12 October 2023
EMA/485899/2023
Human Medicines Division

List of nationally authorised medicinal products

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

Procedure No. PSUSA/00010649/202302

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
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/002	VIATRIS HEALTHCARE LTD	LT
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/003	VIATRIS HEALTHCARE LTD	LT
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/004	VIATRIS HEALTHCARE LTD	LT
Abfen 40 mg/ml geriamoji suspensija	PT/H/2354/001	LT/1/14/3546/001	VIATRIS HEALTHCARE LTD	LT
Actiprofen 200 mg tablets.	not available	PA 0678/061/002	GLAXOSMITHKLINE CONSUMER HEALTHCARE (IRELAND) LTD	IE
ADOLORIN Ibuforte DIREKT 400 mg Suspension zum Einnehmen	SI/H/0214/003	137946	KWIZDA PHARMA GMBH	AT
ADOLORIN® Ibuforte EXPRESS 400 mg Filmtabletten	not available	141241	KWIZDA PHARMA GMBH	AT
ADOLORINI DIREKT 200 mg Suspension zum Einnehmen	SI/H/0214/002	137945	KWIZDA PHARMA GMBH	AT
Advil 200 mg drajeuri	not available	13301/2020/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil 200 mg drajeuri	not available	13301/2020/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil 200 mg drajeuri	not available	13301/2020/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil 200 mg drajeuri	not available	13301/2020/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil 200 mg drajeuri	not available	13301/2020/05	GLAXOSMITHKLINE CONSUMER HEALTHCARE	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
			S.R.L.	
ADVIL 200 mg, comprimé enrobé	not available	34009 332 315 2 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVIL 200 mg, comprimé enrobé	not available	34009 329 593 5 8	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVIL 400 mg, comprimé enrobé	not available	34009 381 710 9 9	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVIL 400 mg, comprimé enrobé	not available	34009 381 711 5 0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
Advil Ultra 200 mg capsule moi	not available	8319/2015/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra 200 mg capsule moi	not available	8319/2015/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/09	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/11	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/07	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/10	GLAXOSMITHKLINE CONSUMER HEALTHCARE	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
			S.R.L.	
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/12	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/08	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/06	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/05	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/13	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte 400 mg capsule moi	not available	13258/2020/14	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/22	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/23	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy	not available	OGYI-T-8476/08	GLAXOSMITHKLINE-	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
kapszula			CONSUMER KFT.	
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/06	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/05	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/24	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/20	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/21	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/25	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/26	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/27	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/28	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/29	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Ultra Forte lágy kapszula	not available	OGYI-T-8476/30	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil, dragees 200 mg	not available	RVG 12471	HALEON NETHERLANDS B.V.	NL
Advil®	not available	2452801	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 627 4 5	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 626 8 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 630 5 6	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-	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
			MALMAISON	
ADVILCAPS 200 mg, capsule molle	not available	34009 381 632 8 5	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 628 0 6	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 631 1 7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 200 mg, capsule molle	not available	34009 381 629 7 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 429-3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 431-8	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 430-1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 426-4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 428-7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 427-0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 432-4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	364 433-0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-	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
			MALMAISON	
ADVILCAPS 400 mg, capsule molle	not available	34009 382 865 6 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILCAPS 400 mg, capsule molle	not available	34009 382 866 2 5	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 100 mg, comprimé enrobé	not available	34009 358 459 1 7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 400 mg, comprimé enrobé	not available	34009 329 595 8 7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 400 mg, comprimé enrobé	not available	34009 329 594 1 9	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 5 %, gel	not available	NL26805	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 5 %, gel	not available	NL26805	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED 5 %, gel	not available	NL26805	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
ADVILMED ENFANTS ET NOURRISSONS 20 mg/1 ml, suspension buvable en flacon	not available	34009 336 406 2 0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
Aktren® 200 mg überzogene Tabletten Ibuprofen	not available	7554.00.00	BAYER VITAL GMBH	DE
Aktren® Forte 400 mg Filmtabletten Ibuprofen	not available	13407.01.00	BAYER VITAL GMBH	DE
Aktren® Spezial 400 mg	not available	37995.00.00	BAYER VITAL GMBH	DE

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Weichkapseln Ibuprofen				
Algifast 400 mg polvo para suspensión oral	not available	75357	LABORATORIO DE APLICACIONES FARMACODINÁMICAS, S.A.-FARDI	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
Algoflex Duo 400 mg/100 mg filmtabletta	DE/H/5544/001/DC	OGYI-T-23652/01	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
Algoflex Duo 400 mg/100 mg filmtabletta	DE/H/5544/001/DC	OGYI-T-23652/02	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
Algoflex Duo 400 mg/100 mg filmtabletta	DE/H/5544/001/DC	OGYI-T-23652/04	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
Algoflex Duo 400 mg/100 mg filmtabletta	DE/H/5544/001	OGYI-T-23652/03	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
Algoflex Gyorsan Ható 400 mg filmtabletta	CZ/H/0184/001	OGYI-T-21112/02	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
Algoflex Gyorsan Ható 400 mg filmtabletta	CZ/H/0184/001	OGYI-T-21112/01	OPELLA HEALTHCARE COMMERCIAL KFT.	HU
ALGOFRENELLE® 1% w/v Κολπικό διάλυμα	not available	121383/03.12.2021	IOULIA AND IRENE TSETI PHARMACEUTICAL LABORATORIES S.A.	GR
ALGOPIRINA FEBBRE E DOLORE 200 mg sospensione orale in bustina, gusto arancia	not available	042178032	SO.SE.PHARM S.R.L.	IT
ALGOPIRINA FEBBRE E DOLORE 200 mg sospensione orale in bustina, gusto arancia	not available	042178044	SO.SE.PHARM S.R.L.	IT
Alvofen Express 400 mg mjúk hylki	IS/H/0205/001	IS/1/13/050/02	ALVOGEN EHF	IS
ANTALGIL 200 mg	not available	027432020	S&R FARMACEUTICI	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
compresse				
ANTALGIL 200 mg compresse	not available	027432032	S&R FARMACEUTICI	IT
ANTARENE 100 mg, comprimé pelliculé	not available	34009 351 999 0 4	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
ANTARENE 100 mg, comprimé pelliculé	not available	34009 351 998 4 3	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
ANTARENE 5 %, gel	not available	34009 355 627 0 8	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
ANTARENE 5 %, gel	not available	34009 364 380 4 0	LABORATOIRE DES REALISATIONS THERAPEUTIQUES ELERTE	FR
ARFEN 10 mg/ml Soluzione vaginale	not available	024635094	S.F. GROUP SRL	IT
ARFEN 400 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	024635106	S.F. GROUP SRL	IT
ARFEN 500 mg Compresse	not available	024635029	S.F. GROUP SRL	IT
Asda Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
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
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	XI
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Boots Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI

