLIN-CHEMIE AG	LV
Ibustar 20 mg/ml geriamoji suspensija, vaikams	DE/H/2463/001	LT/1/11/2632/002	BERLIN-CHEMIE AG	LT
Ibustar 20 mg/ml geriamoji suspensija, vaikams	DE/H/2463/001	LT/1/11/2632/001	BERLIN-CHEMIE AG	LT
MIG Junior 2% 20 mg/ml perorálna suspenzia	DE/H/2463/001	29/0344/11-S	BERLIN-CHEMIE AG	SK
MIG dla dzieci 20 mg/ml, zawiesina doustna	DE/H/2463/001	18904	BERLIN-CHEMIE AG	PL
Ibustar, 20 mg/ml suukaudne suspensioon	DE/H/2463/001	752311	BERLIN-CHEMIE AG	EE
Eudorlin Ibuprofen 20 mg/ml Suspension zum Einnehmen	DE/H/2463/001	78529.00.00	BERLIN-CHEMIE AG	DE
Eudorlin infantil 20 mg/ ml suspensión oral	DE/H/2463/001	76.590	BERLIN-CHEMIE AG	ES
Dolobene® Ibu, 50 mg, Gel Wirkstoff: Ibuprofen	not available	33530.00.00	RECORDATI PHARMA 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
SPIDIFEN 200 mg granulado para solução oral	not available	4617684	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 200 mg granulado para solução oral	not available	5912985	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 400 mg granulado para solução oral	not available	4617981	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 400 mg granulado para solução oral	not available	5912787	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 400 mg granulado para solução oral	not available	4618088	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 400 mg granulado para solução oral	not available	4618187	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Spidifen 600 mg granulado para solução oral (aroma cola-limão)	not available	5717228	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Spidifen 600 mg granulado para solução oral (aroma cola-limão)	not available	5717236	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Spidifen 600 mg granulado para solução oral (aroma cola-limão)	not available	5717244	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ARFEN 500 mg Compresse	not available	024635029	LAB. IT. BIOCHIM. FARM.CO LISAPHARMA S.P.A.	IT
Spidifen 400 mg comprimés pelliculés	NL/H/0341/001	BE244456	ZAMBON NV	BE
Spidifen 400 mg filmomhulde tabletten	NL/H/0341/001	BE244456	ZAMBON NV	BE
Spidifen 400 mg comprimés pelliculés	NL/H/0341/001	2011091290	ZAMBON NV	LU
Spidifen® 400 tablet, filmomhulde tabletten 400 mg	NL/H/0341/001	RVG 25385	ZAMBON NEDERLAND B.V.	NL
SPIDIFEN EF 400 mg comprimido revestido por pelicula	not available	5647177	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN EF 400 mg comprimido revestido por pelicula	not available	4458683	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN EF 400 mg comprimido revestido por pelicula	not available	5472089	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 600 mg granulado para solução oral	not available	5912886	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SPIDIFEN 600 mg granulado para solução oral	not available	4618286	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 600 mg granulado para solução oral	not available	4618385	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Spididol analgesico 200 mg compresse effervescenti	not available	028710010	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL ANALGESICO 200 mg/ml Gocce orali soluzione	not available	028710046	ZAMBON ITALIA S.R.L.	IT
Faspic 400 mg granule pentru soluţie orală	not available	7700/2006/01	ZAMBON S.P.A.	RO
Faspic 400 mg granule pentru soluţie orală	not available	7700/2006/02	ZAMBON S.P.A.	RO
Faspic 400 mg granule pentru soluţie orală	not available	7700/2006/03	ZAMBON S.P.A.	RO
Faspic 600, granule pentru soluţie orală	not available	10106/2017/01	ZAMBON S.P.A.	RO
Faspic 600, granule pentru solutie orala	not available	10106/2017/02	ZAMBON S.P.A.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Faspic 600, granule pentru solutie orala	not available	10106/2017/03	ZAMBON S.P.A.	RO
Faspic 600 mg granule pentru soluţie orală	not available	10106/2017/04	ZAMBON S.P.A.	RO
Faspic 600, granule pentru solutie orala	not available	10106/2017/05	ZAMBON S.P.A.	RO
Faspic 600, granule pentru solutie orala	not available	10106/2017/06	ZAMBON S.P.A.	RO
Spedifen 400, 400 mg, tabletki powlekane	not available	10868	ZAMBON S.P.A.	PL
IBUPROFENE ACRAF 200 mg compresse rivestite.	not available	034178018	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
IBUPROFENE ACRAF 200 mg compresse rivestite.	not available	034178020	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
IBUPROFENE ACRAF 20 g/100 ml gocce orali, soluzione.	not available	034178032	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ibuprofen 600 mg film-coated tablets	not available	PL 44041/0060	NOUMED LIFE SCIENCES	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
Noubid 200 mg film-coated Tablets	not available	PL 44041/0057	NOUMED LIFE SCIENCES	UK
Ibuprofen 200 mg film-coated Tablets	not available	PL 44041/0058	NOUMED LIFE SCIENCES	UK
Ibuprofen 400 mg film-coated tablets	not available	PL 44041/0059	NOUMED LIFE SCIENCES	UK
Ibuprofen Tablets BP 400 mg	not available	PL 20416/0272	CRESCENT PHARMA LIMITED	UK
Rofen™	not available	PL 20416/0272	CRESCENT PHARMA LIMITED	UK
Brufen Paediatric 100mg/5ml oral suspension	not available	PA 2010/2/6	MYLAN IRE HEALTHCARE LIMITED	IE
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 3 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 4 2	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 5 9	ACCORD HEALTHCARE FRANCE SAS	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
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 6 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 7 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 720 8 0	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 1 0	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 0 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 1 7	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 2 4	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 3 1	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 4 8	ACCORD HEALTHCARE FRANCE SAS	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
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 5 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 2 7	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 3 4	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 4 1	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 6 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 7 2	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 8 9	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 721 9 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 300 722 0 2	ACCORD HEALTHCARE FRANCE SAS	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
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 6 2	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 8 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 259 9 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001/DC	34009 550 260 0 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 260 1 3	ACCORD HEALTHCARE FRANCE SAS	FR
Advil®	not available	8577/6-2-2007	PFIZER HELLAS, A.E.	GR
Ibufen 400 mg comprimate filmate	not available	7655/2015/01	ANTIBIOTICE SA	RO
Ibufen 400 mg comprimate filmate	not available	7655/2015/02	ANTIBIOTICE SA	RO
DOLGIT - Creme	not available	1-19151	DOLORGIET GMBH & CO KG	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
doc Ibuprofen Schmerzgel, 5% Gel	not available	1 - 21735	HERMES ARZNEIMITTEL GMBH	AT
Ibuprofen Tablets BP 200mg	not available	PL 39484/0026	FOURRTS (UK) PHARMACARE LIMITED	UK
Ibuprofen Tablets BP 400 mg	not available	PL 39484/0027	FOURRTS (UK) PHARMACARE LIMITED	UK
Ibuprofen Tablets B.P. 600 mg	not available	PL 39484/0028	FOURRTS (UK) PHARMACARE LIMITED	UK
Dolgit gel	not available	29/597/95-C	DOLORGIET GMBH & CO KG	CZ
Ibuprofeno Sandoz 100 mg/5 ml suspensión oral	not available	65529	SANDOZ FARMACÉUTICA, S.A.	ES
Phorpain 5 %w/w Gel	not available	PL 10972/0091	MERCURY PHARMA GROUP LTD	UK
Fenbid 5 %w/w gel	not available	PL 10972/0091	MERCURY PHARMA GROUP LTD	UK
Nurofen for Children 200 mg/5 ml Orange Oral Suspension	UK/H/7028/001/DC	PL 00063/0742	RECKITT BENCKISER HEALTHCARE (UK) 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
IBUPROM SPORT spray, 50 mg/g, aerozol na skórę, roztwór	not available	24970	US PHARMACIA SP. Z O.O.	PL
Ibux 50 mg/g gel	not available	99-553	KARO PHARMA AS	NO
Ibuprofen Nutra Essential 400 mg perorální suspenze v sáčku	CZ/H/0638/002	07/685/15-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra 400 mg perorálna suspenzia vo vrecku	CZ/H/0638/002	07/0093/18-S	NUTRA ESSENTIAL OTC, S.L.	SK
Winadol 400 mg belsoleges szuszpenzió tasakban	CZ/H/0638/002	OGYI-T-23323/04-06	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra 400 mg suspensie orală în plic	CZ/H/0638/002	10548/2018/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 400 mg suspensie orală în plic	CZ/H/0638/002	10548/2018/02	NUTRA ESSENTIAL OTC, S.L.	RO
Winadol 400 mg belsoleges szuszpenzió tasakban	CZ/H/0638/002	OGYI-T-23323/04	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 400 mg belsoleges szuszpenzió tasakban	CZ/H/0638/002	OGYI-T-23323/05	NUTRA ESSENTIAL OTC, S.L.	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
Winadol 400 mg belsoleges szuszpenzió tasakban	CZ/H/0638/002	OGYI-T-23323/06	NUTRA ESSENTIAL OTC, S.L.	HU
Dolgit krém	not available	5284/01	DOLORGIET GMBH & CO KG	HU
VEGETALLUMINA ANTIDOLORE 10% GEL	not available	041734017	PIETRASANTA PHARMA S.P.A.	IT
VEGETALLUMINA ANTIDOLORE 10% GEL	not available	041734029	PIETRASANTA PHARMA S.P.A.	IT
ARFEN 400 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	024635106	LAB. IT. BIOCHIM. FARM.CO LISAPHARMA S.P.A.	IT
Ibuprofen Nutra Essential 200 mg perorální suspenze v sácku	CZ/H/0639/001	07/686/15-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra Essential 200 mg de suspension buvable en sachet	CZ/H/0639/001	BE518657	NUTRA ESSENTIAL OTC, S.L.	BE
BRUFEN 600 mg Granulato effervescente	not available	022593103	MYLAN ITALIA S.R.L.	IT
BRUFEN 600 mg Granulato effervescente	not available	022593178	MYLAN 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
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	SANOFI-AVENTIS GMBH OSTERREICH	AT
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 1 6	SANOFI-AVENTIS FRANCE	FR
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 3 0	SANOFI-AVENTIS 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
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 2 3	SANOFI-AVENTIS FRANCE	FR
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 0 9	SANOFI-AVENTIS FRANCE	FR
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558018	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558020	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558032	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558044	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558057	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ibuprofene Aurobindo Italia 200 mg compresse rivestite con film	PT/H/1776/001	045558069	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Brufen Sem Açúcar 20 mg/ml suspensão oral	not available	5746342	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen Sem Açúcar 20 mg/ml suspensão oral	not available	5746359	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Perdophen 400 mg, comprimés pelliculés	not available	2008099941	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Perdophen 400 mg, comprimés pelliculés	not available	2008099941	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Perdophen 400 mg, comprimés pelliculés	not available	2008099941	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Ibuprofen Seven Plus Pain Relief 200 mg/ 5ml oral suspension	not available	PL35533/0056	ASPIRE PHARMA LIMITED	UK
Cuprofen Ibuprofen Effervescent Tablets 200mg	not available	PL 08215/0111	KENT PHARMACEUTICALS LTD.	UK
Ibuprofen Banner 400 mg kapsel, mjuk	NL/H/4469/001	44549	PATHEON SOFTGELS B.V.	SE
Nurofen Joint & Muscular Pain Relief 200mg Medicated Plaster	DE/H/5067/001	PL 00063/0733	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Patch 200 mg pleister	DE/H/5067/001	BE500080	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
Nurofen Patch 200 mg emplâtre médicamenteux	DE/H/5067/001	BE500080	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen Patch 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	BE500080	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 24-Stunden Schmerzpflaster 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	93368.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Patch 200 mg Emplâtre médicamenteux	DE/H/5067/001	2017040114	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen Durance 200 mg medicated plasters	DE/H/5067/001	PA0979/032/018	RECKITT BENCKISER IRELAND LTD.	IE
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	11221/2018/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	11221/2018/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	11221/2018/02	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	11221/2018/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	11221/2018/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Nurofen 200 mg liečivá náplasť	DE/H/5067/001	29/0369/18-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
NurofenPlast 200 mg, emplâtre médicamenteux	DE/H/5067/001	34009 301 708 5 4	RECKITT BENCKISER HEALTHCARE 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
NurofenPlast 200 mg, emplâtre médicamenteux	DE/H/5067/001	34009 301 708 6 1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen 200 mg léčivá náplast	DE/H/5067/001	29/141/18-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen pleister 200 mg	DE/H/5067/001	RVG 123099	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen, 200 mg, ravimplaaster	DE/H/5067/001	987219	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Mięśnie i Stawy, 200 mg, plaster leczniczy	DE/H/5067/001	25344	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Musc 200 mg emplastro medicamentoso	DE/H/5067/001	5760723	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 200 mg ārstnieciskais plāksteris	DE/H/5067/001	19-0087	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Musc 200 mg emplastro medicamentoso	DE/H/5067/001	5760731	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/140	RECKITT BENCKISER KFT	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
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/136	RECKITT BENCKISER KFT	HU
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/137	RECKITT BENCKISER KFT	HU
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/138	RECKITT BENCKISER KFT	HU
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/139	RECKITT BENCKISER KFT	HU
NUROFLEX Dolori muscolari e articolari, 200 mg cerotto medicato	DE/H/5067/001	047036052	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFLEX Dolori muscolari e articolari, 200 mg cerotto medicato	DE/H/5067/001	047036013	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFLEX Dolori muscolari e articolari, 200 mg cerotto medicato	DE/H/5067/001	047036025	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFLEX Dolori muscolari e articolari, 200 mg cerotto medicato	DE/H/5067/001	047036037	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFLEX Dolori muscolari e articolari, 200 mg cerotto medicato	DE/H/5067/001	047036049	RECKITT BENCKISER HEALTHCARE (ITALIA) 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
Nurofen Durance 200mg φαρμακούχο έμπλαστρο	DE/H/5067/001	109570/19-09-2019	RECKITT BENCKISER HELLAS CHEMICALS ABEE	GR
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/001	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/005	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/002	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/003	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/002	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex 200 mg vaistinis pleistras	DE/H/5067/001	LT/1/19/4400/004	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen 24-Stunden Schmerzpflaster 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	138593	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen 200 mg ljekoviti flaster	DE/H/5067/001	HR-H-092708535	RECKITT BENCKISER (CROATIA) 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
Ibuprofen TABLET 200mg	not available	PL 16028/0161	GALPHARM HEALTHCARE LIMITED	UK
Tesco Health Ibuprofen 200mg Tablets	not available	PL 16028/0161	GALPHARM HEALTHCARE LIMITED	UK
Proff® Schmerzcreme	not available	1192.00.02	DOLORGIET GMBH & CO KG	DE
Espididol 400 mg granulado para solución oral sabor menta	not available	68344	ZAMBON, S.A.U.	ES
DICLOREUM UNIDIE 136 mg cerotto medicato	not available	037184013	ALFASIGMA S.P.A.	IT
DICLOREUM UNIDIE 136 mg cerotto medicato	not available	037184025	ALFASIGMA S.P.A.	IT
DICLOREUM UNIDIE 136 mg cerotto medicato	not available	037184037	ALFASIGMA S.P.A.	IT
DICLOREUM UNIDIE 136 mg cerotto medicato	not available	037184049	ALFASIGMA S.P.A.	IT
DICLOREUM UNIDIE 136 mg cerotto medicato	not available	037184052	ALFASIGMA 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
Brufen Retard 800 mg depottabletter	not available	7614	MYLAN HEALTHCARE NORGE AS	NO
BRUFEN RETARD 800 mg, comprimés à libération prolongée	not available	2002016344	MYLAN EPD BVBA/SPRL	LU
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	2002016344	MYLAN EPD BVBA/SPRL	LU
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (πορτοκάλι)	DE/H/2343/001	021712	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Нурофен за Юноши Портокал 200 mg/5 ml перорална суспензия (Nurofen Junior Orange 200 mg/5 ml oral suspension)	DE/H/2343/001	20150018	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Laranja 40 mg/ml suspensão oral	DE/H/2343/001	5630033	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen narancsízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/001	OGYI-T-6793/115	RECKITT BENCKISER KFT	HU
Nurofen Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen voor Kinderen Suikervrij 4% suspensie voor oraal gebruik	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pour Enfants Sans sucre 4% suspension buvable	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder Zuckerfrei 4% Suspension zum Einnehmen	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pro děti 4% pomeranč perorální suspenze	DE/H/2343/001	07/968/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurodon Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2343/001	77580.00.00	RECKITT BENCKISER DEUTSCHLAND 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
Nurofen narancsízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/001	OGYI-T-6793/103	RECKITT BENCKISER KFT	HU
Nurofen narancsízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/001	OGYI-T-6793/104	RECKITT BENCKISER KFT	HU
Nurofen for Children Six Plus Orange 200mg/5ml Oral Suspension	DE/H/2343/001	PA 979/56/1	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen voor Kinderen Sinaasappel suspensie, suspensie 200mg/5ml	DE/H/2343/001	RVG 104647	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen pre deti 4% pomaranč, perorálna suspenzia	DE/H/2343/001	07/0003/11-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen pour Enfants Sans sucre 4% suspension buvable	DE/H/2343/001	2010120025	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen dla dzieci Junior pomarańczowy, 40 mg/ml, zawiesina doustna	DE/H/2343/001	17854	RECKITT BENCKISER (POLAND) S.A.	PL
Aktren® Forte 400 mg Filmtabletten	not available	13407.01.00	BAYER VITAL GMBH	DE
Momentact 400 mg capsule molli	not available	035618038	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
Momentact 400 mg capsule molli	not available	035618040	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ipren 125 mg suppositorier	not available	15684	MCNEIL SWEDEN AB	SE
Ipren, filmovertrukne tabletter	not available	13445	TAKEDA PHARMA A/S	DK
Ipren, gel 5%	not available	52074	TAKEDA PHARMA A/S	DK
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/23	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/24	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/05	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/21	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/06	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	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
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/20	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/08	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/22	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
MASIPREN 100 mg suspensie orală în plic	AT/H/0678/001	10393/2017/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
MASIPREN 100 mg suspensie orală în plic	AT/H/0678/001	10393/2017/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
MASIPREN 200 mg suspensie orală în plic	AT/H/0678/002	10394/2017/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
MASIPREN 200 mg suspensie orală în plic	AT/H/0678/002	10394/2017/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
MASIPREN 200 mg suspensie orală în plic	AT/H/0678/002	10394/2017/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
MASIPREN 400 mg suspensie orală în plic	AT/H/0678/003	10395/2017/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MASIPREN 400 mg suspensie orală în plic	AT/H/0678/003	10395/2017/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
BRUFEN 400 mg Filmtabletten	not available	BE104876	MYLAN EPD BVBA/SPRL	BE
BRUFEN 400 mg Filmtabletten	not available	BE467093	MYLAN EPD BVBA/SPRL	BE
BRUFEN FORTE 600 mg, Filmtabletten	not available	BE128064	MYLAN EPD BVBA/SPRL	BE
BRUFEN 400 mg, comprimés pelliculés	not available	BE467093	MYLAN EPD BVBA/SPRL	BE
BRUFEN 400 mg, comprimés pelliculés	not available	BE104876	MYLAN EPD BVBA/SPRL	BE
BRUFEN 400 mg, filmomhulde tabletten	not available	BE467093	MYLAN EPD BVBA/SPRL	BE
BRUFEN 400 mg, filmomhulde tabletten	not available	BE104876	MYLAN EPD BVBA/SPRL	BE
BRUFEN FORTE 600 mg, comprimés pelliculés	not available	BE128064	MYLAN EPD BVBA/SPRL	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
BRUFEN FORTE 600 mg, filmomhulde tabletten	not available	BE128064	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	BE147436	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	BE150376	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, comprimés à libération prolongée	not available	BE150376	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, comprimés à libération prolongée	not available	BE147436	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, tabletten met verlengde afgifte	not available	BE150376	MYLAN EPD BVBA/SPRL	BE
BRUFEN RETARD 800 mg, tabletten met verlengde afgifte	not available	BE147436	MYLAN EPD BVBA/SPRL	BE
Dalsy forte 40 mg/ml oralna suspenzija	not available	UP/I-530-09/12-01/625	MYLAN HRVATSKA D.O.O.	HR
Brufen 20 mg/ml sirup	not available	UP/I-530-09/12-01/602	MYLAN 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
DALSY 20 mg/ml sirup	not available	HR-H-407408529	MYLAN HRVATSKA D.O.O.	HR
Brufen Effect forte 40 mg/ml oralna suspenzija	not available	HR-H-330150785	MYLAN HRVATSKA D.O.O.	HR
Brufen 600 mg šumivé granule	not available	29/655/08-C	MYLAN IRE HEALTHCARE LIMITED	CZ
Moment 200 mg compresse masticabili	not available	025669096	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse masticabili	not available	025669108	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ibutop 50 mg/g cream	not available	20010485	DOLORGIET GMBH & CO KG	BG
Ibutop 50 mg/g gel	not available	20010680	DOLORGIET GMBH & CO KG	BG
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/005-009	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/007	PINEWOOD LABORATORIES LIMITED	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
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/006	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/005	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/009	PINEWOOD LABORATORIES LIMITED	LT
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/008	PINEWOOD LABORATORIES LIMITED	LT
Ibuprofen 200mg/5ml Oral Suspension	UK/H/3267/002/E/001	PA0281/088/005	PINEWOOD LABORATORIES LIMITED	IE
Ibuprofen 200mg/5ml Oral Suspension	IE/H/0690/001/DC	04917/0100	PINEWOOD LABORATORIES LIMITED	UK
Phorpain maximum strength 10% w/w Gel	not available	PL 10972/0090	MERCURY PHARMA GROUP LTD	UK
Fenbid 10 %w/w Gel	not available	PL 10972/0090	MERCURY PHARMA GROUP LTD	UK
Ibuleve Maximum Strength Gel	not available	PL 00173/0176	DIOMED DEVELOPMENTS 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
UPFEN 200 mg, comprimé pelliculé	not available	34009 332 158 4 2	UPSA SAS	FR
UPFEN 200 mg, comprimé pelliculé	not available	34009 347 795 5 8	UPSA SAS	FR
UPFEN 200 mg, comprimé pelliculé	not available	34009 331 861 3 5	UPSA SAS	FR
UPFEN 200 mg, comprimé pelliculé	not available	34009 332 159 0 3	UPSA SAS	FR
Ibuprofen Nutra Essential 200 mg perorální suspenze v sácku	CZ/H/0638/001	07/684/15-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra 200 mg perorálna suspenzia vo vrecku	CZ/H/0638/001	07/0092/18-S	NUTRA ESSENTIAL OTC, S.L.	SK