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
Boots Ibuprofen 3 Months Plus 100mg/5ml Oral Suspension Strawberry Flavour	not available	PL 00014/0652	THE BOOTS COMPANY PLC	XI
Boots Ibuprofen 3 Months Plus 100mg/5ml Oral Suspension Strawberry Flavour	not available	PL 00014/0652	THE BOOTS COMPANY PLC	XI
Boots Ibuprofen 3 Months Plus 100mg/5ml Oral Suspension Strawberry Flavour	not available	PL 00014/0859	THE BOOTS COMPANY PLC	XI
Boots Ibuprofen 3 Months Plus 100mg/5ml Suspension Orange Flavour	not available	PL 00014/0648	THE BOOTS COMPANY PLC	XI
Boots Ibuprofen Long Lasting Capsules 200mg	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Boots Max Strength Ibuprofen 10% Gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	XI
Brufedol 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0912/001	023725	VIATRIS HEALTHCARE LTD	CY
Brufedol 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0912/001	93572/12-10-2021	VIATRIS HEALTHCARE LTD	GR
Brufedol 40 mg/ml suspensija iekšķīgai lietošanai	PT/H/2354/001	14-0134	VIATRIS HEALTHCARE LTD	LV
Brufedol 400 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0912/002	023726	VIATRIS HEALTHCARE LTD	CY
Brufedol 400 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0912/002	93573/12-10-2021	VIATRIS HEALTHCARE LTD	GR
Brufedol 600 mg	SE/H/0912/003	023727	VIATRIS HEALTHCARE LTD	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
επικαλυμμένα με λεπτό υμένιο δισκία				
Brufedol 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0912/003	93574/12-10-2021	VIATRIS HEALTHCARE LTD	GR
Brufedol 800 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0912/005	023728	VIATRIS HEALTHCARE LTD	CY
Brufedol 800 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0912/005	93575/12-10-2021	VIATRIS HEALTHCARE LTD	GR
Brufedol Rapid 400 mg filmom obalené tablety	SK/H/0257/002	29/0364/13-S	VIATRIS LIMITED	SK
Brufen	not available	48827/10-09-2009	VIATRIS HEALTHCARE LTD	GR
Brufen	not available	48832/10-09-2009	VIATRIS HEALTHCARE LTD	GR
Brufen	not available	48830/10-09-2009	VIATRIS HEALTHCARE LTD	GR
Brufen 20 mg/ml oral suspension	SE/H/0912/004	9773	VIATRIS AB	SE
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/016	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/018	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/019	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/020	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/017	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/021	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/022	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml peroralna suspenzija	SE/H/0912/004	H/10/01998/023	VIATRIS HEALTHCARE LTD	SI
Brufen 20 mg/ml sirup	not available	29/916/92-S/C	VIATRIS HEALTHCARE LTD	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
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
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/01	VIATRIS HEALTHCARE LTD	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/02	VIATRIS HEALTHCARE LTD	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/04	VIATRIS HEALTHCARE LTD	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/06	VIATRIS HEALTHCARE LTD	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/03	VIATRIS HEALTHCARE LTD	RO
Brufen 20 mg/ml suspensie orală	SE/H/0912/004	7459/2015/05	VIATRIS HEALTHCARE LTD	RO
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
Brufen 200 mg filmdragerade tabletter	SE/H/0912/001	9034	VIATRIS AB	SE
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/01	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/02	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/03	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/04	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/05	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/06	MYLAN EPD 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
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/07	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/08	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/09	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/10	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/11	MYLAN EPD KFT.	HU
Brufen 200 mg filmtabletta	HU/H/0627/001	OGYI-T-23880/12	MYLAN EPD KFT.	HU
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 40 mg/ml suspensie voor oraal gebruik	PT/H/2354/001	BE441847	VIATRIS HEALTHCARE	BE
Brufen 40 mg/ml suspension buvable	PT/H/2354/001	BE441847	VIATRIS HEALTHCARE	BE
Brufen 40 mg/ml suspension buvable	PT/H/2354/001	2014040014	VIATRIS HEALTHCARE	LU
Brufen 40 mg/ml Suspension zum Einnehmen	PT/H/2354/001	BE441847	VIATRIS HEALTHCARE	BE
Brufen 40 mg/ml Suspension zum Einnehmen	PT/H/2354/001	2014040014	VIATRIS HEALTHCARE	LU
Brufen 400 mg brusgranulat	SE/H/1184/001	46807	VIATRIS AB	SE
BRUFEN 400 mg	not available	022593204	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
Compresse rivestite				
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/01	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/02	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/03	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/04	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/05	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/06	VIATRIS HEALTHCARE LTD	RO
Brufen 400 mg comprimate filmate	SE/H/0912/002	7460/2015/07	VIATRIS HEALTHCARE LTD	RO
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 400 mg film-coated tablets	SE/H/0912/002	MA1507/02901	VIATRIS HEALTHCARE LTD	MT
Brufen 400 mg filmdragerade tabletter	SE/H/0912/002	9103	VIATRIS AB	SE
Brufen 400 mg filmom obložene tablete	not available	HR-H-290369954	VIATRIS HRVATSKA D.O.O.	HR
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/004	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/005	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/006	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/007	VIATRIS HEALTHCARE LTD	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/10/01998/008	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/009	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmsko obložene tablete	SE/H/0912/002	H/10/01998/010	VIATRIS HEALTHCARE LTD	SI
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/13	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/14	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/15	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/16	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/17	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/18	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/19	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/20	MYLAN EPD KFT.	HU
Brufen 400 mg filmtabletta	HU/H/0627/002	OGYI-T-23880/21	MYLAN EPD KFT.	HU
BRUFEN 400 mg Filmtabletten	not available	BE104876	VIATRIS HEALTHCARE	BE
BRUFEN 400 mg Filmtabletten	not available	BE467093	VIATRIS HEALTHCARE	BE
BRUFEN 400 mg Filmtabletten	not available	2002016342	VIATRIS HEALTHCARE	LU
Brufen 400 mg granulado efervescente	SE/H/1184/001	5571567	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 400 mg pezsgőgranulátum	SE/H/1184/001	OGYI-T-22073/05	VIATRIS HEALTHCARE LTD	HU
Brufen 400 mg	not available	29/390/92/-S/C	VIATRIS HEALTHCARE LTD	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
potahované tablety				
Brufen 400 mg šumeće granule	not available	HR-H-496875141	VIATRIS HRVATSKA D.O.O.	HR
BRUFEN 400 mg filmom obalené tablety	not available	29/0110/06-S	VIATRIS HEALTHCARE LTD	SK
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 079 5 0	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 257 0 1	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 259 3 0	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 258 7 9	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 260 1 2	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimé pelliculé	not available	34009 362 080 3 2	VIATRIS MEDICAL	FR
BRUFEN 400 mg, comprimés pelliculés	not available	BE467093	VIATRIS HEALTHCARE	BE
BRUFEN 400 mg, comprimés pelliculés	not available	BE104876	VIATRIS HEALTHCARE	BE
BRUFEN 400 mg, comprimés pelliculés	not available	2002016342	VIATRIS HEALTHCARE	LU
BRUFEN 400 mg, filmomhulde tabletten	not available	BE467093	VIATRIS HEALTHCARE	BE
BRUFEN 400 mg, filmomhulde tabletten	not available	BE104876	VIATRIS HEALTHCARE	BE
Brufen 400mg film-coated tablets	not available	PA 2010/2/1	MYLAN IRE HEALTHCARE LIMITED	IE
BRUFEN 600 mg Compresse rivestite	not available	022593216	MYLAN ITALIA S.R.L.	IT
Brufen 600 mg comprimidos revestidos por película	not available	8660811	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen 600 mg	not available	5550389	BGP PRODUCTS	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
comprimidos revestidos por película			UNIPESSOAL, LDA.	
Brufen 600 mg filmdragerade tabletter	SE/H/0912/003	9774	VIATRIS AB	SE
Brufen 600 mg filmom obložene tablete	not available	HR-H-794772028	VIATRIS HRVATSKA D.O.O.	HR
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/011	VIATRIS HEALTHCARE LTD	SI
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/012	VIATRIS HEALTHCARE LTD	SI
Brufen 600 mg filmsko obložene tablete	SE/H/0912/003	H/10/01998/013	VIATRIS HEALTHCARE LTD	SI
Brufen 600 mg granulado efervescente	not available	5550488	BGP PRODUCTS UNIPESSOAL, LDA.	PT
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
Brufen 600 mg šumeće granule	not available	HR-H-548019973	VIATRIS HRVATSKA D.O.O.	HR
Brufen 600 mg šumivé granule	not available	29/655/08-C	VIATRIS HEALTHCARE LTD	CZ
BRUFEN 600 mg šumivý granulát	not available	29/0502/08-S	VIATRIS HEALTHCARE LTD	SK
Brufen 600mg film-coated tablets	not available	PA 2010/2/2	MYLAN IRE HEALTHCARE LIMITED	IE
BRUFEN 800 mg compresse rivestite a rilascio prolungato	not available	022593115	MYLAN ITALIA S.R.L.	IT
Brufen 800 mg ilgstošās darbības tabletes	SE/H/0912/005	11-0251	VIATRIS HEALTHCARE LTD	LV
Brufen 800 mg pailginto atpalaidavimo tabletės	SE/H/0912/005	LT/1/10/1885/008	VIATRIS HEALTHCARE LTD	LT
Brufen 800 mg pailginto atpalaidavimo tabletės	SE/H/0912/005	LT/1/10/1885/007	VIATRIS HEALTHCARE LTD	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
Brufen 800 mg retard tabletta	SE/H/0912/005	OGYI-T-22073/02	VIATRIS HEALTHCARE LTD	HU
Brufen 800 mg retard tabletta	SE/H/0912/005	OGYI-T-22073/01	VIATRIS HEALTHCARE LTD	HU
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
BRUFEN ANALGESICO 200 mg compresse rivestite con film	NL/H/2550/001	042386110	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 200 mg compresse	NL/H/2550/001	042386108	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
rivestite con film				
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
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	NL/H/2550/001	042386223	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
rivestite con film				
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
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 200 mg compresse rivestite con film	NL/H/2550/001	042386591	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 200 mg compresse rivestite con film	NL/H/2550/001	042386603	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386312	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386300	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386324	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse	NL/H/2550/002	042386336	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
rivestite con film				
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386348	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386363	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386351	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386375	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386387	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386399	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386425	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386401	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386413	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386449	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386437	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse	NL/H/2550/002	042386452	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
rivestite con film				
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386464	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386488	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386476	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386502	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386490	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386526	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386514	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386538	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386540	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386553	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386565	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse	NL/H/2550/002	042386577	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
rivestite con film				
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386589	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386627	MYLAN S.P.A.	IT
BRUFEN ANALGESICO 400 mg compresse rivestite con film	NL/H/2550/002	042386615	MYLAN S.P.A.	IT
Brufen Bruis 600 mg, bruisgranulaat	not available	RVG 13331	MYLAN HEALTHCARE B.V.	NL
Brufen cukormentes 40 mg/ml belsőleges szuszpenzió	PT/H/2354/001	OGYI-T-22073/03	VIATRIS HEALTHCARE LTD	HU
Brufen cukormentes 40 mg/ml belsőleges szuszpenzió	PT/H/2354/001	OGYI-T-22073/04	VIATRIS HEALTHCARE LTD	HU
Brufen Effect 20 mg/ml sirup	not available	HR-H-254292502	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect 200 mg šumeće granule	not available	HR-H-053171042	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect 200 mg šumeće granule	not available	HR-H-765431689	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect 400 mg filmom obložene tablete	not available	HR-H-334193871	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect 400 mg šumeće granule	not available	HR-H-071108593	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect forte 40 mg/ml oralna suspenzija	not available	HR-H-330150785	VIATRIS HRVATSKA D.O.O.	HR
Brufen Effect Rapid 684 mg filmom obložene tablete	HU/H/0627/002	HR-H-922117795	VIATRIS LIMITED	HR
BRUFEN FORTE 600 mg, comprimés pelliculés	not available	BE128064	VIATRIS HEALTHCARE	BE
BRUFEN FORTE 600 mg,	not available	2002016343	VIATRIS HEALTHCARE	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
comprimés pelliculés				
BRUFEN FORTE 600 mg, filmomhulde tabletten	not available	BE128064	VIATRIS HEALTHCARE	BE
BRUFEN FORTE 600 mg, Filmtabletten	not available	BE128064	VIATRIS HEALTHCARE	BE
BRUFEN FORTE 600 mg, Filmtabletten	not available	2002016343	VIATRIS HEALTHCARE	LU
Brufen Granules	not available	PL 46302/0007	MYLAN PRODUCTS LIMITED	XI
Brufen Granules 400 mg	SE/H/1184/001	2013110373	VIATRIS HEALTHCARE	LU
Brufen Granules 400 mg, Brausegranulat	SE/H/1184/001	BE434411	VIATRIS HEALTHCARE	BE
Brufen Granules 400 mg, Brausegranulat	SE/H/1184/001	2013110373	VIATRIS HEALTHCARE	LU
Brufen Granules 400 mg, bruisgranulaat	SE/H/1184/001	BE434411	VIATRIS HEALTHCARE	BE
Brufen Granules 400mg, granulés effervescents	SE/H/1184/001	BE434411	VIATRIS HEALTHCARE	BE
BRUFEN GRANULES 600 mg, Brausegranulat	not available	BE158067	VIATRIS HEALTHCARE	BE
BRUFEN GRANULES 600 mg, Brausegranulat	not available	2002016339	VIATRIS HEALTHCARE	LU
BRUFEN GRANULES 600 mg, bruisgranulaat	not available	BE158067	VIATRIS HEALTHCARE	BE
BRUFEN GRANULES 600 mg, granulés effervescents	not available	BE158067	VIATRIS HEALTHCARE	BE
BRUFEN GRANULES 600 mg, granulés effervescents	not available	2002016339	VIATRIS HEALTHCARE	LU
BRUFEN INSTANT 400 mg šumivý granulát	SE/H/1184/001	29/0162/13-S	VIATRIS HEALTHCARE LTD	SK
Brufen Paediatric 100mg/5ml oral suspension	not available	PA 2010/2/6	MYLAN IRE HEALTHCARE LIMITED	IE
BRUFEN RAPID 400 mg	SK/H/0257/002	29/407/13-C	MYLAN IRELAND LIMITED	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
potahované tablety				
Brufen Retard	not available	PL 46302/0010	MYLAN PRODUCTS LIMITED	XI
Brufen Retard 800 mg comprimate cu eliberare prelungită	SE/H/0912/005	7461/2015/01	VIATRIS HEALTHCARE LTD	RO
Brufen Retard 800 mg comprimate cu eliberare prelungită	SE/H/0912/005	7461/2015/02	VIATRIS HEALTHCARE LTD	RO
Brufen Retard 800 mg comprimidos de libertação prolongada	not available	4338281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Retard 800 mg comprimidos de libertação prolongada	not available	4338281	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Retard 800 mg depottabletter	not available	7614	VIATRIS AS	NO
Brufen Retard 800 mg depottabletter	SE/H/0912/005	11238	VIATRIS AB	SE
Brufen Retard 800 mg prolonged-release tablets	SE/H/0912/005	MA1507/02902	VIATRIS HEALTHCARE GMBH	MT
Brufen retard 800 mg tablete s podaljšanim sproščanjem	SE/H/0912/005	H/10/01998/014	VIATRIS HEALTHCARE LTD	SI
Brufen retard 800 mg tablete s podaljšanim sproščanjem	SE/H/0912/005	H/10/01998/015	VIATRIS HEALTHCARE LTD	SI
BRUFEN RETARD 800 mg filmom obalené tablety s predĺženým uvoľňovaním	not available	29/0659/10-S	VIATRIS HEALTHCARE LTD	SK
BRUFEN RETARD 800 mg, comprimés à libération prolongée	not available	BE150376	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800 mg, comprimés à libération prolongée	not available	BE147436	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800	not available	2002016344	VIATRIS HEALTHCARE	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
mg, comprimés à libération prolongée				
Brufen Retard 800 mg, tablet met gereguleerde afgifte	not available	RVG 13478	MYLAN HEALTHCARE B.V.	NL
BRUFEN RETARD 800 mg, tabletten met verlengde afgifte	not available	BE150376	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800 mg, tabletten met verlengde afgifte	not available	BE147436	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	BE147436	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	BE150376	VIATRIS HEALTHCARE	BE
BRUFEN RETARD 800 mg, Tabletten mit verzögerter Wirkstofffreisetzung	not available	2002016344	VIATRIS HEALTHCARE	LU
Brufen Retard 800mg prolonged release tablets	not available	PA 2010/2/7	MYLAN IRE HEALTHCARE LIMITED	IE
Brufen Retard, depottabletter	not available	13416	VIATRIS APS	DK
Brufen Sem Açúcar 20 mg/ml suspensão oral	not available	5746342	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Sem Açúcar 20 mg/ml suspensão oral	not available	5746359	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Sem Açúcar 40 mg/ml suspensão oral	PT/H/2354/001	5475819	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Sem Açúcar 40 mg/ml suspensão oral	PT/H/2354/001	5442132	BGP PRODUCTS UNIPESSOAL, LDA.	PT
BRUFEN sirup	not available	29/0916/92-S	VIATRIS HEALTHCARE LTD	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
100 mg/5 ml sirup				
Brufen SR 800 mg tablete s produljenim oslobađanjem	not available	HR-H-532068337	VIATRIS HRVATSKA D.O.O.	HR
Brufen Suspensão 20 mg/ml suspensão oral	not available	8523209	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufen Syrup	not available	PL 46302/0011	MYLAN PRODUCTS LIMITED	XI
Brufen Tablets 400 mg	not available	PL 46302/0006	MYLAN PRODUCTS LIMITED	XI
Brufen Tablets 600 mg	not available	PL 46302/0009	MYLAN PRODUCTS LIMITED	XI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/001	VIATRIS HEALTHCARE LTD	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/002	VIATRIS HEALTHCARE LTD	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/003	VIATRIS HEALTHCARE LTD	SI
Brufen z okusom jagode 40 mg/ml peroralna suspenzija	PT/H/2354/001	H/12/02001/004	VIATRIS HEALTHCARE LTD	SI
Brufen, 200 mg kihisevad graanulid	EE/H/0295/001	809113	VIATRIS HEALTHCARE LTD	EE
Brufen, 40 mg/ml suukaudne suspensioon	PT/H/2354/001	820713	VIATRIS HEALTHCARE LTD	EE
Brufen, 800 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0912/005	750811	VIATRIS HEALTHCARE LTD	EE
Brufen, filmovertrukne tabletter 400 mg	not available	06689	VIATRIS APS	DK
Brufen, filmovertrukne tabletter 600 mg	not available	10882	VIATRIS APS	DK
Brufen® 400 mg - Filmtabletten	not available	16233	MYLAN ÖSTERREICH GMBH	AT
Brufen® 600 mg -	not available	1-18151	MYLAN ÖSTERREICH GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Filmtabletten				
BRUFEN® Επικαλυμμένο δισκίο 400 mg	not available	5754	VIATRIS HEALTHCARE LTD	CY
Brufenact 200 mg comprimidos revestidos por película	HU/H/0627/001	5810734	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufenact 200 mg comprimidos revestidos por película	HU/H/0627/001	5810742	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufenact 200 mg comprimidos revestidos por película	HU/H/0627/001	5810759	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufenact 400 mg comprimidos revestidos por película	HU/H/0627/002	5810767	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufenact 400 mg comprimidos revestidos por película	HU/H/0627/002	5810775	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brufenact 400 mg comprimidos revestidos por película	HU/H/0627/002	5810809	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Brumare 400 mg putojošās granulas	SE/H/1184/001	13-0078	VIATRIS HEALTHCARE LTD	LV
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/002	VIATRIS HEALTHCARE LTD	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/003	VIATRIS HEALTHCARE LTD	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/001	VIATRIS HEALTHCARE LTD	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/005	VIATRIS HEALTHCARE LTD	LT
Brumare 400 mg šnypščiosios granulės	SE/H/1184/001	LT/1/13/3280/004	VIATRIS HEALTHCARE LTD	LT
Brumare, 400 mg kihisevad graanulid	SE/H/1184/001	805213	VIATRIS HEALTHCARE LTD	EE
Brupro 200mg Film-	not available	PA0074/067/001	ROWA PHARMACEUTICALS	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
coated tablets			LIMITED	
Brupro Max 400mg Film-coated tablets	not available	PA0074/067/002	ROWA PHARMACEUTICALS LIMITED	IE
Brusimp 400 mg filmtabletta	SE/H/0912/002	OGYI-T-23939/01	VIATRIS HEALTHCARE LTD	HU
Brusimp 400 mg filmtabletta	SE/H/0912/002	OGYI-T-23939/02	VIATRIS HEALTHCARE LTD	HU
Brusimp 400 mg filmtabletta	SE/H/0912/002	OGYI-T-23939/03	VIATRIS HEALTHCARE LTD	HU
Brusimp 400 mg filmtabletta	SE/H/0912/002	OGYI-T-23939/04	VIATRIS HEALTHCARE LTD	HU
Brusimp 600 mg filmtabletta	SE/H/0912/003	OGYI-T-23939/05	VIATRIS HEALTHCARE LTD	HU
Brusimp 600 mg filmtabletta	SE/H/0912/003	OGYI-T-23939/06	VIATRIS HEALTHCARE LTD	HU
Buprofess 200 mg pulbere orală	DE/H/4307/001	14551/2022/03	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	14551/2022/02	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	14551/2022/01	STADA M&D S.R.L.	RO
Buprofess 200 mg pulbere orală	DE/H/4307/001	14551/2022/04	STADA M&D S.R.L.	RO
Buprofess 400 mg pulbere orală	DE/H/4307/002	14552/2022/03	STADA M&D S.R.L.	RO
Buprofess 400 mg pulbere orală	DE/H/4307/002	14552/2022/02	STADA M&D S.R.L.	RO
Buprofess 400 mg pulbere orală	DE/H/4307/002	14552/2022/01	STADA M&D S.R.L.	RO
Burana 100 mg mjuka tuggkapslar	not available	32143	ORION CORPORATION	FI
Burana 100 mg pehmeät purukapselit	not available	32143	ORION CORPORATION	FI
Burana 40 mg/ml oraalisuspensio	IE/H/690/1	28530	ORION CORPORATION	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
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596046	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596022	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596059	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596034	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596073	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596010	OPELLA HEALTHCARE ITALY S.R.L.	IT
BUSCOACTFOKUS 400 mg + 100 mg compresse rivestite con film	DE/H/5544/001/DC	047596061	OPELLA HEALTHCARE ITALY S.R.L.	IT
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
υμένιο δισκία				
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Buscofem Extra 400 mg/100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5544/001/DC	53621 /10-06-2021	OPELLA Ε.Π.Ε.	GR
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Calprofen 100mg/5ml Ibuprofen Suspension	not available	PL 15513/0158	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Calprofen 100mg/5ml Oral Suspension Ibuprofen	not available	PL 15513/0147	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Childrens Ibuprofen 100mg/5ml Oral Suspension	not available	PL 16028/0075	GALPHARM HEALTHCARE LIMITED	XI
Cuprofen Maximum Strength Tablets	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Cuprofen Maximum Strength Tablets	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Cuprofen Tablets 400 mg	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Cuprofen Tablets 400 mg	not available	PL 00063/0756	RECKITT BENCKISER	XI