Winadol 200 mg belsőleges szuszpenzió tasakban	CZ/H/0638/001	OGYI-T-23323/01-03	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra 200 mg suspensie orala în plic	CZ/H/0638/001	10547/2018/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	CZ/H/0638/001	10547/2018/02	NUTRA ESSENTIAL OTC, S.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen Nutra 200 mg suspensie orală în plic	CZ/H/0638/001	10547/2018/03	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	CZ/H/0638/001	10547/2018/04	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	CZ/H/0638/001	10547/2018/05	NUTRA ESSENTIAL OTC, S.L.	RO
Winadol 200 mg belsőleges szuszpenzió tasakban	CZ/H/0638/001	OGYI-T-23323/01	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 200 mg belsőleges szuszpenzió tasakban	CZ/H/0638/001	OGYI-T-23323/02	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 200 mg belsőleges szuszpenzió tasakban	CZ/H/0638/001	OGYI-T-23323/03	NUTRA ESSENTIAL OTC, S.L.	HU
Aktren® Spezial 400 mg Weichkapseln	not available	37995.00.00	BAYER VITAL GMBH	DE
Ibukern 600 mg suspensión oral	not available	67.930	KERN PHARMA, S.L.	ES
Ibuleve Speed Relief Max Strength Gel 10% w/w	not available	PL 00173/0404	DIOMED DEVELOPMENTS 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
BRUFEN GRANULES 600 mg, Brausegranulat	not available	BE158067	MYLAN EPD BVBA/SPRL	BE
BRUFEN GRANULES 600 mg, granulés effervescents	not available	BE158067	MYLAN EPD BVBA/SPRL	BE
BRUFEN GRANULES 600 mg, bruisgranulaat	not available	BE158067	MYLAN EPD BVBA/SPRL	BE
Momentact 400 mg compresse rivestite con film	not available	035618053	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con film	not available	035618026	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con film	not available	035618014	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Brufen Retard 800 mg, tablet met gereguleerde afgifte	not available	RVG 13478	MYLAN HEALTHCARE B.V.	NL
Dolgit krém	not available	29/751/92-S/C	DOLORGIET GMBH & CO KG	CZ
Ibuprofen 600 mg Tablets BP	not available	PL 17496/0019	DALKEITH LABORATORIES 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
Ibuprofen 200 mg Coated Tablets	not available	PL 20395/0090	RELON CHEM LIMITED	UK
Ibuprofen Tablets BP 600 mg	not available	PL 20395/0132	RELON CHEM LIMITED	UK
Ibuprofen 100 mg/5 ml oral suspension	not available	PL 20395/0259	RELON CHEM LIMITED	UK
Ibuprofen 400mg Tablets	not available	PL 20395/0080	RELON CHEM LIMITED	UK
Tesco Health Ibuprofen 100 mg/5 ml Oral Suspension	not available	PL 20395/0259	RELON CHEM LIMITED	UK
Well Pharmaceuticals Ibuprofen 200mg Coated Tablets	not available	PL 20395/0090	RELON CHEM LIMITED	UK
Bell's Healthcare Children's Pain and Fever Relief 100 mg/5 ml Oral Suspension	not available	PL 20395/0259	RELON CHEM LIMITED	UK
Ibuprofen Sandoz 100 mg/5 ml suspensie voor oraal gebruik	DE/H/3341/001	BE428601	SANDOZ N.V.	BE
Ibuprofen Sandoz 20 mg/ml belsőleges szuszpenzió	DE/H/3341/001	OGYI-T-22367/01	SANDOZ HUNGÁRIA KFT	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
Ibuprofen Sandoz 20 mg/ml belsőleges szuszpenzió	DE/H/3341/001	OGYI-T-22367/02	SANDOZ HUNGÁRIA KFT	HU
Ibuprofen Sandoz 20 mg/ml belsőleges szuszpenzió	DE/H/3341/001	OGYI-T-22367/03	SANDOZ HUNGÁRIA KFT	HU
Ibuprofeno Sandoz 20 mg/ml, Suspensão oral	DE/H/3341/001	5489752	SANDOZ FARMACÊUTICA LDA.	PT
Flarin Joint & Muscular Pain Relief 200mg Soft Capsules	not available	PL 51724/0002	INFIRST LTD	UK
Ipren 5% gel	not available	18362	MCNEIL SWEDEN AB	SE
Ipren 5% gel	not available	18362	MCNEIL SWEDEN AB	SE
MIG 4% perorálna suspenzia 40 mg/ml perorálna suspenzia	DE/H/2598/002	29/0315/15-S	BERLIN-CHEMIE AG	SK
Berlistar forte 40 mg/ml oralna suspenzija	DE/H/2598/002	HR-H-140469047-01	BERLIN-CHEMIE AG	HR
Berlistar forte 40 mg/ml oralna suspenzija	DE/H/2598/002	HR-H-140469047-02	BERLIN-CHEMIE AG	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
Ibustar forte, 40 mg/ml suukaudne suspensioon	DE/H/2598/002	888015	BERLIN-CHEMIE AG	EE
МИГ джуниър 40 mg/ml перорална суспензия	DE/H/2598/002	20150332	BERLIN-CHEMIE AG	BG
Ibustar 40 mg/ml geriamoji suspensija, vaikams	DE/H/2598/002	LT/1/15/3829/001	BERLIN-CHEMIE AG	LT
Ibustar 40 mg/ml geriamoji suspensija, vaikams	DE/H/2598/002	LT/1/15/3829/002	BERLIN-CHEMIE AG	LT
Ibustar 40 mg/ml geriamoji suspensija, vaikams	DE/H/2598/002	LT/1/15/3829/003	BERLIN-CHEMIE AG	LT
Ibustar 40 mg/ml geriamoji suspensija, vaikams	DE/H/2598/002	LT/1/15/3829/004	BERLIN-CHEMIE AG	LT
Ibustar bērniem 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2598/002	15-0268	BERLIN-CHEMIE AG	LV
EUDORLIN Ibuprofen 40 mg/ml Suspension zum Einnehmen	DE/H/2598/002	79521.00.00	BERLIN-CHEMIE AG	DE
MIG dla dzieci Forte o smaku truskawkowym, 40 mg/ml, zawiesina doustna	DE/H/2598/002	21156	BERLIN-CHEMIE AG	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
doc Ibuprofen Schmerzgel, 5 % Gel	not available	39838.00.00	HERMES ARZNEIMITTEL GMBH	DE
Nurofen Junior Jarðarber 40 mg/ml mixtúra, dreifa	DE/H/2204/002	IS/1/11/119/01	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	IS
Nurofen 40 mg/ml mikstur, suspensjon med jordbærsmak	DE/H/2204/002	08-6425	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nureflex Junior Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2204/002	1-30062	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen Junior Fieber- und Schmerzsaft Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2204/002	76552.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Jordgubb, 40 mg/ml, oral suspension	DE/H/2204/002	41890	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	SE
Brufen Tablets 400mg	not available	PL 46302/0006	MYLAN PRODUCTS LIMITED	UK
Brufen Tablets 600 mg	not available	PL 46302/0009	MYLAN PRODUCTS LIMITED	UK
Brufen Tablets 200 mg	not available	PL 46302/0004	MYLAN PRODUCTS 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
ALGOFRENELLE®, Κολπικό διάλυμα 1% w/v	not available	80517/15	IOULIA AND IRENE TSETI PHARMACEUTICAL LABORATORIES S.A.	GR
Ibufarmalid 200 mg suspensión oral	not available	67.963	FARMALIDER, S.A.	ES
DOLOFAST® 10% Gel Tubo 50 G	not available	029775018	DOMPÉ FARMACEUTICI S.P.A.	IT
ANTALGIL 200 mg compresse	not available	027432020	WELCOME PHARMA SPA	IT
Ibuleve Speed Relief 5% Spray	not available	PL 00173/0160	DIOMED DEVELOPMENTS LIMITED	UK
Dalsy 20 mg/ml suspensión oral	not available	59166	MYLAN IRE HEALTHCARE LIMITED	ES
Moment 200 mg compresse rivestite	not available	025669122	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669146	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669161	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
Moment 200 mg compresse rivestite	not available	025669173	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669185	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669110	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669019	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite	not available	025669072	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ibuprofeno Clover 200 mg capsulas blandas	not available	80987	HC CLOVER PRODUCTOS Y SERVICIOS, S.L.	ES
IBUPROM EFFECT żel, 50 mg/g, żel	not available	23940	US PHARMACIA SP. Z O.O.	PL
Brufen Retard 800 mg comprimidos de libertação prolongada	not available	4338281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Alvofen Express 400 mg mjúk hylki	IS/H/0205/001	IS/1/13/050/02	ALVOGEN IPCO S.AR.L	IS

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
Ibufarmalid 400 mg suspensión oral	not available	68.200	FARMALIDER, S.A.	ES
Perdophen 400 mg, comprimés pelliculés	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN 400 mg, filmomhulde tabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 400 mg, comprimés pelliculés	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 400 mg, comprimés pelliculés	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN 400 mg, filmomhulde tabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN 400 mg, filmomhulde tabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN, 400 mg Filmtabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
PERDOPHEN, 400 mg Filmtabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
PERDOPHEN, 400 mg Filmtabletten	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Espidifen 600 mg granulado para solución oral sabor cola-limón	not available	80719	ZAMBON, S.A.U.	ES
Boots Max Strength Ibuprofen 10% Gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	UK
Nurofen Joint & Back Pain Relief Max Strength 10 % Gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	UK
Ibuprofen 10% w/w gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	UK
Brufen Retard, depottabletter	not available	13416	MYLAN DENMARK APS	DK
Ibuprofen 600 mg Tablets	not available	PL 20416/0293	CRESCENT PHARMA LIMITED	UK
Ibuprofen 200 mg Coated Tablets	not available	PL 03105/0086	BELL SONS & CO (DRUGGISTS) LTD	UK
Ibuprofen 200 mg Coated Tablets	not available	PL 03105/0102	BELL SONS & CO (DRUGGISTS) 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
Well Pharmaceuticals Ibuprofen 200mg Coated Tablets	not available	PL 03105/0086	BELL SONS & CO (DRUGGISTS) LTD	UK
Ibuprofen Sainsbury's 200mg Coated Tablets	not available	PL 03105/0086	BELL SONS & CO (DRUGGISTS) LTD	UK
Ibuprofen Sainsbury's 200mg Coated Tablets	not available	PL 03105/0102	BELL SONS & CO (DRUGGISTS) LTD	UK
BRUFEN 800 mg compresse rivestite a rilascio prolungato	not available	022593115	MYLAN ITALIA S.R.L.	IT
Ibu-LysinHEXAL	DE/H/2593/001	79780.00.00	HEXAL AG	DE
Ibugel Forte 10%	not available	PL 00173/0175	DIOMED DEVELOPMENTS LIMITED	UK
Ibuprofen Patheon 100 mg, zachte kauwcapsules	not available	RVG 123943	PATHEON SOFTGELS B.V.	NL
Aktren® 200 mg überzogene Tabletten	not available	7554.00.00	BAYER VITAL GMBH	DE
Aleve Ibuprofen 200 mg überzogene Tabletten	not available	16301.00.00	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
BRUFEN 400 mg Compresse rivestite	not available	022593204	MYLAN ITALIA S.R.L.	IT
BRUFEN 600 mg Compresse rivestite	not available	022593216	MYLAN ITALIA S.R.L.	IT
Dolgit, 50 mg /g , krem	not available	6443	DOLORGIET GMBH & CO KG	PL
Ibuprofen Nutra Essential 400 mg perorální suspenze v sácku	CZ/H/0639/002	07/687/15-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra Essential 400 mg suspension buvable en sachet	CZ/H/0639/002	BE518675	NUTRA ESSENTIAL OTC, S.L.	BE
STIBUPATCH 136 mg cerotto medicato	not available	037183011	ALFASIGMA S.P.A.	IT
STIBUPATCH 136 mg cerotto medicato	not available	037183023	ALFASIGMA S.P.A.	IT
Ibuprofen proff 5 % Gel Zur Anwendung bei Jugendlichen ab 14 Jahren und Erwachsenen Wirkstoff : Ibuprofen	not available	43767.00.00	DR. THEISS NATURWAREN GMBH	DE
Ibucalm 50 mg/g gel mentolado	not available	70.145	TEVA PHARMA 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
Ibuprofen AL direkt 200 mg Pulver zum Einnehmen	DE/H/4307/001	94256.00.00	ALIUD PHARMA GMBH	DE
Ibuprofen STADA 200 mg perorálny prášok	DE/H/4307/001	07/0226/17-S	STADA ARZNEIMITTEL AG	SK
Buprofess 200 mg pulbere orală	DE/H/4307/001	10209/2017/03	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	10209/2017/02	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	10209/2017/01	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	10209/2017/04	STADA M&D S.R.L.	RO
Ibuprofen STADA 200 mg peroralni prašek	DE/H/4307/001	H/17/02359/002	STADA ARZNEIMITTEL AG	SI
Ibuprofen STADA 200 mg peroralni prašek	DE/H/4307/001	H/17/02359/001	STADA ARZNEIMITTEL AG	SI
Ibuprofen STADA 200 mg peroralni prašek	DE/H/4307/001	H/17/02359/004	STADA ARZNEIMITTEL AG	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
Ibuprofen STADA 200 mg peroralni prašek	DE/H/4307/001	H/17/02359/003	STADA ARZNEIMITTEL AG	SI
Ibudolor Quick, 200 mg, proszek doustny	DE/H/4307/001	24331	STADA ARZNEIMITTEL AG	PL
Ibudolor Direkt 200 mg oralni prašak u vrećici	DE/H/4307/001	HR-H-054613410	STADA D.O.O.	HR
Ibuprofen Apotex 100 mg/5 ml perorální suspenze	SI/H/0162/001	29/308/12-C	APOTEX EUROPE B.V.	CZ
Nurofen for children 100 mg chewable capsules, soft	NL/H/4314/001	PA0979/069/001	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for children 100 mg, chewable capsules, soft	NL/H/4314/001	PL 00063/0721	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen voor Kinderen Sinaasappel 100 mg, kauwcapsules, zacht	NL/H/4314/001	RVG 112524	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/02	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/08	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/09	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/10	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/11	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/12	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/13	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/14	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/15	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	6146/2014/16	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen 200 mg Film-coated Tablets	not available	PL 44673/0084	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	UK
Ibuprofeno Sandoz 40 mg/ml suspensión oral	not available	65865	SANDOZ FARMACÉUTICA, S.A.	ES
Ibuprofen 400mg Tablets	not available	PL 00037/0674	ABBOTT LABORATORIES LIMITED	UK
Brufen Retard	not available	PL 46302/0010	MYLAN PRODUCTS LIMITED	UK
Ibuprofen LIDERFARM 40 mg/ml Suspension zum Einnehmen	not available	87683.00.00	FARMALIDER, S.A.	DE
Ibuprofen LIDERFARM 40 mg/ml Suspension zum Einnehmen	not available	88457.00.00	FARMALIDER, S.A.	DE
EFISEN M 200 mg, comprimate filmate	not available	2357/2010/01	S.C. SOLACIUM PHARMA S.R.L.	RO
Burana 100 mg pehmeät purukapselit	not available	32143	ORION OYJ	FI
Burana 100 mg mjuka tuggkapslar	not available	32143	ORION OYJ	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
Ibudolor Direkt 400 mg oralni prašak u vrecici	DE/H/4307/002	HR-H-991661386	STADA D.O.O.	HR
Ibuprofen AL direkt 400 mg Pulver zum Einnehmen	DE/H/4307/002	94257.00.00	ALIUD PHARMA GMBH	DE
Ibuprofen STADA 400 mg perorálny prášok	DE/H/4307/002	07/0227/17-S	STADA ARZNEIMITTEL AG	SK
Buprofess 400 mg pulbere orală	DE/H/4307/002	10210/2017/03	STADA M&D S.R.L.	RO
Buprofess 400 mg pulbere orală	DE/H/4307/002	10210/2017/02	STADA M&D S.R.L.	RO
Buprofess 400 mg pulbere orală	DE/H/4307/002	10210/2017/01	STADA M&D S.R.L.	RO
Ibuprofen STADA 400 mg peroralni prašek	DE/H/4307/002	H/17/02359/007	STADA ARZNEIMITTEL AG	SI
Ibuprofen STADA 400 mg peroralni prašek	DE/H/4307/002	H/17/02359/005	STADA ARZNEIMITTEL AG	SI
Ibuprofen STADA 400 mg peroralni prašek	DE/H/4307/002	H/17/02359/006	STADA ARZNEIMITTEL AG	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
Ibudolor Quick, 400 mg, proszek doustny	DE/H/4307/002	24332	STADA ARZNEIMITTEL AG	PL
Brupro Max 400mg Film-coated tablets	not available	PA0074/067/002	ROWA PHARMACEUTICALS LIMITED	IE
Brupro 200mg Film-coated tablets	not available	PA0074/067/001	ROWA PHARMACEUTICALS LIMITED	IE
Dalsy 400 mg comprimidos recubiertos con película	not available	54985	MYLAN IRE HEALTHCARE LIMITED	ES
Ibudol 50 mg/g gel	not available	62.054	KERN PHARMA, S.L.	ES
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	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
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Iprensa 400 mg kapslar, mjuka	not available	43284	MCNEIL SWEDEN AB	SE
Iprensa 400 mg kapslar, mjuka	not available	43284	MCNEIL SWEDEN AB	SE
IBUPAS 136 mg cerotto medicato	not available	036439014	ALFASIGMA S.P.A.	IT
IBUPAS 136 mg cerotto medicato	not available	036439026	ALFASIGMA S.P.A.	IT
Ibuprofen Sandoz 800 mg filmomhulde tabletten	not available	BE428382	SANDOZ N.V.	BE
Ibuprofen Sandoz 800 mg filmomhulde tabletten	not available	BE325211	SANDOZ N.V.	BE
Ibumetin 50mg/g Gel	not available	02-1301	TAKEDA AS	NO
Brufen 40 mg/ml Suspension zum Einnehmen	PT/H/2354/001	BE441847	MYLAN EPD BVBA/SPRL	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
Brufen 40 mg/ml suspension buvable	PT/H/2354/001	BE441847	MYLAN EPD BVBA/SPRL	BE
Brufen 40 mg/ml suspensie voor oraal gebruik	PT/H/2354/001	BE441847	MYLAN EPD BVBA/SPRL	BE
Бруфен Форте 40 mg/ml перорална суспензия	PT/H/2354/001	20130323	MYLAN EOOD	BG
Brufen, 40 mg/ml suukaudne suspensioon	PT/H/2354/001	820713	MYLAN IRE HEALTHCARE LIMITED	EE
Brufen cukormentes 40 mg/ml belsőleges szuszpenzió	PT/H/2354/001	OGYI-T-22073/03	MYLAN EPD KFT.	HU
Brufen cukormentes 40 mg/ml belsőleges szuszpenzió	PT/H/2354/001	OGYI-T-22073/04	MYLAN EPD KFT.	HU
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/002	MYLAN IRE HEALTHCARE LIMITED	LT
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/003	MYLAN IRE HEALTHCARE LIMITED	LT
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/004	MYLAN IRE HEALTHCARE LIMITED	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
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/001	MYLAN IRE HEALTHCARE LIMITED	LT
Brufedol 40 mg/ml suspensija iekšķīgai lietošanai	PT/H/2354/001	14-0134	MYLAN IRE HEALTHCARE LIMITED	LV
Brufen Sem Açúcar 40 mg/ml suspensão oral	PT/H/2354/001	5475819	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/001	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/002	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/003	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/004	MYLAN IRE HEALTHCARE LIMITED	SI
Brufen 40 mg/ml suspension buvable	PT/H/2354/001	2014040014	MYLAN EPD BVBA/SPRL	LU
Brufen Sem Açúcar 40 mg/ml suspensão oral	PT/H/2354/001	5442132	BGP PRODUCTS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufen 40 mg/ml Suspension zum Einnehmen	PT/H/2354/001	2014040014	MYLAN EPD BVBA/SPRL	LU
Brufen, filmovertrukne tabletter 400 mg	not available	06689	MYLAN DENMARK APS	DK
Brufen, filmovertrukne tabletter 600 mg	not available	10882	MYLAN DENMARK APS	DK
proff Schmerzgel	not available	1192.00.03	DOLORGIET GMBH & CO KG	DE
Dolgit 50 mg/g kreem	not available	147796	DOLORGIET GMBH & CO KG	EE
ADVIL 400 mg, comprimé enrobé	not available	34009 381 710 9 9	PFIZER SANTE FAMILIALE	FR
ADVILMED 100 mg, comprimé enrobé	not available	358 459-1	PFIZER SANTE FAMILIALE	FR
ADVIL 200 mg, comprimé enrobé	not available	34009 329 593 5 8	PFIZER SANTE FAMILIALE	FR
ADVILMED 400 mg, comprimé enrobé	not available	329 594-1	PFIZER SANTE FAMILIALE	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
ADVILMED 400 mg, comprimé enrobé	not available	329 595-8	PFIZER SANTE FAMILIALE	FR
ADVIL 400 mg, comprimé enrobé	not available	34009 381 711 5 0	PFIZER SANTE FAMILIALE	FR
ADVIL 200 mg, comprimé enrobé	not available	34009 332 315 2 1	PFIZER SANTE FAMILIALE	FR
Dolgit gél	not available	5285/01	DOLORGIET GMBH & CO KG	HU
Ibuprofeno Clover 600 mg capsulas blandas	not available	80976	HC CLOVER PRODUCTOS Y SERVICIOS, S.L.	ES
Ibugel	not available	PL 00173/0050	DIOMED DEVELOPMENTS LIMITED	UK
Ibuprofen Banner 100 mg, zachte kauwcapsules	NL/H/3903/001	RVG 112525	PATHEON SOFTGELS B.V.	NL
Ibuprofen Banner 100 mg, soft chewable capsules	NL/H/3903/001	MA1328/00101	PATHEON SOFTGELS B.V.	MT
Ibuprofen Banner 100 mg, soft chewable capsules	NL/H/3903/001	PA22773/001/001	PATHEON SOFTGELS B.V.	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
Ibuprofen Banner 100 mg meke kapsule za žvakanje	NL/H/3903/001	HR-H-911536382	PATHEON SOFTGELS B.V.	HR
Ibuprofen Banner 100 mg mīkstās košļājamās kapsulas	NL/H/3903/001	19-0200	PATHEON SOFTGELS B.V.	LV
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/01	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/02	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/03	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/04	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/05	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/06	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/07	PATHEON SOFTGELS B.V.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/08	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/09	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/10	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/11	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/12	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/13	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/14	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/15	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/16	PATHEON SOFTGELS B.V.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/17	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg Weichkapseln zum Zerbeißen	NL/H/3903/001	139259	PATHEON SOFTGELS B.V.	AT
Ibuprofen Banner 100 mg měkké žvýkací tobolky	NL/H/3903/001	07/125/19-C	PATHEON SOFTGELS B.V.	CZ
Ibuprofen Banner 100 mg Weichkapseln zum Zerbeißen	NL/H/3903/001	2203445.00.00	PATHEON SOFTGELS B.V.	DE
Pathofen 100 mg, zachte kauwcapsules	NL/H/4843/001	RVG 123962	PATHEON SOFTGELS B.V.	NL
Pathofen 100 mg Weichkapseln zum Zerbeißen	NL/H/4843/001	2204317.00.00	PATHEON SOFTGELS B.V.	DE
Ozonol 200 mg comprimidos revestidos	not available	8782235	PERRIGO PORTUGAL LDA	PT
Ozonol 200 mg comprimidos revestidos	not available	8782227	PERRIGO PORTUGAL LDA	PT
Ozonol 200 mg comprimidos revestidos	not available	8782243	PERRIGO PORTUGAL LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ozonol 200 mg comprimidos revestidos	not available	5467725	PERRIGO PORTUGAL LDA	PT
Ozonol 200 mg comprimidos revestidos	not available	8782219	PERRIGO PORTUGAL LDA	PT
Ibuprofen 200mg Tablets	not available	PL 31308/0013	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen 200mg Tablets	not available	PL 31308/0013	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen 200mg Tablets	not available	PL 31308/0016	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen 400mg film-coated Tablets	not available	PL 31308/0040	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen 200mg Tablets	not available	PL 31308/0014	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen 400mg Tablets	not available	PL 31308/0035	BRUNEL HEALTHCARE MANUFACTURING LIMITED	UK
Ibuprofen Sandoz 40 mg/ml suspensie voor oraal gebruik	DE/H/3341/002	BE428617	SANDOZ N.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
Ibux 600 mg tabletter	not available	7069	KARO PHARMA AS	NO
Algiasdin retard 600 mg comprimidos de liberación prolongada	not available	61609	ISDIN, S.A.	ES
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386012	SANOFI S.P.A	IT
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386036	SANOFI S.P.A	IT
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386024	SANOFI S.P.A	IT
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386063	SANOFI S.P.A	IT
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386051	SANOFI S.P.A	IT
Ibuprofene Sanofi 200 mg polvere orale in bustina	CZ/H/0654/001	045386048	SANOFI S.P.A	IT
Moment Orosolubile 200 mg polvere orale	IT/H/0745/001	046054019	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
Moment Orosolubile 200 mg polvere orale	IT/H/0745/001	046054021	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment Orosolubile 200 mg polvere orale	IT/H/0745/001	046054033	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment Orosolubile 400 mg polvere orale	IT/H/0745/002	046054045	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment Orosolubile 400 mg polvere orale	IT/H/0745/002	046054058	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment Orosolubile 400 mg polvere orale	IT/H/0745/002	046054060	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ibudol 400 mg suspensión oral	not available	68194	KERN PHARMA, S.L.	ES
Brufen Bruis 600 mg, bruisgranulaat	not available	RVG 13331	MYLAN HEALTHCARE B.V.	NL
Ibuprofen 400 mg Solution For Infusion	PT/H/1862/001	PL 08553/0619	DR. REDDY'S LABORATORIES (UK) LTD.	UK
Ibuprofen 600 mg Solution For Infusion	PT/H/1862/002	PL 08553/0632	DR. REDDY'S LABORATORIES (UK) 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
Ibuflam® Schmerzgel Wirkstoff: 5 % Ibuprofen	not available	13563.00.01	CHEMISCHE FABRIK KREUSSLER & CO. GMBH	DE
Ibuprofen 100mg/5ml Oral Suspension	not available	04917/0044	PINEWOOD LABORATORIES LIMITED	UK
Calprofen 100mg/5ml Oral Suspension Ibuprofen	not available	PL 15513/0147	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324032	ZENTIVA ITALIA SRL	IT
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324018	ZENTIVA ITALIA SRL	IT
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324020	ZENTIVA ITALIA SRL	IT
Rapidophen 40 mg/ml belsőleges szuszpenzió	not available	OGYI-T-22440/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Rapidophen 40 mg/ml belsőleges szuszpenzió	not available	OGYI-T-22440/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Rapidophen 40 mg/ml belsőleges szuszpenzió	not available	OGYI-T-22440/04	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
Rapidophen 40 mg/ml belsőleges szuszpenzió	not available	OGYI-T-22440/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Ibuprofen Pliva 400 mg filmom obložene tablete	not available	HR-H-948734629	PLIVA HRVATSKA D.O.O.	HR
Ibuxin Rapid 684 mg filmom obložene tablete	not available	HR-H-064892441	PLIVA HRVATSKA D.O.O.	HR
Boots Ibuprofen 3 Months Plus 100mg/5ml Oral Suspension Strawberry Flavour	not available	PL 00014/0859	THE BOOTS COMPANY PLC	UK
Ibuprofen 600mg tablets	not available	PL 00037/0675	ABBOTT LABORATORIES LIMITED	UK