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
			HEALTHCARE (UK) LTD	
DALSY 20 mg/ml sirup	not available	HR-H-407408529	VIATRIS HRVATSKA D.O.O.	HR
Dalsy 20 mg/ml suspensión oral	not available	59166	MYLAN IRE HEALTHCARE LIMITED	ES
Dalsy 200 mg granulado efervescente	not available	63990	MYLAN IRE HEALTHCARE LIMITED	ES
Dalsy 40 mg/ml suspensión oral	not available	69726	MYLAN IRE HEALTHCARE LIMITED	ES
Dalsy forte 40 mg/ml oralna suspenzija	not available	HR-H-093886315	VIATRIS HRVATSKA D.O.O.	HR
Dalsydol 400 mg comprimidos recubiertos con película.	not available	54985	MYLAN IRE HEALTHCARE LIMITED	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
DIMIDON 400 mg, comprimidos revestidos por pelicula	not available	5564034	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por pelicula	not available	5564042	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por pelicula	not available	5564059	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
DIMIDON 400 mg, comprimidos revestidos por pelicula	not available	5564026	LABORATÓRIOS BASI – INDÚSTRIA FARMACÊUTICA, S.A.	PT
doc Ibuprofen	not available	1- 21735	HERMES ARZNEIMITTEL	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
Schmerzgel, 5% Gel			GMBH	
doc® Ibuprofen Schmerzgel, 5 % Gel	not available	39838.00.00	HERMES ARZNEIMITTEL GMBH	DE
Dolalgial ibuprofeno/cafeína 400mg /100mg comprimidos recubiertos con película	DE/H/5544/001/DC	85244	OPELLA HEALTHCARE SPAIN, S.L	ES
DOLGIT - Creme	not available	1-19151	DOLORGIET GMBH & CO KG	AT
Dolgit 50 mg/g gel	not available	29/597/95-C	DOLORGIET GMBH & CO KG	CZ
Dolgit 50 mg/g kreem	not available	147796	DOLORGIET GMBH & CO KG	EE
Dolgit akut Schmerzgel, 50 mg / g Gel	not available	1 - 21736	DOLORGIET GMBH & CO KG	AT
Dolgit gél	not available	5285/01	DOLORGIET GMBH & CO KG	HU
DOLGIT GEL 50 mg/g dermálny gél	not available	29/0851/95-S	DOLORGIET GMBH & CO KG	SK
Dolgit Gel, 50 mg/g, żel	not available	R/6758	DOLORGIET GMBH & CO KG	PL
Dolgit krém	not available	29/751/92-S/C	DOLORGIET GMBH & CO KG	CZ
Dolgit krém	not available	5284/01	DOLORGIET GMBH & CO KG	HU
DOLGIT KRÉM 50 mg/g dermálny krém	not available	29/0751/92-S	DOLORGIET GMBH & CO KG	SK
Dolgit, 50 mg /g krem	not available	6443	DOLORGIET GMBH & CO KG	PL
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
Dolobene® Ibu, 50 mg, Gel Wirkstoff: Ibuprofen	not available	33530.00.00	RECORDATI PHARMA GMBH	DE
DOLOFAST® 10% Gel Tubo 50 G	not available	029775018	DOMPÉ FARMACEUTICI S.P.A.	IT
Easofen Max Strength 400 mg Film-coated Tablets	not available	PA0126/060/002	CLONMEL HEALTHCARE 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
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
Espididol 400 mg granulado para solución oral sabor menta	not available	68344	ZAMBON, S.A.U.	ES
Espidifen 600 mg granulado para solución oral sabor albaricoque	not available	68222	ZAMBON, S.A.U.	ES
Espidifen 600 mg granulado para solución oral sabor cola-limón	not available	80719	ZAMBON, S.A.U.	ES
Essential Waitrose Long Lasting Pain Relief 200mg Modified Release Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Eudorlin Ibuprofen 20 mg/ml Suspension zum Einnehmen	DE/H/2463/001	78529.00.00	BERLIN-CHEMIE AG	DE
EUDORLIN Ibuprofen 40 mg/ml Suspension zum Einnehmen	DE/H/2598/002	79521.00.00	BERLIN-CHEMIE AG	DE
Extra Strength Ibuprofen 400mg Tablets	not available	PL 17907/0161	BRISTOL LABORATORIES LIMITED	XI
Faspic 400 mg granule pentru soluţie orală	not available	10105/2017/01	ZAMBON S.P.A.	RO
Faspic 400 mg granule pentru soluţie orală	not available	10105/2017/02	ZAMBON S.P.A.	RO
Faspic 400 mg granule pentru soluţie orală	not available	10105/2017/03	ZAMBON S.P.A.	RO
Faspic 600, granule pentru solutie orala	not available	10106/2017/02	ZAMBON S.P.A.	RO
Faspic 600, granule	not available	10106/2017/03	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
pentru solutie orala				
Faspic 600, granule pentru solutie orala	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
Faspic 600, granule pentru soluţie orală	not available	10106/2017/01	ZAMBON S.P.A.	RO
FEDOTASK 200 mg polvere orale in bustina	not available	046903011	S.F. GROUP SRL	IT
FEDOTASK 200 mg polvere orale in bustina	not available	046903023	S.F. GROUP SRL	IT
FEDOTASK 200 mg polvere orale in bustina	not available	046903050	S.F. GROUP SRL	IT
FEDOTASK 200 mg polvere orale in bustina	not available	046903062	S.F. GROUP SRL	IT
Fenbid 10 %w/w Gel	not available	PL 10972/0090	MERCURY PHARMA GROUP LTD	XI
Fenbid 5 %w/w gel	not available	PL 10972/0091	MERCURY PHARMA GROUP LTD	XI
Fenbid Forte 10% Gel	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	XI
Fenopine for Children Six Plus Strawberry 200 mg/5 ml ORAL SUSPENSION	not available	PA0281/088/006	PINEWOOD LABORATORIES LIMITED	IE
Fenpaed Ibuprofen 100 mg/5ml Oral Suspension	not available	04917/0082	PINEWOOD LABORATORIES LIMITED	XI
Flarin 200 mg soft capsules	not available	PL 51724/0001	INFIRST LTD	XI
Flarin Joint & Muscular Pain Relief 200 mg Soft Capsules	not available	PL 51724/0002	INFIRST LTD	XI
FROBEN DOLORE E FEBBRE 200 mg	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
Granulato Effervescente				
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
Galprofen Long Lasting Ibuprofen Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Geloprofen 400 mg comprimidos recubiertos con película	not available	66.327	FERRER INTERNACIONAL, S.A.	ES
Geloprofen Rapid 400 mg polvo para suspensión oral	not available	83227	FERRER INTERNACIONAL, S.A.	ES
GINENORM 0,1 g SOLUZIONE VAGINALE	not available	029135023	AESCULAPIUS FARMACEUTICI S.R.L.	IT
GINENORM 0,1 g SOLUZIONE VAGINALE	not available	029135035	AESCULAPIUS FARMACEUTICI S.R.L.	IT
GINENORM 1 g POLVERE PER SOLUZIONE VAGINALE	not available	029135011	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Ibalgin DUO 400 mg/100 mg comprimate filmate	DE/H/5544/001/DC	13115/2020/02	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin DUO 400 mg/100 mg comprimate filmate	DE/H/5544/001	13115/2020/01	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin krém 50 mg/1 g	not available	29/0051/02-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin krém 50 mg/1 g	not available	29/0051/02-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	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 Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg filmom obalené tablety	DE/H/5544/001	07/0162/20-S	OPELLA HEALTHCARE SLOVAKIA S.R.O.	SK
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Plus 400 mg/100 mg potahované tablety	DE/H/5544/001/DC	07/106/18-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/02	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/04	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/03	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg	CZ/H/0184/001	7102/2014/06	OPELLA HEALTHCARE	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
comprimate filmate			ROMANIA SRL	
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/10	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/05	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/12	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/11	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/07	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/09	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/08	OPELLA HEALTHCARE ROMANIA SRL	RO
Ibalgin Rapid 400 mg comprimate filmate	CZ/H/0184/001	7102/2014/01	OPELLA HEALTHCARE ROMANIA SRL	RO
IBALGIN RAPID 400 mg filmom obložene tablete	not available	HR-H-029787657-01	OPELLA HEALTHCARE FRANCE SAS	HR
IBALGIN RAPID 400 mg filmom obložene tablete	not available	HR-H-029787657-01	OPELLA HEALTHCARE FRANCE SAS	HR
IBALGIN RAPID 400 mg filmom obložene tablete	not available	HR-H-029787657-01	OPELLA HEALTHCARE FRANCE SAS	HR
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH 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
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
Ibalgin Rapid 400 mg potahované tablety	CZ/H/0184/001	29/581/09-C	OPELLA HEALTHCARE CZECH S.R.O	CZ
IbuARISTO 800 mg Filmtabletten	DE/H/5687/004	2202406.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ibucalm 50 mg/g gel mentolado	not available	70145	TEVA PHARMA S.L.U.,	ES
Ibucalmin 200 mg comprimate filmate	not available	13082/2020/02	LAROPHARM SRL	RO
Ibucalmin 200 mg comprimate filmate	not available	13082/2020/01	LAROPHARM SRL	RO
Ibudol 400 mg suspensión oral	not available	68194	KERN PHARMA, S.L.	ES
Ibudol 50 mg/g gel	not available	62.054	KERN PHARMA, S.L.	ES
Ibudol Pediátrico 200 mg suspensión oral	not available	68.086	KERN PHARMA, S.L.	ES
Ibudol roll-on 50 mg/g gel envase con aplicador de bola	not available	72.989	KERN PHARMA, S.L.	ES
Ibudolor Direkt 200 mg oralni prašak u vrećici	DE/H/4307/001	HR-H-054613410	STADA D.O.O.	HR
Ibudolor Direkt 400 mg oralni prašak u vrećici	DE/H/4307/002	HR-H-991661386	STADA D.O.O.	HR
Ibufarmalid 200 mg suspensión oral	not available	67.963	FARMALIDER, S.A.	ES
Ibufarmalid 400 mg	not available	68.200	FARMALIDER, S.A.	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
suspensión oral				
Ibufarmalid 600 mg suspensión oral	not available	67.931	FARMALIDER, S.A.	ES
Ibufem 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Ibufem 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Ibufem 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Íbúfen 800mg Film-coated tablet	IS/H/0425/004	IS/1/09/068/04	TEVA B.V	IS
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuflam-Lysin 400 mg Filmtabletten	CZ/H/0184/001	84271.00.00	A. NATTERMANN & CIE. 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
Ibugel	not available	PL 00173/0050	DIOMED DEVELOPMENTS LIMITED	XI
Ibugel 5% w/w Gel	not available	PA23128/011/001	DERMAL LABORATORIES (IRELAND) LIMITED	IE
Ibugel Forte 10%	not available	PL 00173/0175	DIOMED DEVELOPMENTS LIMITED	XI
Ibukern 600 mg suspensión oral	not available	67.930	KERN PHARMA, S.L.	ES
Ibuleve Max Strength Pain Relief 10% Gel	not available	PL 00173/0404	DIOMED DEVELOPMENTS LIMITED	XI
Ibuleve Maximum Strength Gel	not available	PL 00173/0176	DIOMED DEVELOPMENTS LIMITED	XI
Ibuleve Pain Relief 5% Gel/Ibuderm 5% Gel	not available	PL 00173/0060	DIOMED DEVELOPMENTS LIMITED	XI
Ibuleve Pain Relief 5% Spray	not available	PL 00173/0160	DIOMED DEVELOPMENTS LIMITED	XI
Ibuleve/Ibuleve Sports Gel	not available	PL 00173/0413	DIOMED DEVELOPMENTS LIMITED	XI
Ibu-LysinHEXAL 400 mg Filmtabletten	DE/H/2593/001	79780.00.00	HEXAL AG	DE
IBU-LYSIN-ratiopharm® 684 mg Filmtabletten 400 mg Ibuprofen	DE/H/2590/001	79779.00.00	RATIOPHARM GMBH	DE
IBU-LYSIN-ratiopharm® 684 mg Filmtabletten 400 mg Ibuprofen	DE/H/2590/001	0687/11080034	RATIOPHARM GMBH	LU
Ibumetin 50 mg/g gel	not available	02-1301	ORIFARM HEALTHCARE A/S	NO
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 "B. Braun", infusionsvæske, opløsning	ES/H/0390/001	57165	B.BRAUN MELSUNGEN AG	DK
Ibuprofen "B. Braun", infusionsvæske,	ES/H/0392/001/DC	57161	B.BRAUN MELSUNGEN AG	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
opløsning				
Ibuprofen "B. Braun", infusionsvæske, opløsning 200 mg	ES/H/0392/002/DC	62135	B.BRAUN MELSUNGEN AG	DK
Ibuprofen (als lysine) Mylan OTC 200 mg, filmomhulde tabletten	NL/H/2550/001	RVG 111362	MYLAN PHARMACEUTICALS LIMITED	NL
Ibuprofen (als lysine) Mylan OTC 400 mg, filmomhulde tabletten	NL/H/2550/002	RVG 111363	MYLAN PHARMACEUTICALS LIMITED	NL
Ibuprofen 10% w/w gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	XI
Ibuprofen 10% w/w Gel	not available	PL 10972/0090	MERCURY PHARMA GROUP LTD	XI
Ibuprofen 100mg/5ml Oral Suspension	not available	PL 29831/0695	WOCKHARDT UK LTD	XI
Ibuprofen 100mg/5ml Oral Suspension	not available	04917/0044	PINEWOOD LABORATORIES LIMITED	XI
Ibuprofen 200 mg Caplets	not available	PL 17907/0078	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200 mg film-coated Tablets	not available	PL 44041/0058	NOUMED LIFE SCIENCES	XI
Ibuprofen 200 mg Film-coated Tablets	not available	PL 44673/0084	HALEON UK TRADING LIMITED	XI
Ibuprofen 200 mg solution for infusion	ES/H/0392/002/DC	PL 03551/0155	B.BRAUN MELSUNGEN AG	XI
Ibuprofen 200 mg Tablets	not available	PL 17907/0077	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200 mg tablets	not available	PL 17907/0160	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200mg Caplets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg Caplets	not available	PL 17907/0077	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200mg Liquicaps	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI

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	XI
Ibuprofen 200mg Tablets	not available	PL 31308/0013	BRUNEL HEALTHCARE MANUFACTURING LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 31308/0014	BRUNEL HEALTHCARE MANUFACTURING LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 31308/0016	BRUNEL HEALTHCARE MANUFACTURING LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 16028/0161	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 16028/0027	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 17907/0078	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200mg tablets	not available	PL 17907/0505	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200mg tablets	not available	PL 17907/0159	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg Tablets	not available	PL 16028/0028	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 200mg/5ml Oral Suspension	UK/H/3267/002/E/001	PA0281/088/005	PINEWOOD LABORATORIES LIMITED	IE
Ibuprofen 200mg/5ml Oral Suspension	not available	04917/0099	PINEWOOD LABORATORIES LIMITED	XI
Ibuprofen 400 mg film-coated tablets	not available	PL 44041/0059	NOUMED LIFE SCIENCES	XI
Ibuprofen 400 mg Solution for Infusion	DE/H/6040/001	PL 08828/0287	FRESENIUS KABI LIMITED	XI
Ibuprofen 400 mg solution for infusion	ES/H/0390/001	PL 03551/0152	B.BRAUN MELSUNGEN AG	XI
Ibuprofen 400mg Caplets	not available	PL 16028/0040	GALPHARM HEALTHCARE	XI