Nurofen dla dzieci Forte truskawkowy, 40 mg/ml, zawiesina doustna	DE/H/2206/002	22449	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/025	RECKITT BENCKISER (POLAND) S.A.	LT
Nuroflex cu aroma de capsuni 40 mg/ml suspensie orala	DE/H/2206/002	9341/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de capsuni 40 mg/ml suspensie orala	DE/H/2206/002	9341/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nuroflex cu aroma de capsuni 40 mg/ml suspensie orala	DE/H/2206/002	9341/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de capsuni 40 mg/ml suspensie orala	DE/H/2206/002	9341/2016/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nuroflex cu aroma de capsuni 40 mg/ml suspensie orala	DE/H/2206/002	9341/2016/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Нурофен за Деца Форте Ягода 200 mg/5 ml перорална суспензия	DE/H/2206/002	20100671	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Forte Strawberry, 40 mg/ml, suukaudne suspensioon	DE/H/2206/002	707210	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Junior Fieber- und Schmerzsaft Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2206/002	76554.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen for Children Strawberry 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2206/002	10-0477	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/017	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/018	RECKITT BENCKISER (POLAND) S.A.	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
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/019	RECKITT BENCKISER (POLAND) S.A.	LT
Junipro 40 mg/ml suspensión oral sabor fresa	DE/H/2206/002	73048	RECKITT BENCKISER HEALTHCARE S.A.	ES
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/020	RECKITT BENCKISER (POLAND) S.A.	LT
proff Schmerzcreme	not available	0473/85/03/0258	DOLORGIET GMBH & CO KG	LU
Brufen Granules	not available	PL 46302/0007	MYLAN PRODUCTS LIMITED	UK
Nurofen Immedia 400 mg Weichkapseln	NL/H/4313/001/DC	78463.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen ® Express 400 mg, καψάκιο, μαλακό	NL/H/4313/001/DC	21486	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/03	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofencaps 400 mg, capsule molle	NL/H/4313/001	NL38369	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen rapid 400 mg Weichkapseln	NL/H/4313/001/DC	1-31144	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen Fastine Liquid Caps 400 mg, capsule, zacht	NL/H/4313/001	RVG 105812	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Express Forte, 400 mg, kapsułki, miękkie	NL/H/4313/001	19888	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001/DC	5437330	RECKITT BENCKISER HEALTHCARE, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001/DC	5437348	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001/DC	5437355	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001/DC	5437363	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 400 mg Capsule, soft	NL/H/4313/001	PL 00063/0615	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Нурофен Експрес Форте 400 mg капсули, меки Nurofen Express Forte 400 mg capsules, soft	NL/H/4313/001/DC	20120227	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001/DC	OGYI-T-6793/110	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001/DC	OGYI-T-6793/111	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001/DC	OGYI-T-6793/112	RECKITT BENCKISER KFT	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
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001/DC	OGYI-T-6793/113	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001/DC	OGYI-T-6793/114	RECKITT BENCKISER KFT	HU
Nurofencaps 400 mg, capsule molle	NL/H/4313/001	34009 221 338 4 3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofencaps 400 mg, capsule molle	NL/H/4313/001	34009 221 334 9 2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofencaps 400 mg, capsule molle	NL/H/4313/001	34009 221 336 1 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFENCAPS 400 mg, capsule molle	NL/H/4313/001	34009 221 339 0 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFENCAPS 400 mg, capsule molle	NL/H/4313/001	34009 221 335 5 3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFENCAPS 400 mg, capsule molle	NL/H/4313/001	34009 221 337 8 2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860014	RECKITT BENCKISER HEALTHCARE (ITALIA) 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
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860026	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860038	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860040	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860053	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860115	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860065	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860077	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860089	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860127	RECKITT BENCKISER HEALTHCARE (ITALIA) 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
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860091	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001/DC	041860103	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen pour Enfants100 mg, capsules molles à mâcher	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen voor Kinderen 100 mg, zachte kauwcapsules	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder 100 mg, Weichkapseln zum Zerbeissen	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 100 mg tyggekapsler, myke	UK/H/5966/001	15-10700	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nurofen 100 mg purukapseli, pehmeä	UK/H/5966/001	33426	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	FI
Nurofen pour Enfants 100 mg, capsules molles à mâcher	BE/H/0320/001	2017070240	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen 100 mg tuggkapsel, mjuk	UK/H/5966/001	52973	RECKITT BENCKISER NORDIC A/S	SE

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
Nurofen for children 100 mg, μασώμενα καψάκια, μαλακά Πορτοκάλι	BE/H/0320/001	35066/28-3-2019	RECKITT BENCKISER HELLAS CHEMICALS ABEE	GR
Nurofen for children 100 mg, μασώμενα καψάκια, μαλακά Πορτοκάλι	BE/H/0320/001	022496	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Nurofen Junior Appelsín 40 mg/ml mixtúra, dreifa	DE/H/2204/001	IS/1/11/118/01	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	IS
Nurofen 40 mg/ml mikstur, suspensjon med appelsinsmak	DE/H/2204/001	08-6424	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nureflex Junior Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2204/001	1-30061	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2204/001	76551.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Apelsin, 40 mg/ml, oral suspension	DE/H/2204/001	41889	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	SE
Fennings Ibuprofen For Children 100mg/5ml Oral Suspension	not available	PL 20416/0535	CRESCENT PHARMA LIMITED	UK
Ibuprofen 100mg/5ml Oral Suspension	not available	PL 20416/0535	CRESCENT PHARMA 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
Ibuprofène Mylan 200 mg, comprimé enrobé	not available	NL 26874	MYLAN S.A.S	FR
Ibuleve Speed Relief Gel	not available	PL 00173/0060	DIOMED DEVELOPMENTS LIMITED	UK
ANTARENE 5 %, gel	not available	34009 355 627 0 8	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
Dalsy 40 mg/ml suspensión oral	not available	69726	MYLAN IRE HEALTHCARE LIMITED	ES
Moment 200 mg sospensione orale in bustine.	not available	025669336	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669348	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669351	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669363	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669375	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
Moment 200 mg sospensione orale in bustine.	not available	025669387	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669399	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669401	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg sospensione orale in bustine.	not available	025669413	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669223	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669235	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669247	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669250	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669262	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
Moment 200 mg compresse orodispersibili	not available	025669274	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669286	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669298	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669300	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669312	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse orodispersibili	not available	025669324	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Perdofemina 400 mg, comprimés pelliculés	not available	2008099940	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Perdofemina 400 mg, comprimés pelliculés	not available	2008099940	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Perdofemina 400 mg, comprimés pelliculés	not available	2008099940	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
DIMIDON 400 mg, comprimidos revestidos por película	not available	5564034	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por película	not available	5564042	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por película	not available	5564059	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por película	not available	5564026	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
ANTARENE 5 %, gel	not available	34009 355 627 0 8	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
Dalsy 200 mg granulado efervescente	not available	63990	MYLAN IRE HEALTHCARE LIMITED	ES
Ibuprofen (als lysine) Mylan 400 mg, filmomhulde tabletten	NL/H/2576/002	RVG 111365	MYLAN B.V.	NL
Ibuprofen (als lysine) Mylan 200 mg, filmomhulde tabletten	NL/H/2576/001	RVG 111364	MYLAN B.V.	NL
Brufedol Rapid 200 mg filmom obalené tablety	NL/H/2576/001	29/0363/13-S	MYLAN IRELAND LIMITED	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brufedol Rapid 400 mg filmom obalené tablety	NL/H/2576/002	29/0364/13-S	MYLAN IRELAND LIMITED	SK
BRUFEN RAPID 400 mg potahované tablety	NL/H/2576/002	29/407/13-C	MYLAN IRELAND LIMITED	CZ
BRUFEN RETARD 800 mg filmom obalené tablety s predĺženým uvoľňovaním	not available	29/0659/10-S	MYLAN IRE HEALTHCARE LIMITED	SK
BRUFEN sirup 100 mg/5 ml sirup	not available	29/0916/92-S	MYLAN IRE HEALTHCARE LIMITED	SK
Brufen® 400 mg - Filmtabletten	not available	16233	MYLAN ÖSTERREICH GMBH	AT
Brufen® 600 mg - Filmtabletten	not available	1-18151	MYLAN ÖSTERREICH GMBH	AT
Spalt Forte 400 mg Weichkapseln	not available	9032.01.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
Spalt Migräne 400 mg Weichkapseln	not available	12732.00.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
Spalt Mobil 400 mg Weichkapseln	not available	9632.01.00	PFIZER CONSUMER HEALTHCARE 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
Spalt Kopfschmerz 200 mg Weichkapseln	not available	16823.00.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
Brufen 400mg film-coated tablets	not available	PA 2010/2/1	MYLAN IRE HEALTHCARE LIMITED	IE
Brufen 600mg film-coated tablets	not available	PA 2010/2/2	MYLAN IRE HEALTHCARE LIMITED	IE
FASPIC 200, 200 mg granulado para solução oral	not available	5220686	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 200, 200 mg granulado para solução oral	not available	5471388	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 200, 200 mg granulado para solução oral	not available	5220785	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 200, 200 mg granulado para solução oral	not available	5220884	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 400, 400 mg granulado para solução oral	not available	5220983	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 400, 400 mg granulado para solução oral	not available	5471487	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
FASPIC 400, 400 mg granulado para solução oral	not available	5221080	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 400, 400 mg granulado para solução oral	not available	5221189	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 600, 600 mg granulado para solução oral	not available	5221288	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 600, 600 mg granulado para solução oral	not available	5471586	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 600, 600 mg granulado para solução oral	not available	5221387	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