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
			LIMITED	
Ibuprofen 400mg Caplets	not available	PL 16028/0040	GALPHARM HEALTHCARE LIMITED	XI
Ibuprofen 400mg Tablets	not available	PL: 31308/0035	BRUNEL HEALTHCARE MANUFACTURING LIMITED	XI
Ibuprofen 600 mg film-coated tablets	not available	PL 44041/0060	NOUMED LIFE SCIENCES	XI
Ibuprofen 600 mg solution for infusion.	ES/H/0392/001	PL 03551/0153	B.BRAUN MELSUNGEN AG	XI
Ibuprofen AL direkt 200 mg Pulver zum Einnehmen	DE/H/4307/001	94256.00.00	ALIUD PHARMA GMBH	DE
Ibuprofen AL direkt 400 mg Pulver zum Einnehmen	DE/H/4307/002	94257.00.00	ALIUD PHARMA GMBH	DE
Ibuprofen Altan, 400 mg, roztwór do infuzji	PT/H/1732/001	25888	ALTAN PHARMA LTD	PL
Ibuprofen Altan, 600 mg, roztwór do infuzji	PT/H/1732/002	25889	ALTAN PHARMA LTD	PL
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen and Caffeine Sanofi 400 mg/100 mg	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS	XI

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
film-coated tablets			SANOFI	
Ibuprofen and Caffeine Sanofi 400 mg/100 mg film-coated tablets	DE/H/5544/001/DC	PL 53886/0034	OPELLA HEALTHCARE UK LIMITED, TRADING AS SANOFI	XI
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002/DC	140180	B.BRAUN MELSUNGEN AG	AT
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002/DC	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002/DC	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002/DC	2203479.00.00	B.BRAUN MELSUNGEN AG	DE
Ibuprofen B. Braun 200 mg Infusionslösung	ES/H/0392/002	2021110224	B.BRAUN MELSUNGEN AG	LU
Ibuprofen B. Braun 200 mg infusionsvätska, lösning	ES/H/0392/002/DC	36701	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 200 mg infusionsvätska, lösning	ES/H/0392/002/DC	58662	B.BRAUN MELSUNGEN AG	SE
Ibuprofen B. Braun 200 mg infusioonilahus	ES/H/0392/002	1001020	B.BRAUN MELSUNGEN AG	EE
Ibuprofen B. Braun 200 mg infusjonsvæske, oppløsning	ES/H/0392/002/DC	18-12653	B.BRAUN MELSUNGEN AG	NO
Ibuprofen B. Braun 200 mg infuusioneste, liuos	ES/H/0392/002/DC	36701	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 200 mg infuzní roztok	ES/H/0392/002/DC	07/530/18-C	B.BRAUN MELSUNGEN AG	CZ
Ibuprofen B. Braun 200 mg infúzny roztok	ES/H/0392/002/DC	07/0041/20-S	B.BRAUN MELSUNGEN AG	SK
Ibuprofen B. Braun 200 mg oldatos infúzió	ES/H/0392/002/DC	OGYI-T-23186/03	B.BRAUN MELSUNGEN AG	HU
Ibuprofen B. Braun 200 mg oplossing voor infusie	ES/H/0392/002/DC	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200	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
mg oplossing voor infusie				
Ibuprofen B. Braun 200 mg šķīdums infūzijām	ES/H/0392/002/DC	20-0016	B.BRAUN MELSUNGEN AG	LV
Ibuprofen B. Braun 200 mg soluţie perfuzabilă	ES/H/0392/002/DC	13087/2020/01	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 200 mg soluţie perfuzabilă	ES/H/0392/002/DC	13087/2020/02	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 200 mg solution for infusion	ES/H/0392/002/DC	PA0736/041/003	B.BRAUN MELSUNGEN AG	IE
Ibuprofen B. Braun 200 mg, solution pour perfusion	ES/H/0392/002/DC	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 200 mg, solution pour perfusion	ES/H/0392/002/DC	BE557600	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 400 mg Infusionslösung	ES/H/0390/001	138239	B.BRAUN MELSUNGEN AG	AT
Ibuprofen B. Braun 400 mg Infusionslösung	ES/H/0390/001	BE512240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 400 mg Infusionslösung	ES/H/0390/001	BE512240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 400 mg Infusionslösung	ES/H/0390/001	97002.00.00	B.BRAUN MELSUNGEN AG	DE
Ibuprofen B. Braun 400 mg Infusionslösung	ES/H/0390/001	2019070164	B.BRAUN MELSUNGEN AG	LU
Ibuprofen B. Braun 400 mg infusionsvätska, lösning	ES/H/0390/001	33914	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 400 mg infusionsvätska, lösning	ES/H/0390/001	54208	B.BRAUN MELSUNGEN AG	SE
Ibuprofen B. Braun 400 mg infusjonsvæske, oppløsning	ES/H/0390/001	15-11009	B.BRAUN MELSUNGEN AG	NO
Ibuprofen B. Braun 400 mg infuusioneste, liuos	ES/H/0390/001	33914	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 400 mg infuzinis tirpalas	ES/H/0390/001	LT/1/17/4131/001	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 400 mg infuzinis tirpalas	ES/H/0390/001	LT/1/17/4131/002	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 400 mg infuzinis tirpalas	ES/H/0390/001	LT/1/17/4131/001	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 400 mg infuzinis tirpalas	ES/H/0390/001	LT/1/17/4131/002	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 400 mg infuzní roztok	ES/H/0390/001	07/866/15-C	B.BRAUN MELSUNGEN AG	CZ
Ibuprofen B. Braun 400 mg infúzny roztok	ES/H/0390/001	07/0154/17-S	B.BRAUN MELSUNGEN AG	SK
Ibuprofen B. Braun 400 mg oldatos infúzió	ES/H/0390/001	OGYI-T-23186/01	B.BRAUN MELSUNGEN AG	HU
Ibuprofen B. Braun 400 mg oplossing voor infusie	ES/H/0390/001	BE512240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 400 mg oplossing voor infusie	ES/H/0390/001	BE512240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 400 mg oplossing voor infusie	ES/H/0390/001	RVG 118611	B.BRAUN MELSUNGEN AG	NL
Ibuprofen B. Braun 400 mg raztopina za infundiranje	ES/H/0390/001	H/17/02363/001	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun 400 mg raztopina za infundiranje	ES/H/0390/001	H/17/02363/002	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun 400 mg šķīdums infūzijām	ES/H/0390/001	17-0102	B.BRAUN MELSUNGEN AG	LV
Ibuprofen B. Braun 400 mg soluție perfuzabilă	ES/H/0390/001	10023/2017/01	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 400 mg soluție perfuzabilă	ES/H/0390/001	10023/2017/02	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 400 mg solution for infusion	ES/H/0390/001	PA0736/041/001	B.BRAUN MELSUNGEN AG	IE
Ibuprofen B. Braun 400 mg solution pour	ES/H/0390/001	BE512240	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
perfusion				
Ibuprofen B. Braun 400 mg solution pour perfusion	ES/H/0390/001	BE512240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 600 mg Infusionslösung	ES/H/0392/001/DC	138240	B.BRAUN MELSUNGEN AG	AT
Ibuprofen B. Braun 600 mg Infusionslösung	ES/H/0392/001/DC	BE516240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 600 mg Infusionslösung	ES/H/0392/001/DC	BE516240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 600 mg Infusionslösung	ES/H/0392/001	97003.00.00	B.BRAUN MELSUNGEN AG	DE
Ibuprofen B. Braun 600 mg Infusionslösung	ES/H/0392/001	2019070165	B.BRAUN MELSUNGEN AG	LU
Ibuprofen B. Braun 600 mg infusionsvätska, lösning	ES/H/0392/001/DC	33913	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 600 mg infusionsvätska, lösning	ES/H/0392/001	54209	B.BRAUN MELSUNGEN AG	SE
Ibuprofen B. Braun 600 mg infusjonsvæske, oppløsning	ES/H/0392/001	15-11010	B.BRAUN MELSUNGEN AG	NO
Ibuprofen B. Braun 600 mg infuusioneste, liuos	ES/H/0392/001/DC	33913	B.BRAUN MELSUNGEN AG	FI
Ibuprofen B. Braun 600 mg infuzinis tirpalas	ES/H/0392/001	LT/1/17/4131/004	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 600 mg infuzinis tirpalas	ES/H/0392/001	LT/1/17/4131/003	B.BRAUN MELSUNGEN AG	LT
Ibuprofen B. Braun 600 mg infuzní roztok	ES/H/0392/001/DC	07/867/15-C	B.BRAUN MELSUNGEN AG	CZ
Ibuprofen B. Braun 600 mg infúzny roztok	ES/H/0392/001/DC	07/0155/17-S	B.BRAUN MELSUNGEN AG	SK
Ibuprofen B. Braun 600 mg oldatos infúzió	ES/H/0392/001/DC	OGYI-T-23186/02	B.BRAUN MELSUNGEN AG	HU
Ibuprofen B. Braun 600	ES/H/0392/001/DC	BE516240	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
mg oplossing voor infusie				
Ibuprofen B. Braun 600 mg oplossing voor infusie	ES/H/0392/001/DC	BE516240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 600 mg oplossing voor infusie	ES/H/0392/001	RVG 118618	B.BRAUN MELSUNGEN AG	NL
Ibuprofen B. Braun 600 mg raztopina za infundiranje	ES/H/0392/001/DC	H/17/02364/001	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun 600 mg raztopina za infundiranje	ES/H/0392/001/DC	H/17/02364/002	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun 600 mg šķīdums infūzijām	ES/H/0392/001/DC	17-0110	B.BRAUN MELSUNGEN AG	LV
Ibuprofen B. Braun 600 mg soluție perfuzabilă	ES/H/0392/001	10070/2017/01	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 600 mg soluție perfuzabilă	ES/H/0392/001	10070/2017/02	B.BRAUN MELSUNGEN AG	RO
Ibuprofen B. Braun 600 mg solution for infusion	ES/H/0392/001/DC	PA0736/041/002	B.BRAUN MELSUNGEN AG	IE
Ibuprofen B. Braun 600 mg solution pour perfusion	ES/H/0392/001/DC	BE516240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun 600 mg solution pour perfusion	ES/H/0392/001/DC	BE516240	B.BRAUN MELSUNGEN AG	BE
Ibuprofen B. Braun za otroke 200 mg raztopina za infundiranje	ES/H/0392/002/DC	H/17/02364/003	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun za otroke 200 mg raztopina za infundiranje	ES/H/0392/002/DC	H/17/023/004	B.BRAUN MELSUNGEN AG	SI
Ibuprofen B. Braun, 200 mg/50 mL, roztwór do infuzji	ES/H/0392/002/DC	25841	B.BRAUN MELSUNGEN AG	PL
Ibuprofen B. Braun, 400 mg infusioonilahus	ES/H/0390/001	935517	B.BRAUN MELSUNGEN AG	EE

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, 400 mg/100 mL, roztwór do infuzji	ES/H/0390/001	25190	B.BRAUN MELSUNGEN AG	PL
Ibuprofen B. Braun, 600 mg infusioonilahus	ES/H/0392/001/DC	937417	B.BRAUN MELSUNGEN AG	EE
Ibuprofen B. Braun, 600 mg/100 mL, roztwór do infuzji	ES/H/0392/001	25191	B.BRAUN MELSUNGEN AG	PL
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
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	NL/H/3903/001	12778/2019/10	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
mg capsule moi masticabile				
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
Ibuprofen Banner 100 mg capsule moi masticabile	NL/H/3903/001	12778/2019/17	PATHEON SOFTGELS B.V.	RO
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/001	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/002	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/003	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/004	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100	NL/H/3903/001	LT/1/20/4569/005	PATHEON SOFTGELS B.V.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg kramtomosios minkštosios kapsulės				
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/006	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/007	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/008	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/009	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/010	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/011	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/012	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/013	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/014	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/015	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100 mg kramtomosios minkštosios kapsulės	NL/H/3903/001	LT/1/20/4569/016	PATHEON SOFTGELS B.V.	LT
Ibuprofen Banner 100	NL/H/3903/001	LT/1/20/4569/017	PATHEON SOFTGELS B.V.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg kramtomosios minkštosios kapsulės				
Ibuprofen Banner 100 mg meke kapsule za žvakanje	NL/H/3903/001	HR-H-911536382	PATHEON SOFTGELS B.V.	HR
Ibuprofen Banner 100 mg měkké žvýkací tobolky	NL/H/3903/001	07/125/19-C	PATHEON SOFTGELS B.V.	CZ
Ibuprofen Banner 100 mg mīkstās košļājamās kapsulas	NL/H/3903/001	19-0200	PATHEON SOFTGELS B.V.	LV
Ibuprofen Banner 100 mg soft chewable capsules	NL/H/3903/001	PA22773/001/001	PATHEON SOFTGELS B.V.	IE
Ibuprofen Banner 100 mg Weichkapseln zum Zerbeißen	NL/H/3903/001	139259	PATHEON SOFTGELS B.V.	AT
Ibuprofen Banner 100 mg, soft chewable capsules	NL/H/3903/001	MA1328/00101	PATHEON SOFTGELS B.V.	MT
Ibuprofen Banner 100 mg, zachte kauwcapsules	NL/H/3903/001	RVG 112525	PATHEON SOFTGELS B.V.	NL
Ibuprofen Banner 400 mg kapsel, mjuk	NL/H/4469/001	44549	PATHEON SOFTGELS B.V.	SE
Ibuprofen Banner, 100 mg, kapsułki do żucia, elastyczne	NL/H/3903/001	26150	PATHEON SOFTGELS B.V.	PL
Ibuprofen Billev 800 mg Filmtabletten	PT/H/2609/003	7000532.00.00	ZENTIVA PHARMA GMBH	DE
Ibuprofen Billev farmacija vzhod 800 mg Filmtabletten	PT/H/2719/003	7005756.00.00	ZENTIVA PHARMA GMBH	DE
Ibuprofen Farmalider 20 mg/ml oral suspension	FI/H/0799/001	50518	FARMALIDER, S.A.	DK
Ibuprofen Farmalider 20	FI/H/0799/001	47788	FARMALIDER, S.A.	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
mg/ml oral suspension				
Ibuprofen FARMALIDER 20 mg/ml peroralna suspenzija	SI/H/0162/001/DC	5363-I-1601/12	FARMALIDER, S.A.	SI
Ibuprofen FARMALIDER 20 mg/ml peroralna suspenzija	SI/H/0162/001/DC	5356-I-1602/12	FARMALIDER, S.A.	SI
Ibuprofen FARMALIDER 20 mg/ml peroralna suspenzija	SI/H/0162/001/DC	5363-I-1603/12	FARMALIDER, S.A.	SI
Ibuprofen FARMALIDER 20 mg/ml, zawiesina doustna	SI/H/0162/001/DC	21257	FARMALIDER, S.A.	PL
Ibuprofen Farmalider 40 mg/ml oraalisuspensio	FI/H/0799/002	30596	FARMALIDER, S.A.	FI
Ibuprofen Farmalider 40 mg/ml oral suspension	FI/H/0799/002	50519	FARMALIDER, S.A.	DK
Ibuprofen Farmalider 40 mg/ml oral suspension	FI/H/0799/002	47789	FARMALIDER, S.A.	SE
Ibuprofen Fresenius Kabi 400 mg Infusionslösung	DE/H/6040/001	BE578497	FRESENIUS KABI NV/SA	BE
Ibuprofen Fresenius Kabi 400 mg oplossing voor infusie	DE/H/6040/001	BE578497	FRESENIUS KABI NV/SA	BE
Ibuprofen Fresenius Kabi 400 mg oplossing voor infusie	DE/H/6040/001	RVG 124247	FRESENIUS KABI NEDERLAND B.V.	NL
Ibuprofen Fresenius Kabi 400 mg solution pour perfusion	DE/H/6040/001	BE578497	FRESENIUS KABI NV/SA	BE
Ibuprofen Hermes, 50 mg/g, Gel	not available	43768.00.00	HERMES ARZNEIMITTEL GMBH	DE
Ibuprofen infirst 200 mg soft capsule	not available	PL 51724/0003	INFIRST LTD	XI
Ibuprofen Kabi 400 mg Infusionslösung	DE/H/6040/001	140620	FRESENIUS KABI AUSTRIA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen Kabi 400 mg Infusionslösung	DE/H/6040/001	2203294.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Ibuprofen Kabi 400 mg infuzní roztok	DE/H/6040/001	07/514/18-C	FRESENIUS KABI S.R.O.	CZ
Ibuprofen Kabi 400 mg oldatos infúzió	DE/H/6040/001	OGYI-T-23781/01	FRESENIUS KABI HUNGARY KFT.	HU
Ibuprofen Kabi 400 mg oldatos infúzió	DE/H/6040/001	OGYI-T-23781/03	FRESENIUS KABI HUNGARY KFT.	HU
Ibuprofen Kabi 400 mg oldatos infúzió	DE/H/6040/001	OGYI-T-23781/02	FRESENIUS KABI HUNGARY KFT.	HU
Ibuprofen Kabi 400 mg raztopina za infundiranje	DE/H/6040/001	H/21/02859/001	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Ibuprofen Kabi 400 mg raztopina za infundiranje	DE/H/6040/001	H/21/02859/003	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Ibuprofen Kabi 400 mg raztopina za infundiranje	DE/H/6040/001	H/21/02859/002	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Ibuprofen Kabi 400 mg soluție perfuzabilă	DE/H/6040/001	13592/2020/01	FRESENIUS KABI ROMANIA SRL	RO
Ibuprofen Kabi 400 mg soluție perfuzabilă	DE/H/6040/001	13592/2020/02	FRESENIUS KABI ROMANIA SRL	RO
Ibuprofen Kabi 400 mg soluție perfuzabilă	DE/H/6040/001	13592/2020/03	FRESENIUS KABI ROMANIA SRL	RO
Ibuprofen Kabi 400 mg solution for infusion	DE/H/6040/001	85580	FRESENIUS KABI ESPAÑA S.A.U.	ES
Ibuprofen Kabi 400 mg, infúzny roztok	DE/H/6040/001	07/0252/20-S	FRESENIUS KABI S.R.O.	SK
Ibuprofen Kabi, 4 mg/ml, roztwór do infuzji	DE/H/6040/001	26183	FRESENIUS KABI POLSKA SP. Z O.O.	PL
Ibuprofen Nutra 200 mg perorálna suspenzia vo vrecku	HU/H/0828/001	07/0092/18-S	NUTRA ESSENTIAL OTC, S.L.	SK
Ibuprofen Nutra 200 mg suspensie orala în plic	HU/H/0828/001	10547/2018/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	HU/H/0828/001	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	HU/H/0828/001	10547/2018/03	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	HU/H/0828/001	10547/2018/04	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 200 mg suspensie orală în plic	HU/H/0828/001	10547/2018/05	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 400 mg perorálna suspenzia vo vrecku	HU/H/0828/002	07/0093/18-S	NUTRA ESSENTIAL OTC, S.L.	SK
Ibuprofen Nutra 400 mg suspensie orală în plic	HU/H/0828/002	10548/2018/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra 400 mg suspensie orală în plic	HU/H/0828/002	10548/2018/02	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential 100 mg belsőleges szuszpenzió tasakban	SI/H/0214/001	OGYI-T-23167/04	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 100 mg belsőleges szuszpenzió tasakban	SI/H/0214/001	OGYI-T-23167/05	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 100 mg geriamoji suspensija paketelyje	SI/H/0214/001	LT/1/17/4134/001	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 100 mg geriamoji suspensija paketelyje	SI/H/0214/001	LT/1/17/4134/002	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 100 mg perorálna suspenzia vo vrecku	SI/H/0214/001	07/0293/17-S	NUTRA ESSENTIAL OTC, S.L.	SK
Ibuprofen Nutra Essential 100 mg peroralna suspenzija v	SI/H/0214/001/DC	H/18/02475/001	NUTRA ESSENTIAL OTC, S.L.	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
vrecici (v nadaljevanju: Ibuprofen Nutra Essential 100 mg)				
Ibuprofen Nutra Essential 100 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 100 mg)	SI/H/0214/001/DC	H/18/02475/002	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 100 mg suspensija iekšķīgai lietošanai paciņā	SI/H/0214/001	17-0213	NUTRA ESSENTIAL OTC, S.L.	LV
Ibuprofen Nutra Essential 100 mg Suspension zum Einnehmen	SI/H/0214/001	137944	NUTRA ESSENTIAL OTC, S.L.	AT
Ibuprofen Nutra Essential 100 mg Suspension zum Einnehmen im Beutel	SI/H/0214/001	97819.00.00	NUTRA ESSENTIAL OTC, S.L.	DE
Ibuprofen Nutra Essential 100 mg suukaudne suspensioon kotikeses	SI/H/0214/001	949317	NUTRA ESSENTIAL OTC, S.L.	EE
Ibuprofen Nutra Essential 200 mg belsoleges szuszpenzió tasakban	SI/H/0214/002	OGYI-T-23167/06	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 200 mg belsoleges szuszpenzió tasakban	SI/H/0214/002	OGYI-T-23167/07	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 200 mg belsoleges szuszpenzió	SI/H/0214/002	OGYI-T-23167/08	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
tasakban				
Ibuprofen Nutra Essential 200 mg de suspension buvable en sachet	FR/H/0786/001	BE518657	NUTRA ESSENTIAL OTC, S.L.	BE
Ibuprofen Nutra Essential 200 mg geriamoji suspensija paketėlyje	SI/H/0214/002	LT/1/17/4134/003	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 200 mg geriamoji suspensija paketėlyje	SI/H/0214/002	LT/1/17/4134/004	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 200 mg geriamoji suspensija paketėlyje	SI/H/0214/002	LT/1/17/4134/005	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 200 mg perorálna suspenzia vo vrecku	SI/H/0214/002	07/0294/17-S	NUTRA ESSENTIAL OTC, S.L.	SK
Ibuprofen Nutra Essential 200 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 200 mg)	SI/H/0214/002/DC	H/18/02475/005	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 200 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 200 mg)	SI/H/0214/002/DC	H/18/02475/004	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 200 mg peroralna suspenzija v	SI/H/0214/002/DC	H/18/02475/003	NUTRA ESSENTIAL OTC, S.L.	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
vrečici (v nadaljevanju: Ibuprofen Nutra Essential 200 mg)				
Ibuprofen Nutra Essential 200 mg suspensie voor oraal gebruik in sachet	FR/H/0786/001	BE518657	NUTRA ESSENTIAL OTC, S.L.	BE
Ibuprofen Nutra Essential 200 mg suspensija iekšķīgai lietošanai paciņā	SI/H/0214/002	17-0214	NUTRA ESSENTIAL OTC, S.L.	LV
Ibuprofen Nutra Essential 200 mg Suspension zum Einnehmen im Beutel	FR/H/0786/001	BE518657	NUTRA ESSENTIAL OTC, S.L.	BE
Ibuprofen Nutra Essential 200 mg Suspension zum Einnehmen im Beutel	SI/H/0214/002	97820.00.00	NUTRA ESSENTIAL OTC, S.L.	DE
Ibuprofen Nutra Essential 200 mg suukaudne suspensioon kotikeses	SI/H/0214/002	949417	NUTRA ESSENTIAL OTC, S.L.	EE
Ibuprofen Nutra Essential 400 mg belsoleges szuszpenzió tasakban	SI/H/0214/003	OGYI-T-23167/09	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 400 mg belsoleges szuszpenzió tasakban	SI/H/0214/003	OGYI-T-23167/10	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra Essential 400 mg belsoleges szuszpenzió tasakban	SI/H/0214/003	OGYI-T-23167/11	NUTRA ESSENTIAL OTC, S.L.	HU
Ibuprofen Nutra	SI/H/0214/003	LT/1/17/4134/006	NUTRA ESSENTIAL OTC, S.L.	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
Essential 400 mg geriamoji suspensija paketėlyje				
Ibuprofen Nutra Essential 400 mg geriamoji suspensija paketėlyje	SI/H/0214/003	LT/1/17/4134/007	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 400 mg geriamoji suspensija paketėlyje	SI/H/0214/003/DC	LT/1/17/4134/008	NUTRA ESSENTIAL OTC, S.L.	LT
Ibuprofen Nutra Essential 400 mg perorálna suspenzia vo vrecku	SI/H/0214/003	07/0295/17-S	NUTRA ESSENTIAL OTC, S.L.	SK
Ibuprofen Nutra Essential 400 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 400 mg)	SI/H/0214/003/DC	H/18/02475/008	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 400 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 400 mg)	SI/H/0214/003/DC	H/18/02475/006	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 400 mg peroralna suspenzija v vrečici (v nadaljevanju: Ibuprofen Nutra Essential 400 mg)	SI/H/0214/003/DC	H/18/02475/007	NUTRA ESSENTIAL OTC, S.L.	SI
Ibuprofen Nutra Essential 400 mg suspensie voor oraal	FR/H/0786/002	BE518675	NUTRA ESSENTIAL OTC, S.L.	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
gebruik in sachet				
Ibuprofen Nutra Essential 400 mg suspensija iekšķīgai lietošanai paciņā	SI/H/0214/003	17-0215	NUTRA ESSENTIAL OTC, S.L.	LV
Ibuprofen Nutra Essential 400 mg suspension buvable en sachet	FR/H/0786/002	BE518675	NUTRA ESSENTIAL OTC, S.L.	BE
Ibuprofen Nutra Essential 400 mg Suspension zum Einnehmen im Beutel	FR/H/0786/002	BE518675	NUTRA ESSENTIAL OTC, S.L.	BE
Ibuprofen Nutra Essential 400 mg Suspension zum Einnehmen im Beutel	SI/H/0214/003	97821.00.00	NUTRA ESSENTIAL OTC, S.L.	DE
Ibuprofen Nutra Essential 400 mg suukaudne suspensioon kotikeses	SI/H/0214/003	949517	NUTRA ESSENTIAL OTC, S.L.	EE
Ibuprofen Nutra Essential OTC 100 mg suspensie orală în plic	SI/H/0214/001	10393/2017/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential OTC 100 mg suspensie orală în plic	SI/H/0214/001	10393/2017/02	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential OTC 200 mg suspensie orală în plic	SI/H/0214/002	10394/2017/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential OTC 200 mg suspensie orală în plic	SI/H/0214/002	10394/2017/02	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential OTC 200 mg suspensie orală în plic	SI/H/0214/002	10394/2017/03	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 Essential OTC 400 mg suspensie orală în plic	SI/H/0214/003	10395/2017/01	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential OTC 400 mg suspensie orală în plic	SI/H/0214/003	10395/2017/02	NUTRA ESSENTIAL OTC, S.L.	RO
Ibuprofen Nutra Essential S.L. 100 mg perorální suspenze v sáčku	SI/H/0214/001/DC	07/662/16-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra Essential S.L. 200 mg perorální suspenze v sáčku	SI/H/0214/002/DC	07/663/16-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Nutra Essential S.L. 400 mg perorální suspenze v sáčku	SI/H/0214/003	07/664/16-C	NUTRA ESSENTIAL OTC, S.L.	CZ
Ibuprofen Patheon 100 mg, zachte kauwcapsules	not available	RVG 123943	PATHEON SOFTGELS B.V.	NL
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
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
Ibuprofen Sanias 200 mg, filmomhulde tabletten	NL/H/4164/001	RVG 101773	AUROBINDO PHARMA B.V.	NL
Ibuprofen Sanias 400	NL/H/4164/002	RVG 101775	AUROBINDO PHARMA 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
mg, filmomhulde tabletten				
Ibuprofen Seven Plus Pain Relief 200 mg/ 5ml oral suspension	not available	PL35533/0056	ASPIRE PHARMA LIMITED	XI
Ibuprofen Skillpharma 400 mg granule pentru solutie orala	CZ/H/1073/001	14708/2022/01	SKILLPHARMA S.R.L.	RO
IBUPROFEN SKILLPHARMA 400 mg granule pro perorální roztok	CZ/H/1073/001	07/455/20-C	SKILLPHARMA S.R.L.	CZ
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
Ibuprofen STADA 200 mg peroralni prašek	DE/H/4307/001	H/17/02359/003	STADA ARZNEIMITTEL AG	SI
Ibuprofen STADA 200 mg perorálny prášok	DE/H/4307/001	07/0226/17-S	STADA ARZNEIMITTEL AG	SK
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
Ibuprofen STADA 400 mg perorálny prášok	DE/H/4307/002	07/0227/17-S	STADA ARZNEIMITTEL AG	SK
Ibuprofen Tablets 400 mg.	not available	PL 17907/0079	BRISTOL LABORATORIES LIMITED	XI
Ibuprofen Tablets BP 400 mg	not available	PL 00063/0756	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Ibuprofen Teva 5 % gel	not available	BE179715	TEVA PHARMA BELGIUM 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
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 Twelve Plus Pain Relief 200mg/5ml oral suspension	not available	PL35533/0034	ASPIRE PHARMA LIMITED	XI
Ibuprofen US Pharmacia 200 mg comprimate filmate	PL/H/0709/001-002/DC	14367/2022/01-06	US PHARMACIA SP. Z O.O.	RO
Ibuprofen US Pharmacia 200 mg comprimate filmate	PL/H/0709/001-002/DC	14367/2022/01-06	US PHARMACIA SP. Z O.O.	RO
Ibuprofen Zentiva 800 mg filmdragerade tabletter	SE/H/2058/003	60149	ZENTIVA, K.S.	SE
Ibuprofen Zentiva 800 mg kalvopäällysteiset tabletit	SE/H/2058/003	37741	ZENTIVA, K.S.	FI
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001/DC	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001/DC	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001/DC	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001/DC	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
Ibuprofen/Coffein Sanofi 400 mg/100 mg	DE/H/5544/001/DC	2201926.00.00	A. NATTERMANN & CIE. 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
Filmtabletten				
Ibuprofen/Coffein Sanofi 400 mg/100 mg Filmtabletten	DE/H/5544/001	2201926.00.00	A. NATTERMANN & CIE. GMBH	DE
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 3 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 4 2	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 5 9	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 6 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 7 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 720 8 0	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 1 0	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 0 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 1 7	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 2 4	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé	IE/H/0755/001	34009 550 259 3 1	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
pelliculé				
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 4 8	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 5 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 2 7	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 3 4	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 4 1	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 6 5	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 7 2	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 8 9	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 721 9 6	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 300 722 0 2	ACCORD HEALTHCARE FRANCE SAS	FR
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	IE/H/0755/001	34009 550 259 8 6	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
pelliculé				
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	34009 550 259 9 3	ACCORD HEALTHCARE FRANCE SAS	FR
IBUPROFENE ACCORD 200 mg, comprimé pelliculé	IE/H/0755/001	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
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553173	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553108	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553110	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553122	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553134	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg compresse rivestite con film	DE/H/5687/004	047553146	ARISTO PHARMA GMBH (ART 57)	IT
Ibuprofene Aristo Pharma 800 mg	DE/H/5687/004	047553159	ARISTO PHARMA GMBH (ART 57)	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
compresse rivestite con film				
IBUPROFENE ARROW LAB 200 mg, comprimé pelliculé	not available	NL48215	ARROW GENERIQUES	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
IBUPROFENE B. BRAUN 200 mg, solution pour perfusion	ES/H/0392/002	34009 302 001 3 1	B.BRAUN MELSUNGEN AG	FR
IBUPROFENE B. BRAUN 200 mg, solution pour perfusion	ES/H/0392/002	34009 550 713 7 2	B.BRAUN MELSUNGEN AG	FR
Ibuprofene B. Braun 400 mg Soluzione per infusione	ES/H/0390/001	045126012	B.BRAUN MELSUNGEN AG	IT
Ibuprofene B. Braun 400 mg Soluzione per infusione	ES/H/0390/001	045126024	B.BRAUN MELSUNGEN AG	IT
IBUPROFENE B. BRAUN	ES/H/0390/001	34009 550 560 6 5	B.BRAUN MELSUNGEN AG	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
400 mg/100 mL, solution pour perfusion				
IBUPROFENE B. BRAUN 400 mg/100 mL, solution pour perfusion	ES/H/0390/001	34009 550 560 5 8	B.BRAUN MELSUNGEN AG	FR
IBUPROFENE B. BRAUN 600 mg/100 mL, solution pour perfusion	ES/H/0392/001/DC	34009 550 560 7 2	B.BRAUN MELSUNGEN AG	FR
IBUPROFENE B. BRAUN 600 mg/100 mL, solution pour perfusion	ES/H/0392/001/DC	34009 550 560 8 9	B.BRAUN MELSUNGEN AG	FR
Ibuprofene B. Braun Melsungen 200 mg soluzione per infusione	ES/H/0392/002/DC	045125034	B.BRAUN MELSUNGEN AG	IT
Ibuprofene B. Braun Melsungen 200 mg soluzione per infusione	ES/H/0392/002/DC	045125046	B.BRAUN MELSUNGEN AG	IT
Ibuprofene B. Braun Melsungen 600 mg Soluzione per infusione	ES/H/0392/001/DC	045125010	B.BRAUN MELSUNGEN AG	IT
Ibuprofene B. Braun Melsungen 600 mg Soluzione per infusione	ES/H/0392/001/DC	045125022	B.BRAUN MELSUNGEN AG	IT
Ibuprofene Banner 100 mg capsules molles à mâcher	NL/H/3903/001	34009 302 174 6 7	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg capsules molles à mâcher	NL/H/3903/001	34009 302 174 7 4	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 1 1	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 2 8	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100	NL/H/3903/001	34009 302 373 3 5	PATHEON SOFTGELS B.V.	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
mg, capsule molle à mâcher				
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 5 9	PATHEON SOFTGELS B.V.	FR
IBUPROFENE BANNER 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 6 6	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 7 3	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 8 0	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 373 9 7	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 1 0	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 3 4	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 2 7	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 4 1	PATHEON SOFTGELS B.V.	FR
IBUPROFENE BANNER 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 5 8	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100 mg, capsule molle à mâcher	NL/H/3903/001	34009 302 374 7 2	PATHEON SOFTGELS B.V.	FR
Ibuprofene Banner 100	NL/H/3903/001	34009 302 374 6 5	PATHEON SOFTGELS B.V.	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
mg, capsule molle à mâcher				
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130087	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400955048398	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130100	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400955048411	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130070	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130094	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au	FR/H/0786/001/DC	3400930130087	BIOGARAN	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
maltitol liquide				
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400955048398	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130100	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400955048411	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130070	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 200 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/001/DC	3400930130094	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400930130124	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400955048459	BIOGARAN	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 BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400955048480	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400955048466	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400955048442	BIOGARAN	FR
IBUPROFENE BIOGARAN CONSEIL 400 mg, suspension buvable en sachet édulcorée au maltitol liquide	FR/H/0786/002/DC	3400955048473	BIOGARAN	FR
IBUPROFENE MYLAN ITALIA 400 mg compresse rivestite	not available	042995011	MYLAN ITALIA S.R.L.	IT
Ibuprofene Skillpharma 400 mg granulato per soluzione orale	CZ/H/1073/001	050101017	SKILLPHARMA S.R.L.	IT
IBUPROFENE VIATRIS 200 mg, comprimé enrobé	not available	NL 26874	VIATRIS SANTE	FR
Ibuprofene Welcome Pharma 200 mg sospensione orale in bustina, gusto arancia	not available	041997038	WELCOME PHARMA SPA	IT
Ibuprofene Welcome Pharma 200 mg sospensione orale in	not available	041997040	WELCOME PHARMA 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
bustina, gusto arancia				
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324032	ZENTIVA ITALIA S.R.L.	IT
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324018	ZENTIVA ITALIA S.R.L.	IT
IBUPROFENE ZENTIVA ITALIA 200 mg compresse rivestite con film	not available	042324020	ZENTIVA ITALIA S.R.L.	IT
IBUPROFENE ZF 400 mg, comprimé pelliculé	not available	34009 417 592 0 8	ZYDUS FRANCE	FR
IBUPROFENE ZF 400 mg, comprimé pelliculé	not available	34009 417 591 4 7	ZYDUS FRANCE	FR
IBUPROFENE ZF 400 mg, comprimé pelliculé	not available	34009 417 593 7 6	ZYDUS FRANCE	FR
IBUPROFENE ZF 400 mg, comprimé pelliculé	not available	34009 417 594 3 7	ZYDUS FRANCE	FR
Ibuprofeno + Cafeína Aspegic 400 mg + 100 mg comprimidos revestidos por película	DE/H/5544/001	5797808	OPELLA HEALTHCARE PORTUGAL UNIPESSOAL LDA.	PT
Ibuprofeno B. Braun 400 mg Solução para perfusão	ES/H/0390/001	5710637	B.BRAUN MELSUNGEN AG	PT
Ibuprofeno B. Braun 400 mg Solução para perfusão	ES/H/0390/001	5710645	B.BRAUN MELSUNGEN AG	PT
Ibuprofeno B. Braun 400 mg solución para perfusion	ES/H/0390/001	82170	B.BRAUN MELSUNGEN AG	ES
Ibuprofeno B. Braun 600 mg solução para	ES/H/0392/001	5714928	B.BRAUN MELSUNGEN AG	PT