FASPIC 600, 600 mg granulado para solução oral	not available	5221486	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Brufen 400 mg potahované tablety	not available	29/390/92/-S/C	MYLAN IRE HEALTHCARE LIMITED	CZ
Ibuprofen 200mg Tablets	not available	PL 43461/0004	FLAMINGO PHARMA UK LTD	UK
Ibuprofen 200mg Tablets BP	not available	PL 43461/0007	FLAMINGO PHARMA UK 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
Ibuprofen 400mg Tablets BP	not available	PL 43461/0005	FLAMINGO PHARMA UK LTD	UK
EDENIL 0,1 g soluzione vaginale	not available	027741040	TEOFARMA S.R.L.	IT
EDENIL 0,1 g soluzione vaginale	not available	027741014	TEOFARMA S.R.L.	IT
EDENIL 1 g polvere per soluzione vaginale	not available	027741038	TEOFARMA S.R.L.	IT
GINENORM 0,1 g SOLUZIONE VAGINALE	not available	029135023	AESCULAPIUS FARMACEUTICI S.R.L.	IT
GINENORM 1 g POLVERE PER SOLUZIONE VAGINALE	not available	029135011	AESCULAPIUS FARMACEUTICI S.R.L.	IT
GINENORM 0,1 g SOLUZIONE VAGINALE	not available	029135035	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Solibu 400 mg Solução para perfusão	PT/H/1732/001	5747266	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Ibuprofene Dr. Reddy's 400 mg soluzione per infusione	PT/H/1732/001	045826017	DR. REDDY'S 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
Ibuprofene Dr. Reddy's 400 mg soluzione per infusione	PT/H/1732/001	045826029	DR. REDDY'S S.R.L.	IT
Ibuprofene Dr. Reddy's 600 mg soluzione per infusione	PT/H/1732/002	045826031	DR. REDDY'S S.R.L.	IT
Ibuprofene Dr. Reddy's 600 mg soluzione per infusione	PT/H/1732/002	045826043	DR. REDDY'S S.R.L.	IT
Peribu 400 mg solución para perfusión.	PT/H/1732/001	84556	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	ES
Solibu 400 mg Solução para perfusão	PT/H/1732/001	5747258	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Solibu 600 mg Solução para perfusão	PT/H/1732/002	5747274	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Solibu 600 mg Solução para perfusão	PT/H/1732/002	5747308	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Solibu 400 mg Infusionslösung	PT/H/1732/001	98473.00.00	ALTAN PHARMA LTD	DE
Solibu 600 mg Infusionslösung	PT/H/1732/002	2201636.00.00	ALTAN PHARMA LTD	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
Solibu 600 mg solución para perfusión.	PT/H/1732/002	83680	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	ES
Ibuprofen Hermes, 50 mg/g, Gel	not available	43768.00.00	HERMES ARZNEIMITTEL GMBH	DE
Ibuprofen 200mg Tablets	not available	PL 17907/0078	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen 200mg Caplets	not available	PL 17907/0077	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen 200mg tablets	not available	PL 17907/0505	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen 200 mg Tablets	not available	PL 17907/0077	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen 200 mg Caplets	not available	PL 17907/0078	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen Tablets 400 mg.	not available	PL 17907/0079	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Ibuprofen 200mg tablets	not available	PL 17907/0159	BRISTOL LABORATORIES LTD (BERKHAMSTED)	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
Ibuprofen 200 mg tablets	not available	PL 17907/0160	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Extra Strength Ibuprofen 400mg Tablets	not available	PL 17907/0161	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Nurofen Rapid Forte 400 mg meke kapsule	not available	HR-H-023698858	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 400 mg obložene tablete	not available	UP/I-530-09/12-01/127	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 200 Fastcaps, 200 mg capsules, zacht	not available	BE442346	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 200 Fastcaps, 200 mg capsules molles	not available	BE442346	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 200 Fastcaps 200 mg Weichkapseln	not available	BE442346	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen for Children Cold & Flu & Pain Orange 100mg/5ml Oral Suspension	not available	PA 979/66/01	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for Children Cold & Flu & Pain Strawberry 100mg/5ml Oral Suspension	not available	PA 979/67/01	RECKITT BENCKISER IRELAND LTD.	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
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero	not available	034102386	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero	not available	034102398	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero	not available	034102400	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero	not available	034102412	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102424	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102436	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102448	RECKITT BENCKISER HEALTHCARE (ITALIA) 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
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102451	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/03	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/12	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/101	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/102	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen 200mg Liquid Capsules	not available	PA 979/32/12	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Rapid Relief Maximum Strength 400mg Liquid Capsules	not available	PA 979/32/13	RECKITT BENCKISER IRELAND LTD.	IE
NUREFLEX 200 mg, comprimé enrobé	not available	339 861-2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 200 mg, comprimé enrobé	not available	360 919-6	RECKITT BENCKISER HEALTHCARE 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
NUREFLEX 200 mg, comprimé enrobé	not available	360 920-4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	339 643-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	384 142-1 OU 34009 384 142 1 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	360 763-6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	360 917-3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 647-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 648-1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 649-8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 650-6	RECKITT BENCKISER HEALTHCARE 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
NUROFEN 400 mg, comprimé enrobé	not available	368 651-2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 652-9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 653-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 654-1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	381 420-0 OU 34009 381 420 0 6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	381 419-2 OU 34009 381 419 2 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	381 421-7 OU 34009 381 421 7 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN FEBBRE E DOLORE Bambini 100mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102020	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen Express 684mg Caplets	not available	PL 00063/0384	RECKITT BENCKISER HEALTHCARE (UK) 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
NUREFLEX 400 mg, comprimé enrobé	not available	364 936-2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 937-9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 938-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 939-1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 948-0	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 947-4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	341 684-7	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 945-1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 946-8	RECKITT BENCKISER HEALTHCARE 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
NUREFLEX 400 mg, comprimé enrobé	not available	364 943-9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 944-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 941-6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	364 942-2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen narancsízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/99	RECKITT BENCKISER KFT	HU
Nurofen narancsízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/100	RECKITT BENCKISER KFT	HU
Nurofen for Children Orange Baby	not available	MA096/01102	RECKITT BENCKISER HEALTHCARE (UK) LTD	MT
Nurofen for Children 3 months to 9 years Orange	not available	MA096/01102	RECKITT BENCKISER HEALTHCARE (UK) LTD	MT
NUROFEN 200 Soditabs 200 mg comprimés enrobés	not available	2014090203	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
NUROFEN 200 Fastcaps, 200 mg capsules molles	not available	2014090205	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400 Soditabs 400 mg comprimés enrobés	not available	2014090204	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen Express 200 mg drajeuri	not available	12930/2020/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/07	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Express 200 mg drajeuri	not available	12930/2020/08	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/09	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/10	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	6139/2014/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	6139/2014/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	6139/2014/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	6139/2014/05	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN 200 mg drajeuri	not available	8883/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/04	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/05	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/06	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN 200 mg drajeuri	not available	8883/2016/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Forte 400 mg drajeuri	not available	8889/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Forte 400 mg drajeuri	not available	8889/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN PENTRU COPII cu aromă de căpşuni 100 mg/5 ml suspensie orală	not available	9005/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN PENTRU COPII cu aromă de portocale 100 mg/5 ml suspensie orală	not available	9006/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN PENTRU COPII cu aromă de portocale 100 mg/5 ml suspensie orală	not available	9006/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Long Lasting Pain Relief 300mg Prolonged Release Capsules	not available	PL 00063/0378	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen 200mg Dragees	not available	14712	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen rapid 400 mg - Filmtabletten	not available	1-23773	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen 200 mg - Dragees	not available	14712	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen rapid 400 mg Filmtabletten	not available	1-23773	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
NUROFEN VOOR KINDEREN 200 mg omhulde tabletten	not available	BE189935	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR ENFANTS 200 mg comprimés enrobés.	not available	BE189935	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder 200 mg überzogene Tabletten	not available	BE189935	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
NUROFEN 200, 200 mg omhulde tabletten	not available	BE132011	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 200, 200 mg comprimés enrobés	not available	BE132011	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 200, 200 mg überzogene Tabletten	not available	BE132011	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 Fastcaps, 400 mg capsules, zacht	not available	BE305724	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 Fastcaps, 400 mg capsules molles	not available	BE305724	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 400 Fastcaps 400 mg Weichkapseln	not available	BE305724	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Cuprofen Tablets 400 mg	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Ibuprofen Tablets BP 400 mg	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Cuprofen Maximum Strength Tablets	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) 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