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perfusão				
Ibuprofeno B. Braun 600 mg solução para perfusão	ES/H/0392/001	5714936	B.BRAUN MELSUNGEN AG	PT
Ibuprofeno B. Braun 600 mg solución para perfusión	not available	80720	B.BRAUN MEDICAL, S.A.	ES
Ibuprofeno B. Braun pediátrico 200 mg solución para perfusión	ES/H/0392/002/DC	85161	B.BRAUN MELSUNGEN AG	ES
Ibuprofeno B.Braun 600 mg concentrado para solución para perfusión	not available	79593	B.BRAUN MEDICAL, S.A.	ES
Ibuprofeno Braun 400 mg solución para perfusión	not available	82808	B.BRAUN MEDICAL, S.A.	ES
Ibuprofeno Braun 600 mg solución para perfusión	ES/H/0392/001/DC	82171	B.BRAUN MELSUNGEN AG	ES
Ibuprofeno Clover 200 mg capsulas blandas.	not available	80987	HC CLOVER PRODUCTOS Y SERVICIOS, S.L.	ES
Ibuprofeno Clover 600 mg capsulas blandas	not available	80976	HC CLOVER PRODUCTOS Y SERVICIOS, S.L.	ES
Ibuprofeno Farmalider 20 mg/ml suspensão oral	SI/H/0162/001/DC	5442108	FARMALIDER, S.A.	PT
Ibuprofeno Farmalider 20 mg/ml suspensão oral	SI/H/0162/001/DC	5442116	FARMALIDER, S.A.	PT
Ibuprofeno Farmalider 20 mg/ml suspensão oral	SI/H/0162/001/DC	5442124	FARMALIDER, S.A.	PT
Ibuprofeno Farmalider 400 mg concentrado para solución para perfusión	not available	83570	FARMALIDER, S.A.	ES
Ibuprofeno Kabi 400 mg Solução para Perfusão	DE/H/6040/001	5808407	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ibuprofeno Kabi 400 mg	DE/H/6040/001	5808415	FRESENIUS KABI PHARMA	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
Solução para Perfusão			PORTUGAL, LDA.	
Ibuprofeno Kabi 400 mg Solução para Perfusão	DE/H/6040/001	5808423	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ibuprofeno Stadapharm 400 mg cápsulas blandas	not available	80.975	LABORATORIO STADA, S.L.	ES
Ibuprofeno Worldrugs 40 mg/ml suspensão oral	not available	5677166	LDP TORLAN LDA.	PT
Ibuprofenum US Pharmacia 200 mg apvalkotās tabletes	PL/H/0709/001-002/DC	22-0032	US PHARMACIA SP. Z O.O.	LV
Ibuprofenum US Pharmacia 200 mg apvalkotās tabletes	PL/H/0709/001-002/DC	22-0032	US PHARMACIA SP. Z O.O.	LV
Ibuprofenum US Pharmacia, 200 mg ohukese polümeerikattega tabletid	PL/H/0709/001-002/DC	1054822	US PHARMACIA SP. Z O.O.	EE
Ibuprofenum US Pharmacia, 200 mg ohukese polümeerikattega tabletid	PL/H/0709/001-002/DC	1054822	US PHARMACIA SP. Z O.O.	EE
IBUPROM EFFECT żel, 50 mg/g, żel	not available	23940	US PHARMACIA SP. Z O.O.	PL
IBUPROM MAX RAPID, 400 mg, tabletki powlekane	not available	26606	US PHARMACIA SP. Z O.O.	PL
IBUPROM REGULAR, 200 mg, tabletki powlekane	PL/H/0709/002	26994	US PHARMACIA SP. Z O.O.	PL
IBUPROM REGULAR, 200 mg, tabletki powlekane	PL/H/0709/002	26994	US PHARMACIA SP. Z O.O.	PL
IBUPROM SPORT spray, 50 mg/g, aerozol na skórę, roztwór	not available	24970	US PHARMACIA SP. Z O.O.	PL
IBUPROM ULTRAMAX	not available	26756	US PHARMACIA 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
SPRINT, 600 mg, kapsułki, miękkie				
IBUPROM ULTRAMAX SPRINT, 600 mg, kapsułki, miękkie	not available	26756	US PHARMACIA SP. Z O.O.	PL
IBU-ratiopharm® direkt 200 mg Pulver zum Einnehmen	DE/H/6731/001	94807.00.00	RATIOPHARM GMBH	DE
IBU-SPA, 400 mg + 100 mg, tabletki powlekane	DE/H/5544/001/DC	25950	OPELLA HEALTHCARE POLAND SP. Z O.O.	PL
IBU-SPA, 400 mg + 100 mg, tabletki powlekane	DE/H/5544/001/DC	25950	OPELLA HEALTHCARE POLAND SP. Z O.O.	PL
Ibuspray	not available	PL 00173/0150	DIOMED DEVELOPMENTS LIMITED	XI
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
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
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	DE/H/2598/002	LT/1/15/3829/004	BERLIN-CHEMIE AG	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
geriamoji suspensija, vaikams				
Ibustar bērniem 100 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2463/001	11-0290	BERLIN-CHEMIE AG	LV
Ibustar bērniem 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2598/002	15-0268	BERLIN-CHEMIE AG	LV
Ibustar forte, 40 mg/ml suukaudne suspensioon	DE/H/2598/002	888015	BERLIN-CHEMIE AG	EE
Ibustar, 20 mg/ml suukaudne suspensioon	DE/H/2463/001	752311	BERLIN-CHEMIE AG	EE
Ibutop 5% geel	not available	280299	DOLORGIET GMBH & CO KG	EE
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
Ibutop Gel 50 mg/g gel	not available	7878/2015/01	DOLORGIET GMBH & CO KG	RO
Ibutop, creme	not available	18905	DOLORGIET GMBH & CO KG	DK
IbuViva 200mg/5ml geriamoji suspensiija	IE/H/0690/001/DC	LT/1/10/2162/005-009	PINEWOOD LABORATORIES LIMITED	LT
Ibux 40 mg/ml mikstur, suspensjon	IE/H/0690/001	18-12289	KARO PHARMA AS	NO
Ibux 50 mg/g gel	not available	99-553	KARO PHARMA AS	NO
Ibux 600 mg tabletter, filmdrasjerte	not available	0000-07069	KARO PHARMA AS	NO
ibuxin rapid 400 mg filmuhúðaðar töflur	IS/H/0353/001	IS/1/12/080/01	RATIOPHARM GMBH	IS
Ibuxin rapid 400 mg tabletti, kalvopäällysteinen	DE/H/2590/001	28002	RATIOPHARM GMBH	FI
Ibuxin Rapid 684 mg filmom obložene tablete	not available	HR-H-064892441	PLIVA HRVATSKA D.O.O.	HR
Imofen 20 mg/ml suspensão oral	not available	5677158	CUIDAFARMA, LDA.	PT
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 1 6	OPELLA 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
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 3 0	OPELLA HEALTHCARE FRANCE SAS	FR
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 2 3	OPELLA HEALTHCARE FRANCE SAS	FR
IPRAFEINE 400 mg/100 mg, comprimé pelliculé	DE/H/4270/001	34009 300 749 0 9	OPELLA HEALTHCARE FRANCE SAS	FR
Ipren 125 mg suppositorier	not available	15684	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
Ipren 20 mg/ml, oral suspension	not available	19260	MCNEIL SWEDEN AB	SE
Ipren 5% gel	not available	18362	MCNEIL SWEDEN AB	SE
Ipren 5% gel	not available	18362	MCNEIL SWEDEN AB	SE
Ipren 60 mg suppositorier	not available	46199	MCNEIL SWEDEN AB	SE
Ipren, filmovertrukne tabletter	not available	13445	MCNEIL DENMARK APS	DK
Iprensa 400 mg kapslar, mjuka	not available	43284	MCNEIL SWEDEN AB	SE
Iprensa 400 mg kapslar, mjuka	not available	43284	MCNEIL SWEDEN AB	SE
JOINTRAL 100 mg/g gel	not available	040608010	LAB. IT. BIOCHIM. FARM.CO LISAPHARMA S.P.A.	IT
Junifen 40 mg/ml suspensión oral sabor naranja	DE/H/2206/001	73.047	RECKITT BENCKISER HEALTHCARE S.A.	ES
Junipro 40 mg/ml suspensión oral sabor fresa	DE/H/2206/002	73048	RECKITT BENCKISER HEALTHCARE S.A.	ES
Kidofen max, 250 mg/5 mL, zawiesina doustna	not available	26590	AFLOFARM FARMACJA POLSKA SP. Z O.O.	PL
LASONIL ANTIDOLORE	not available	042154029	BAYER SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
10% gel				
LASONIL ANTIDOLORE 10% gel	not available	042154017	BAYER SPA	IT
Lloydspharmacy Ibuprofen 200mg Long Lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
MIG 4% perorálna suspenzia 40 mg/ml perorálna suspenzia	DE/H/2598/002	29/0315/15-S	BERLIN-CHEMIE AG	SK
MIG dla dzieci Forte o smaku truskawkowym, 40 mg/ml, zawiesina doustna	DE/H/2598/002	21156	BERLIN-CHEMIE AG	PL
MIG dla dzieci, 20 mg/ml, zawiesina doustna	DE/H/2463/001	18904	BERLIN-CHEMIE AG	PL
MIG Junior 2% 20 mg/ml perorálna suspenzia	DE/H/2463/001	29/0344/11-S	BERLIN-CHEMIE AG	SK
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
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
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
Moment 200 mg compresse masticabili	not available	025669096	AZIENDE CHIMICHE RIUNITE ANGELINI	IT