UniChem Extra Strength Ibuprofen 400mg Tablets	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
NUROFEN 400, 400 mg omhulde tabletten	not available	BE189926	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN VOOR KINDEREN SUIKERVRIJ 2% suspensie voor oraal gebruik.	not available	BE191021	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR ENFANTS SANS SUCRE 2% suspension buvable	not available	BE191021	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder zuckerfrei 2% Suspension zum Einnehmen	not available	BE191021	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Нурофен за Деца с вкус на Портокал 100 mg/5 ml перорална суспензия Nurofen for Children Orange 100 mg/5 ml oral suspension	not available	20000158	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen For Children	not available	13609	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Nurofen Junior Fiebersaft Orange 20 mg/ml Suspension zum Einnehmen	not available	39181.00.00	RECKITT BENCKISER DEUTSCHLAND 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
Nurofen for Children Orange 100mg/5ml Oral Suspension	not available	PA 979/32/1	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Febbre e Dolore Bambini 100mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102018	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen Febbre e Dolore Bambini 100mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102020	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN bērniem 100 mg/5 ml suspensija iekšķīgai lietošanai	not available	01-0450	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen for children Orange	not available	MA096/01102	RECKITT BENCKISER HEALTHCARE (UK) LTD	MT
Nurofen voor Kinderen suikervrije suspensie	not available	RVG 19838	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen dla dzieci, 100 mg/5 ml, zawiesina doustna	not available	4567	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen For Children Orange Singles	not available	PL 00063/0669	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Orange	not available	PL 00063/0668	RECKITT BENCKISER (HEALTHCARE) UK 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
Nurofen for Children 3 months to 9 years Orange	not available	PL 00063/0668	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Orange Baby	not available	PL 00063/0668	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/79	RECKITT BENCKISER KFT	HU
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/80	RECKITT BENCKISER KFT	HU
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/81	RECKITT BENCKISER KFT	HU
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/82	RECKITT BENCKISER KFT	HU
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/83	RECKITT BENCKISER KFT	HU
Nurofen 125 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/08	RECKITT BENCKISER KFT	HU
Nurofen 125 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/09	RECKITT BENCKISER KFT	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
Nurofen 125 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/10	RECKITT BENCKISER KFT	HU
Nurofen 125 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/11	RECKITT BENCKISER KFT	HU
Nurofen for Children Meltlets 100mg Orodispersible Tablets	not available	PA 979/32/5	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Active 200 mg orodispersible tablets (Нурофен Актив 200 mg таблетки, диспергиращи се в устата)	not available	20030694	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Non-Aqua 200 mg szájban diszpergálódó tabletta	not available	OGYI-T-6793/84	RECKITT BENCKISER KFT	HU
Nurofen Meltlets Lemon	not available	PL 00063/0382	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen 200mg Liquicaps Pharmacy Only	not available	PL 00063/0654	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Express 200mg Liquid Capsules	not available	PL 00063/0654	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Express Period Pain 200mg soft capsules	not available	PL 00063/0646	RECKITT BENCKISER HEALTHCARE (UK) 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
Ibuprofen 200mg Liquicaps	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen Express 200mg Liquid Capsules	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen 200mg Liquid Capsules	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Нурофен 200 mg обвити таблетки	not available	9800355	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen 200 mg obalené tablety	not available	07/376/92-S/C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen, 200 mg kaetud tabletid	not available	218498	RECKITT BENCKISER (POLAND) S.A.	EE
NUREFLEX 200 mg, comprimé enrobé	not available	339 860-6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	339 642-9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/67	RECKITT BENCKISER KFT	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
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/68	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/69	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/70	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/71	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/72	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/73	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/74	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/91	RECKITT BENCKISER KFT	HU
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/92	RECKITT BENCKISER KFT	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
Nurofen 200mg Coated Tablets	not available	PA 979/32/6	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN 200 mg compresse rivestite	not available	025634015	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN 200 mg compresse rivestite	not available	025634041	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN 200 mg compresse rivestite	not available	025634092	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen 200 mg apvalkotās tabletes	not available	01-0449	RECKITT BENCKISER (POLAND) S.A.	LV
NUROFEN 200 mg dengtos tabletės	not available	LT/1/97/0271/003	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen 200 tablet, omhulde tabletten 200 mg	not available	RVG 10674	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen, 200 mg, tabletki powlekane	not available	R/7068	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen 200 mg comprimidos revestidos	not available	4348686	RECKITT BENCKISER HEALTHCARE, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen 200 mg comprimidos revestidos	not available	4348785	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 200 mg comprimidos revestidos	not available	4348884	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 200 mg comprimidos revestidos	not available	4348983	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 200 mg obalené tablety	not available	29/0376/92-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen 200 mg Tablets	not available	PL 00063/0388	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen 200 mg Tablets	not available	PL 00063/0385	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
NUROFEN LYSINE 200 mg, filmomhulde tabletten	not available	BE239005	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN LYSINE 200 mg, comprimés pelliculés	not available	BE239005	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen Lysine 200 mg Filmtabletten	not available	BE239005	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
NUROFEN 400 FASTTABS 400 mg, filmomhulde tabletten.	not available	BE238551	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 FASTTABS 400 mg, comprimés pelliculés.	not available	BE238551	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 400 Fasttabs 400mg, Filmtabletten	not available	BE238551	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 Soditabs 400 mg omhulde tabletten	not available	BE415292	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 Soditabs 400 mg comprimés enrobés	not available	BE415292	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 400 Soditabs 400 mg überzogene Tabletten	not available	BE415292	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen Migraine Pain 342 mg film-coated tablets (Нурофен за Мигрена 342 mg филмиранитаблетки)	not available	20070006/07.03.2012	RECKITT BENCKISER (ROMANIA) SRL	BG
Нурофен Фемина Форте 512 mg обвити таблетки	not available	20100648	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Neo 200 mg Obalené tablety	not available	07/812/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN NEO FEMINA 400 mg obalené tablety	not available	07/813/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen Forte Express, 400mg kaetud tabletid	not available	743111	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Immedia 400 mg Filmtabletten	not available	43917.01.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/41	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/44	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/47	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/50	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/53	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/56	RECKITT BENCKISER KFT	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
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/59	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/62	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte bevont tabletta	not available	OGYI-T-6793/65	RECKITT BENCKISER KFT	HU
Nurofen Express Maximum Strength 400 mg Tablets	not available	PA 979/32/11	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Forte Express 400 mg apvalkotās tabletes	not available	08-0205	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Omhulde Tabletten 200, omhulde tabletten 200 mg	not available	RVG 23518	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen 400 tablet Female, omhulde tablet 400 mg	not available	RVG 29736	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen 400 tablet Migraine, omhulde tablet 400 mg	not available	RVG 29737	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Flextin 200 mg tablet ovaal, omhulde tabletten	not available	RVG 100659	RECKITT BENCKISER HEALTHCARE B.V.	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
Nurofen Flextin 400 mg tablet, omhulde tabletten	not available	RVG 100657	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Forte Express, 400mg tabletki powlekane	not available	14316	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Musc 400 mg comprimidos revestidos	not available	5296678	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Musc 400 mg comprimidos revestidos	not available	5296702	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Musc 400 mg comprimidos revestidos	not available	5296728	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Expres 400 mg Obalené tablety	not available	07/0063/09-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen Migraine Pain	not available	PL 00063/0380	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Tension Headache	not available	PL 00063/0380	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Express 342mg Caplets	not available	PL 00063/0380	RECKITT BENCKISER HEALTHCARE (UK) 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
Nurofen Maximum Strength Migraine Pain 684mg Caplets	not available	PL 00063/0384	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Pain Relief Caplets 256mg Tablets	not available	00063/0411	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen Express 256 mg Caplets	not available	00063/0374	RECKITT BENCKISER (UK) LIMITED	UK
Nurofen 400 tablet, omhulde tabletten 400 mg	not available	RVG 22100	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Forte 400mg tabletki powlekane	not available	4937	RECKITT BENCKISER (POLAND) S.A.	PL
NUROFEN 400, 400 mg comprimidos revestidos	not available	4283586	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4283685	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4284386	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4283784	RECKITT BENCKISER HEALTHCARE, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN 400, 400 mg comprimidos revestidos	not available	4283883	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4283982	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4284089	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4284188	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN 400, 400 mg comprimidos revestidos	not available	4284287	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN Rapid 400 mg Capsules mäkké kapsuly	not available	29/0302/08-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen 400 mg obalené tablety	not available	29/0054/13-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
NUROFEN 5 %, gel	not available	357 295-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/04	RECKITT BENCKISER KFT	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
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/05	RECKITT BENCKISER KFT	HU
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/06	RECKITT BENCKISER KFT	HU
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/07	RECKITT BENCKISER KFT	HU
Nurofen 5% w/w Gel	not available	PA 979/32/3	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN 400, 400 mg comprimés enrobés	not available	BE189926	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400, 400 mg überzogene Tabletten	not available	BE189926	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 200 Soditabs 200 mg omhulde tabletten	not available	BE415283	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 200 Soditabs 200 mg comprimés enrobés	not available	BE415283	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 200 Soditabs 200 mg überzogene Tabletten	not available	BE415283	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
Нурофен Х2 256 mg обвити таблетки	not available	20100647	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Express 200mg, kaetud tabletid	not available	743211	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/14	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/17	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/20	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/23	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/26	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/29	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/32	RECKITT BENCKISER KFT	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
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/35	RECKITT BENCKISER KFT	HU
Nurofen Rapid bevont tabletta	not available	OGYI-T-6793/38	RECKITT BENCKISER KFT	HU
Nurofen Express 200mg Tablets	not available	PA 979/32/10	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Express 200mg apvalkotās tabletes	not available	09-0056	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Long Lasting 300mg Prolonged Release Hard Capsules	not available	PA 979/32/15	RECKITT BENCKISER IRELAND LTD.	IE
Нурофен Форте 400 mg обвити таблетки	not available	9800354	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Forte, 400 mg kaetud tabletid	not available	450704	RECKITT BENCKISER (POLAND) S.A.	EE
NUREFLEX 400 mg, comprimé enrobé	not available	364 935-6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	368 646-9	RECKITT BENCKISER HEALTHCARE 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
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/93	RECKITT BENCKISER KFT	HU
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/94	RECKITT BENCKISER KFT	HU
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/95	RECKITT BENCKISER KFT	HU
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/96	RECKITT BENCKISER KFT	HU
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/97	RECKITT BENCKISER KFT	HU
Nurofen Forte 400 mg bevont tabletta	not available	OGYI-T-6793/98	RECKITT BENCKISER KFT	HU
NUROFEN 400 mg compresse rivestite	not available	025634128	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN 400 mg compresse rivestite	not available	025634130	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen Forte 400 mg apvalkotās tabletes	not available	03-0422	RECKITT BENCKISER (POLAND) S.A.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN FORTE 400 mg dengtos tabletės	not available	LT/1/97/0271/001	RECKITT BENCKISER (POLAND) S.A.	LT
NUROFEN FORTE 400 mg dengtos tabletės	not available	LT/1/97/0271/002	RECKITT BENCKISER (POLAND) S.A.	LT
NUROFEN RAPID 400 mg měkké tobolky	not available	29/158/08-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen 400 Orange bruisgranulaat, bruisgranulaat 400 mg per sachet	not available	RVG 19423	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Gel 50 mg/g gel	not available	4977989	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Gel 50 mg/g gel	not available	4978086	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Extra Strength 400 mg Liquid Capsules	not available	PL 00063/0653	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Express 400 mg Liquid Capsules	not available	PL 00063/0653	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Advance 200mg Tablets	not available	PA 979/75/1	RECKITT BENCKISER IRELAND LTD.	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
Nurofen Flextin 200 mg tablet, omhulde tabletten	not available	RVG 100658	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Express, 200 mg, tabletki powlekane	not available	14446	RECKITT BENCKISER (POLAND) S.A.	PL
NUROFEN XPRESS 200 mg comprimidos revestidos	not available	5296579	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN XPRESS 200 mg comprimidos revestidos	not available	5296603	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN XPRESS 200 mg comprimidos revestidos	not available	5296629	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Expres 200 mg Obalené tablety	not available	07/0062/09-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen Pain Relief 256mg Tablets	not available	00063/0373	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Express 256 mg Tablets	not available	00063/0372	RECKITT BENCKISER (UK) LIMITED	UK
NUROFEN FORTE EXPRESS 400 mg dengtos tabletės	not available	LT/1/97/0271/023	RECKITT BENCKISER (POLAND) S.A.	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
Nurofen 200, 200 mg comprimés enrobés	not available	0209/11041123	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN POUR ENFANTS SANS SUCRE 2% suspension buvable	not available	0209/10040751	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400 Fastcaps, 400 mg capsules molles	not available	0209/11041125	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400, 400 mg comprimés enrobés	not available	0209/11041124	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN LYSINE 200 mg, comprimés pelliculés	not available	2010080859	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400 FASTTABS 400 mg, comprimés pelliculés	not available	0209/03070141	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen 200 mg Tablets	not available	MA096/01103	RECKITT BENCKISER HEALTHCARE (UK) LTD	MT
Nurofen for Children Sachets 100mg/5ml Oral Suspension	not available	PA 979/32/8	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN EXPRESS 200 mg dengtos tabletės	not available	LT/1/97/0271/022	RECKITT BENCKISER (POLAND) S.A.	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
Nurofen Back Pain 300mg Sustained Release Capsules Nurofen Long Lasting Pain Relief 300mg Prolonged Release Capsules	not available	PL 00063/0378	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen, 50 mg/g, żel	not available	10062	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Xpress 200 mg C ₸₀psulas moles	not available	5734207	RECKITT BENCKISER HEALTHCARE, LDA.	PT
NUROFEN VOOR KINDEREN SUIKERVRIJ rood 2% suspensie voor oraal gebruik.	not available	BE281644	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR ENFANTS SANS SUCRE rouge 2% suspension buvable	not available	BE281644	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder zuckerfrei Rot 2% Suspension zum Einnehmen	not available	BE281644	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Нурофен за деца с вкус на Ягода 100mg/5ml перорална суспензия	not available	20060193	RECKITT BENCKISER (ROMANIA) SRL	BG
Nurofen Junior Fiebersaft Erdbeer 20 mg/ml Suspension zum	not available	62150.00.00	RECKITT BENCKISER DEUTSCHLAND 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
Einnehmen				
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/02	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen for Children Strawberry 100mg/5ml Oral Suspension	not available	PA 979/32/9	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN FEBBRE E DOLORE Bambini 100mg/5ml sospensione orale gusto fragola senza | zucchero	not available	034102259	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE Bambini 100mg/5ml sospensione orale gusto fragola senza | zucchero	not available	034102261	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE Bambini 100mg/5ml sospensione orale gusto fragola senza | zucchero	not available	034102246	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E DOLORE Bambini 100mg/5ml sospensione orale gusto fragola senza | zucchero	not available	034102273	RECKITT BENCKISER HEALTHCARE (ITALIA) 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
Nurofen 200 tablet ovaal, omhulde tabletten 200 mg	not available	RVG 25190	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen 200 mg Caplets	not available	PL 00063/0386	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen 200 mg Caplets	not available	PL 00063/0387	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen for Children 3 months to 9 years Strawberry	not available	PL 00063/0667	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Cold, Pain and Fever Strawberry Flavour 100mg/5mL Oral Suspension	not available	PL 00063/0667	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Strawberry 3 Months to 12 years	not available	PL 00063/0666	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Strawberry	not available	PL 00063/0666	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen for Children Strawberry Singles	not available	PL 00063/0670	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
NUROFEN POUR ENFANTS 200 mg comprimés enrobés.	not available	0209/10040753	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	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
NUROFEN POUR ENFANTS SANS SUCRE rouge 2% suspension buvable	not available	0209/10040752	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/04-07	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen 125 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/08-11	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen pentru copii cu aromă de portocale 100 mg/5 ml suspensie orală	not available	9006/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN EXPRESS 200 mg dengtos tabletės	not available	LT/1/97/0271/021	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen narancsízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/01	RECKITT BENCKISER KFT	HU
Nurofen 200 mg Coated Tablets	not available	9734	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY
Nurofen Max Strength Pain Relief 512mg tablets	not available	00063/0412	RECKITT BENCKISER (HEALTHCARE) UK LTD	UK
Nurofen Xpress 200 mg Cápsulas moles	not available	5732441	RECKITT BENCKISER HEALTHCARE, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen For Children	not available	MA0190/00402	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	MT
Nurofen 200mg Liquid Capsules	not available	PA 979/32/12	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Rapid 200 mg meke kapsule	not available	HR-H-821340280	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 200 mg obložene tablete	not available	HR-H-151406116	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen za djecu 100mg/5 ml oralna suspenzija	not available	HR-H-846940596	RECKITT BENCKISER (CROATIA) D.O.O.	HR
LASONIL ANTIDOLORE 10% gel	not available	042154029	BAYER SPA	IT
LASONIL ANTIDOLORE 10% gel	not available	042154017	BAYER SPA	IT
ARFEN 10 mg/ml Soluzione vaginale	not available	024635094	LAB. IT. BIOCHIM. FARM.CO LISAPHARMA S.P.A.	IT
Ibufarmalid, 200 mg/ 10 ml, zawiesina doustna	not available	21882	FARMALIDER, S.A.	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
Ibutop ® Gel	not available	PL 41188/0031	JENSONR+ LTD.	UK
IBALGIN RAPID 400 mg filmom obložene tablete	not available	HR-H-029787657-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Dolgit® Schmerzgel	not available	12779.00.00	DOLORGIET GMBH & CO KG	DE
Dolgit® Schmerzgel, 50 mg/g Gel	not available	473/96/07/0422	DOLORGIET GMBH & CO KG	LU
Moment 200 mg granulato per soluzione orale	not available	025669211	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Ibuprofen Sanias 400 mg, filmomhulde tabletten	NL/H/4164/002	RVG 101775	AUROBINDO PHARMA B.V.	NL
Ibuprofen Sanias 200 mg, filmomhulde tabletten	NL/H/4164/001	RVG 101773	AUROBINDO PHARMA B.V.	NL
Íbúfen 800mg Film-coated tablet	IS/H/0425/004	IS/1/09/068/04	ACTAVIS GROUP PTC EHF.	IS
Ibuprofen Actavis 800mg Film-coated tablet	IS/H/0425/004	MA628/07104	ACTAVIS GROUP PTC EHF.	MT