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			FRANCESCO - A.C.R.A.F. S.P.A.	
Moment 200 mg compresse masticabili	not available	025669108	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
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	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
			FRANCESCO - A.C.R.A.F. S.P.A.	
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
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	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	025669072	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Moment 200 mg compresse rivestite.	not available	025669122	AZIENDE CHIMICHE RIUNITE ANGELINI	IT

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			FRANCESCO - A.C.R.A.F. S.P.A.	
Moment 200 mg compresse rivestite.	not available	025669161	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 granulato per soluzione orale	not available	025669211	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
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
Moment 200 mg sospensione orale in	not available	025669387	AZIENDE CHIMICHE RIUNITE ANGELINI	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
bustine.			FRANCESCO - A.C.R.A.F. S.P.A.	
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 Orosolubile 200 mg polvere orale	IT/H/0745/001	046054019	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
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	IT

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			FRANCESCO - A.C.R.A.F. S.P.A.	
Momentact 400 mg capsule molli	not available	035618038	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg capsule molli	not available	035618040	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con film	not available	035618154	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con film	not available	035618166	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
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
Momentact 400 mg compresse rivestite con film.	not available	035618178	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con	not available	035618180	AZIENDE CHIMICHE RIUNITE ANGELINI	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
film.			FRANCESCO - A.C.R.A.F. S.P.A.	
Momentact 400 mg compresse rivestite con film.	not available	035618192	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg compresse rivestite con film.	not available	035618204	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momentact 400 mg sospensione orale in bustine	not available	035618065	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
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	not available	035618127	AZIENDE CHIMICHE RIUNITE ANGELINI	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
bustine			FRANCESCO - A.C.R.A.F. S.P.A.	
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
Morrisons Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Neobrufen 400 mg Brausegranulat	SE/H/1184/001	1-31789	MYLAN ÖSTERREICH GMBH	AT
Neobrufen 600 mg granulado efervescente	not available	61610	MYLAN IRE HEALTHCARE LIMITED	ES
Neobrufen retard 800 mg comprimidos de liberación prolongada	not available	61618	MYLAN IRE HEALTHCARE LIMITED	ES
Noubid 200 mg film-coated Tablets	not available	PL 44041/0057	NOUMED LIFE SCIENCES	XI
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 936 2 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 937 9 7	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 938 5 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 939 1 9	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	34009 364 947 4 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 341 684 7 5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg,	not available	34009 364 945 1 0	RECKITT BENCKISER	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
comprimé enrobé			HEALTHCARE FRANCE	
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 946 8 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 943 9 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 944 5 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 941 6 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 942 2 0	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 364 935 6 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUREFLEX 400 mg, comprimé enrobé	not available	34009 301 861 4 5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nureflex Junior Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2204/002	1-30062	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nureflex Junior Orange 100 mg Weichkapseln zum Zerbeißen	DE/H/6014/001	137572	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nureflex Junior Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2204/001	1-30061	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nureflex Junior Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2204/001	1-30061	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurodon Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2343/001	77580.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen ® 200mg Καψάκια, μαλακά	DE/H/1482/001	20918	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen ® 200mg Μαλακές Κάψουλες	DE/H/1482/001	2075015	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
Nurofen ® Express 400	NL/H/4313/001	21486	RECKITT BENCKISER	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
mg, καψάκιο, μαλακό			HELLAS HEALTHCARE SA	
Nurofen 100 mg Cápsulas moles para mastigar	DE/H/6014/001	5697016	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 100 mg Cápsulas moles para mastigar	DE/H/6014/001	5697024	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 100 mg Cápsulas moles para mastigar	DE/H/6014/001	5697008	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 100 mg Cápsulas moles para mastigar	DE/H/6014/001	5697032	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen 100 mg tyggekapsler, myke	BE/H/0320/001	15-10700	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
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
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 200 Fastcaps 200 mg Weichkapseln	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 capsules molles	not available	2014090205	RECKITT BENCKISER HEALTHCARE (BELGIUM)	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			SA/NV	
Nurofen 200 Fastcaps, 200 mg zachte capsules	not available	BE442346	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 200 mg - Dragees	not available	14712	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
NUROFEN 200 mg apvalkotas tabletes	not available	01-0449	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen 200 mg ārstnieciskais plāksteris	DE/H/5067/001	19-0078	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen 200 mg ārstnieciskais plāksteris	DE/H/5067/001	19-0087	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen 200 mg bevont tabletta	not available	OGYI-T-6793/67	RECKITT BENCKISER KFT	HU
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
Nurofen 200 mg Caplets	not available	PL 00063/0386	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen 200 mg Caplets	not available	PL 00063/0387	RECKITT BENCKISER	XI

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
			HEALTHCARE (UK) LTD	
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 comprimidos revestidos	not available	4348686	RECKITT BENCKISER HEALTHCARE, LDA.	PT
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 dengtos tabletės	not available	LT/1/97/0271/003	RECKITT BENCKISER (POLAND) S.A.	LT
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
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 200 mg emplastru medicamentos	DE/H/5067/001	14199/2021/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	14199/2021/01	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 200 mg emplastru medicamentos	DE/H/5067/001	14199/2021/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	14199/2021/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg emplastru medicamentos	DE/H/5067/001	14199/2021/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen 200 mg gyógyszeres tapasz	DE/H/5067/001	OGYI-T-6793/140	RECKITT BENCKISER KFT	HU
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
Nurofen 200 mg léčivá náplast	DE/H/5067/001	29/141/18-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen 200 mg liečivá náplasť	DE/H/5067/001	29/0369/18-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen 200 mg ljekoviti flaster	DE/H/5067/001	HR-H-092708535	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 200 mg obalené tablety	not available	07/376/92-S/C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen 200 mg obložene tablete	not available	HR-H-151406116	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 200 mg Tablets	not available	PL 00063/0388	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen 200 mg Tablets	not available	PL 00063/0385	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
NUROFEN 200 mg obalené tablety	not available	29/0376/92-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	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
NUROFEN 200 mg, comprimé enrobé	not available	34009 339 643 5 1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	34009 384 142 1 9	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	34009 360 763 6 5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	34009 360 917 3 3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 200 mg, comprimé enrobé	not available	34009 339 642 9 0	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen 200 tablet ovaal, omhulde tabletten 200 mg	not available	RVG 25190	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen 200 tablet, omhulde tabletten 200 mg	not available	RVG 10674	RECKITT BENCKISER HEALTHCARE B.V.	NL
NUROFEN 200, 200 mg comprimés enrobés	not available	BE132011	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen 200, 200 mg comprimés enrobés	not available	0209/11041123	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 200, 200 mg omhulde tabletten	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 200mg Coated Tablets	not available	9734	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen 200mg Coated Tablets	not available	PA 979/32/6	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen 200mg Dragees	not available	14712	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen 200mg Liquicaps Pharmacy Only	not available	PL 00063/0654	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI

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 Liquid Capsules	not available	PA 979/32/12	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen 200mg Liquid Capsules	not available	PA 979/32/12	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen 200mg Liquid Capsules	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen 24-Stunden Schmerzpflaster 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	138593	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen 24-Stunden Schmerzpflaster 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	138593	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen 24-Stunden Schmerzpflaster 200 mg wirkstoffhaltiges Pflaster	DE/H/5067/001	93368.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen 40 mg/ml mikstur, suspensjon med appelsinsmak	DE/H/2204/001	08-6424	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nurofen 40 mg/ml mikstur, suspensjon med appelsinsmak	DE/H/2204/001	08-6424	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nurofen 40 mg/ml mikstur, suspensjon med jordbærsmak	DE/H/2204/002	08-6425	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	NO
Nurofen 40 mg/ml suspensión oral	not available	65.526	RECKITT BENCKISER HEALTHCARE S.A.	ES
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 Fastcaps 400 mg Weichkapseln	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

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 Fastcaps, 400 mg capsules molles	not available	0209/11041125	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400 Fastcaps, 400 mg capsules, zacht	not available	BE305724	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 FASTTABS 400 mg, comprimés pelliculés	not available	0209/03070141	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN 400 FASTTABS 400 mg, comprimés pelliculés.	not available	BE238551	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400 FASTTABS 400 mg, filmomhulde tabletten.	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 mg Capsule, soft	NL/H/4313/001	PL 00063/0615	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
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 400 mg comprimidos recubiertos	not available	67.968	RECKITT BENCKISER HEALTHCARE S.A.	ES
Nurofen 400 mg obalené tablety	not available	29/0054/13-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen 400 mg obložene tablete	not available	UP/I-530-09/12-01/127	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen 400 mg Weichkapseln	NL/H/4313/001	78463.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
NUROFEN 400 mg, comprimé enrobé	not available	34009 368 647 5 7	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg,	not available	34009 368 649 8 6	RECKITT BENCKISER	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
comprimé enrobé			HEALTHCARE FRANCE	
NUROFEN 400 mg, comprimé enrobé	not available	34009 368 650 6 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	34009 368 652 9 7	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	34009 368 653 5 8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	34009 381 420 0 6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	34009 381 419 2 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN 400 mg, comprimé enrobé	not available	34009 368 646 9 6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
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 400 tablet, omhulde tabletten 400 mg	not available	RVG 22100	RECKITT BENCKISER HEALTHCARE B.V.	NL
NUROFEN 400, 400 mg comprimés enrobés	not available	BE189926	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN 400, 400 mg comprimés enrobés	not available	0209/11041124	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
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
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

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	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 400, 400 mg omhulde tabletten	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 400mg Tablets	not available	PL 00063/0722	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen 400mg Tablets	not available	PL 00063/0722	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen 400mg Tablets	not available	PL 00063/0722	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen 5% w/w Gel	not available	PA 979/32/3	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen 60 mg végbélkúp gyermekeknek	not available	OGYI-T-6793/04	RECKITT BENCKISER KFT	HU
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 Advance 200mg Tablets	not available	PA0979/075/001	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Apelsin, 40 mg/ml, oral suspension	DE/H/2204/001	41889	RECKITT BENCKISER HEALTHCARE	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
			(SCANDINAVIA) A/S	
Nurofen Apelsin, 40 mg/ml, oral suspension	DE/H/2204/001	41889	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	SE
Nurofen Back Pain 300mg Sustained Release Capsules	not available	PL 00063/0378	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
NUROFEN bērniem 100 mg/5 ml suspensija iekšķīgai lietošanai	not available	01-0450	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen dla dzieci Forte pomarańczowy, 40 mg/ml, zawiesina doustna	DE/H/2206/001	22448	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen dla dzieci Forte truskawkowy, 40 mg/ml, zawiesina doustna	DE/H/2206/002	22449	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen dla dzieci Junior pomarańczowy, 40 mg/ml, zawiesina doustna	DE/H/2343/001	17854	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen dla dzieci Junior truskawkowy, 40 mg/ml, zawiesina doustna	DE/H/2343/002	17855	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen dla dzieci Junior, 100 mg, kapsułki do żucia, elastyczne	PL/H/0602/001	23941	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Durance 200 mg medicated plasters	DE/H/5067/001	PA0979/032/018	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Durance 200mg φαρμακούχα έμπλαστρα	DE/H/5067/001	022903	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen Durance 200mg φαρμακούχο έμπλαστρο	DE/H/5067/001	109570/19-09-2019	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió	not available	OGYI-T-6793/03	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	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
gyermekeknek				
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/12	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/101	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/102	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	HU
Nurofen eperízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/02	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	HU
Nurofen eperízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/002	OGYI-T-6793/116	RECKITT BENCKISER KFT	HU
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 Expres 200 mg Obalené tablety	not available	07/0062/09-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen Expres 400 mg Obalené tablety	not available	07/0063/09-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen Express 200 mg	not available	09-0056	RECKITT BENCKISER	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
apvalkotās tabletes			(POLAND) S.A.	
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
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 Express 200 mg capsule moi	DE/H/1482/001	7253/2014/01	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN EXPRESS 200 mg dengtos tabletės	not available	LT/1/97/0271/022	RECKITT BENCKISER (POLAND) S.A.	LT
NUROFEN EXPRESS 200 mg dengtos tabletės	not available	LT/1/97/0271/021	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Express 200 mg drajeuri	not available	12930/2020/01	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	12930/2020/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200 mg drajeuri	not available	12930/2020/09	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express 200mg Liquid Capsules	not available	PL 00063/0654	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI

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 200mg Liquid Capsules	not available	PL 00063/0648	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Express 200mg Tablets.	not available	PA 979/32/10	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Express 256 mg Caplets	not available	00063/0374	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Express 256 mg Tablets	not available	00063/0372	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Express 342mg Caplets	not available	PL 00063/0380	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Express 400 mg Liquid Capsules	not available	PL 00063/0653	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Express 684mg Caplets	not available	PL 00063/0384	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Express Femina, 200 mg, kapsułki, miękkie	DE/H/1482/001	16931	RECKITT BENCKISER (POLAND) S.A.	PL
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
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
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express Forte 400 mg capsule moi	NL/H/4313/001	11360/2019/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Express Forte, 400 mg, kapsułki, miękkie	NL/H/4313/001	19888	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Express Maximum Strength 400	not available	PA 979/32/11	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
mg Tablets.				
Nurofen Express Period Pain 200mg soft capsules	not available	PL 00063/0646	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Express, 200 mg kaetud tabletid	not available	743211	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Express, 200 mg, tabletki powlekane	not available	14446	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Extra Strength 400 mg Liquid Capsules	not available	PL 00063/0653	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Fastine Liquid Caps 200 mg, capsule, zacht	DE/H/1482/001	RVG 102120	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Fastine Liquid Caps 400 mg, capsule, zacht	NL/H/4313/001	RVG 105812	RECKITT BENCKISER HEALTHCARE B.V.	NL
NUROFEN FEBBRE E DOLORE 200 mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102451	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
NUROFEN FEBBRE E DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero	not available	034102386	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN FEBBRE E	not available	034102398	RECKITT BENCKISER	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
DOLORE 200mg/5ml sospensione orale gusto fragola senza zucchero			HEALTHCARE (ITALIA) SPA	
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 Bambini 100mg/5ml sospensione orale gusto arancia senza zucchero	not available	034102020	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
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 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

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 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
Nurofen Flextin 200 mg tablet ovaal, omhulde tabletten	not available	RVG 100659	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Flextin 200 mg tablet, omhulde tabletten	not available	RVG 100658	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Flextin 400 mg tablet, omhulde tabletten	not available	RVG 100657	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen for children	not available	MA0190/00402	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD	MT
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	XI
Nurofen for children 100 mg, μασώμενα καψάκια, μαλακά (Πορτοκάλι)	BE/H/0320/001	22496	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen for Children 200 mg/5 ml Orange Oral Suspension	not available	PL00063/0742	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen for Children 200 mg/5 ml Strawberry Oral Suspension	not available	PL 00063/0743	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen for Children 3	not available	PL 00063/0668	RECKITT BENCKISER	XI

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
months to 9 years Orange			(HEALTHCARE) UK LTD	
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
Nurofen for Children Meltlets 100mg Orodispersible Tablets	not available	PA 979/32/5	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN FOR CHILDREN oral suspension 100mg/5ml	not available	13609	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen for Children Orange	not available	PL 00063/0668	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen for Children Orange 100mg/5ml Oral Suspension	not available	PA 979/32/1	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for Children Orange 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2206/001	10-0476	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen for Children Orange Baby	not available	PL 00063/0668	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen for Children Orange Flavoured 100mg Chewable Capsule, Soft	not available	PL 00063/0723	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen For Children Orange Singles	not available	PL 00063/0669	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen for Children Sachets 100mg/5ml Oral Suspension	not available	PA 979/32/8	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for Children Six	DE/H/2343/001	PA 979/56/1	RECKITT BENCKISER	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
Plus Orange 200mg/5ml Oral Suspension			IRELAND LTD.	
Nurofen for Children Six Plus Strawberry 200mg/5ml Oral Suspension	DE/H/2343/002	PA 979/56/2	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for Children Strawberry	not available	PL 00063/0666	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen for Children Strawberry 100mg/5ml Oral Suspension	not available	PA 979/32/9	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen for Children Strawberry 200 mg/5 ml suspensija iekšķīgai lietošanai	DE/H/2206/002	10-0477	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen for Children Strawberry 3 Months to 12 years	not available	PL 00063/0666	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen for Children Strawberry Singles	not available	PL 00063/0670	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
NUROFEN FORTE 400 mg apvalkotās tabletes	not available	03-0422	RECKITT BENCKISER (POLAND) S.A.	LV
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 FORTE 400 mg dengtos tabletės	not available	LT/1/97/0271/001	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 400 mg dengtos tabletės	not available	LT/1/97/0271/002	RECKITT BENCKISER (POLAND) S.A.	LT
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 Forte 400 mg obložene tablete	not available	HR-H-934065674	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen Forte Express 400 mg apvalkotās tabletes	not available	08-0205	RECKITT BENCKISER (POLAND) S.A.	LV
NUROFEN FORTE EXPRESS 400 mg dengtos tabletės	not available	LT/1/97/0271/023	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Express, 400 mg kaetud tabletid	not available	743111	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/024	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Orange 40 mg/ml geriamoji suspensija	DE/H/2206/001	LT/1/97/0271/0130093421~30~BOT-PETAC-CRC5	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
Nurofen Forte Orange, 40 mg/ml, suukaudne suspensioon	DE/H/2206/001	707310	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Forte Strawberry 40 mg/ml	DE/H/2206/002	LT/1/97/0271/017-LT/1/97/0271/020/,	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
geriamoji suspensija		LT1/97/0271/25		
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
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/019	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Strawberry 40 mg/ml geriamoji suspensija	DE/H/2206/002	LT/1/97/0271/020	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Forte Strawberry, 40 mg/ml, suukaudne suspensioon	DE/H/2206/002	707210	RECKITT BENCKISER (POLAND) S.A.	EE
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
Nurofen Forte, 400 mg kaetud tabletid	not available	450704	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Forte, 400 mg, tabletki powlekane	not available	4937	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen für Kinder 100 mg, Weichkapseln zum Zerbeissen	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen für Kinder 200 mg überzogene Tabletten.	not available	BE189935	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
Nurofen für Kinder Zuckerfrei 4%	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM)	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Suspension zum Einnehmen			SA/NV	
Nurofen für Kinder zuckerfrei Rot 2% Suspension zum Einnehmen	not available	BE281644	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 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 Immedia 200 mg Weichkapseln	DE/H/1482/001	71862.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Immedia 400 mg Filmtabletten	not available	43917.01.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/05	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Immedia Ultra 400 mg drajeuri	not available	12516/2019/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Joint & Back Pain Relief Max Strength 10 % Gel	not available	PL 10972/0089	MERCURY PHARMA GROUP LTD	XI
Nurofen Joint & Muscular Pain Relief 200mg Medicated Plaster	not available	PL 00063/0733	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI

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 Joint & Muscular Pain Relief 200mg Medicated Plaster	DE/H/5067/001	PL 00063/0733	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Jordgubb, 40 mg/ml, oral suspension	DE/H/2204/002	41890	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	SE
Nurofen Junior 100 mg cápsulas blandas masticables	DE/H/6014/001	82519	RECKITT BENCKISER HEALTHCARE S.A.	ES
Nurofen Junior 100 mg cápsulas blandas masticables	DE/H/6014/001	82519	RECKITT BENCKISER HEALTHCARE S.A.	ES
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 Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/01	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/02	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/04	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/05	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	14118/2021/06	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/07	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/08	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/09	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/10	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/11	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/12	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/13	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/14	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	14118/2021/15	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen Junior 7 ani+ cu aromă de portocale 100 mg capsule moi masticabile	NL/H/4314/001	14118/2021/16	RECKITT BENCKISER (ROMANIA) SRL	RO
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 Junior Appelsín 40 mg/ml mixtúra, dreifa	DE/H/2204/001	IS/1/11/118/01	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	IS
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 Junior cu aromă de portocale 40 mg/ml suspensie orală	DE/H/2343/001	9338/2016/01	RECKITT BENCKISER (ROMANIA) SRL	RO
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/H/2343/001	9338/2016/03	RECKITT BENCKISER	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
de portocale 40 mg/ml suspensie orală			(ROMANIA) SRL	
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 Junior Fieber- und Schmerzsaft Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2204/002	76552.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
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 Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2204/001	76551.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
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 Junior Fieber- und Schmerzsaft Orange 40 mg/ml Suspension zum Einnehmen	DE/H/2206/001	76553.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen Junior Fiebersaft Erdbeer 20 mg/ml Suspension zum Einnehmen	not available	62150.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
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 Junior Jarðarber 40 mg/ml mixtúra, dreifa	DE/H/2204/002	IS/1/11/119/01	RECKITT BENCKISER HEALTHCARE (SCANDINAVIA) A/S	IS
Nurofen Junior Kaudragee Orange	DE/H/6014/001	95058.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
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
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	PL/H/0602/001	OGYI-T-6793/130	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
narancsízű 100 mg lágy rágókapszula				
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
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
Nurofen Laranja 40 mg/ml suspensão oral	DE/H/2343/001	5630033	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Long Lasting 300mg Prolonged Release Hard Capsules	not available	PA 979/32/15	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Long Lasting Pain Relief 300mg Prolonged Release Capsules	not available	PL 00063/0378	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Lysine 200 mg Filmtabletten	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	not available	2010080859	RECKITT BENCKISER	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
mg, comprimés pelliculés			HEALTHCARE (BELGIUM) SA/NV	
NUROFEN LYSINE 200 mg, filmomhulde tabletten	not available	BE239005	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen Max Strength Pain Relief 512mg tablets	not available	00063/0412	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Maximum Strength Migraine Pain 684mg Caplets	not available	PL 00063/0384	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Meltlets Lemon	not available	PL 00063/0382	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Miesnie i Stawy Forte, 400 mg, tabletki powlekane	not available	14316	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Mięśnie i Stawy, 200 mg, plaster leczniczy	DE/H/5067/001	25344	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Migraine Pain	not available	PL 00063/0380	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
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
Nurofen Morango 40 mg/ml suspensão oral	DE/H/2343/002	5630157	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Musc 200 mg emplastro medicamentoso	DE/H/5067/001	5760723	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Musc 200 mg emplastro medicamentoso	DE/H/5067/001	5760731	RECKITT BENCKISER HEALTHCARE, LDA.	PT
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

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 Musc 400 mg comprimidos revestidos	not available	5296728	RECKITT BENCKISER HEALTHCARE, LDA.	PT
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 narancsízű 20 mg/ml belsőleges szuszpenzió gyermekeknek	not available	OGYI-T-6793/01	RECKITT BENCKISER KFT	HU
Nurofen narancsízű 40 mg/ml belsőleges szuszpenzió gyermekeknek	DE/H/2343/001	OGYI-T-6793/115	RECKITT BENCKISER KFT	HU
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 NEO 200 MG obalené tablety	not available	07/812/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
NUROFEN NEO FEMINA 400 mg obalené tablety	not available	07/813/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen Non-Aqua 100 mg szájban diszpergálódó tabletta gyermekeknek	not available	OGYI-T-6793/79	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 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 Non-Aqua 200 mg szájban diszpergálódó tabletta	not available	OGYI-T-6793/84	RECKITT BENCKISER KFT	HU
Nurofen Omhulde Tabletten 200, omhulde tabletten 200 mg	not available	RVG 23518	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen Orange 100 mg kramtomosios minkštosios kapsulės	PL/H/0602/001	LT/1/16/4016/001-LT/1/16/4016/016	RECKITT BENCKISER (POLAND) S.A.	LT
Nurofen Orange 100 mg mīkstās košļājamās kapsulas	PL/H/0602/001	17-0025	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Orange, 100 mg pehmed närimiskapslid	PL/H/0602/001	925516	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen Pain Relief 256mg Tablets	not available	00063/0373	RECKITT BENCKISER HEALTHCARE (UK) LTD	XI
Nurofen Pain Relief Caplets 256mg Tablets	not available	00063/0411	RECKITT BENCKISER (HEALTHCARE) UK LTD	XI
Nurofen Patch 200 mg emplâtre médicamenteux	DE/H/5067/001	BE500080	RECKITT BENCKISER HEALTHCARE (BELGIUM)	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			SA/NV	
Nurofen Patch 200 mg Emplâtre médicamenteux	DE/H/5067/001	2017040114	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen Patch 200 mg pleister	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 pediátrico 20 mg/ml suspension oral sabor fresa	not available	82.998	RECKITT BENCKISER HEALTHCARE S.A.	ES
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
NUROFEN PENTRU COPII cu aromă de portocale 100 mg/5 ml suspensie orală	not available	9006/2016/02	RECKITT BENCKISER (ROMANIA) SRL	RO
NUROFEN PENTRU COPII cu aromă de portocale 100 mg/5 ml suspensie orală	not available	9006/2016/03	RECKITT BENCKISER (ROMANIA) SRL	RO
Nurofen pleister 200 mg	DE/H/5067/001	RVG 123099	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen pleister 200 mg	DE/H/5067/001	RVG 123099	RECKITT BENCKISER HEALTHCARE B.V.	NL
NUROFEN POUR ENFANTS 200 mg comprimés enrobés.	not available	BE189935	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR	not available	0209/10040753	RECKITT BENCKISER	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
ENFANTS 200 mg comprimés enrobés.			HEALTHCARE (BELGIUM) SA/NV	
NUROFEN POUR ENFANTS SANS SUCRE 2% suspension buvable	not available	BE191021	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR ENFANTS SANS SUCRE 2% suspension buvable	not available	0209/10040751	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen pour Enfants Sans sucre 4% suspension buvable	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pour Enfants Sans sucre 4% suspension buvable	DE/H/2343/001	2010120025	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
NUROFEN POUR ENFANTS SANS SUCRE rouge 2% suspension buvable	not available	BE281644	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
NUROFEN POUR ENFANTS SANS SUCRE rouge 2% suspension buvable	not available	0209/10040752	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen pour Enfants Sans sucre rouge 4% suspension buvable	DE/H/2343/002	BE378831	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pour Enfants Sans sucre rouge 4% suspension buvable	DE/H/2343/002	2010120026	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen pour Enfants100 mg, capsules molles à mâcher	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen pour Enfants100 mg, capsules molles à mâcher	BE/H/0320/001	2017070240	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	LU
Nurofen pre deti 4 % jahoda	DE/H/2343/002	07/0004/11-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL.	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
perorálna suspenzia			S.R.O	
Nurofen pre deti 4 % pomaranc perorálna suspenzia	DE/H/2343/001	07/0003/11-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
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 pro děti 4 % pomeranč perorální suspenze	DE/H/2343/001	07/968/10-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen Rapid 200 mg Capsules	DE/H/1482/001	07/025/11-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
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 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 Rapid 200 mg meke kapsule	not available	HR-H-821340280	RECKITT BENCKISER (CROATIA) D.O.O.	HR
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

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 400 mg - Filmtabletten	not available	1-23773	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen Rapid 400 mg cápsulas blandas	not available	68.517	RECKITT BENCKISER HEALTHCARE S.A.	ES
NUROFEN Rapid 400 mg Capsules mäkké kapsuly	not available	29/0302/08-S	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	SK
Nurofen rapid 400 mg Filmtabletten	not available	1-23773	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
NUROFEN RAPID 400 mg měkké tobolky	not available	29/158/08-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
Nurofen rapid 400 mg Weichkapseln	NL/H/4313/001	1-31144	RECKITT BENCKISER DEUTSCHLAND GMBH	AT
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001	OGYI-T-6793/110	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001	OGYI-T-6793/111	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001	OGYI-T-6793/112	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001	OGYI-T-6793/113	RECKITT BENCKISER KFT	HU
Nurofen Rapid Forte 400 mg lágy kapszula	NL/H/4313/001	OGYI-T-6793/114	RECKITT BENCKISER KFT	HU
Nurofen Rapid Relief Maximum Strength 400mg Liquid Capsules	not available	PA 979/32/13	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Schmerz- und Fiebersaft Erdbeer 40 mg/ml Suspension zum Einnehmen	DE/H/2343/002	77581.00.00	RECKITT BENCKISER DEUTSCHLAND GMBH	DE
Nurofen voor Kinderen 100 mg, zachte kauwcapsules	BE/H/0320/001	BE503022	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen voor Kinderen Aardbei suspensie,	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
suspensie 200mg/5ml				
Nurofen voor Kinderen Sinaasappel 100 mg, kauwcapsules, zacht	NL/H/4314/001	RVG 112524	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen voor Kinderen Sinaasappel suspensie, suspensie 200mg/5ml	DE/H/2343/001	RVG 104647	RECKITT BENCKISER HEALTHCARE B.V.	NL
Nurofen voor Kinderen Suikervrij 4% suspensie voor oraal gebruik	DE/H/2343/001	BE378822	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen voor Kinderen Suikervrij rood 4% suspensie voor oraal gebruik	DE/H/2343/002	BE378831	RECKITT BENCKISER HEALTHCARE (BELGIUM) SA/NV	BE
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001	5437330	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001	5437348	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001	5437355	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen Xpress 400 mg Cápsulas moles	NL/H/4313/001	5437363	RECKITT BENCKISER HEALTHCARE, LDA.	PT
Nurofen za djecu 100mg/5 ml oralna suspenzija	not available	HR-H-846940596	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (φράουλα)	DE/H/2343/002	21713	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (φράουλα)	DE/H/2343/002	27900/20-4-12	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (πορτοκάλι)	DE/H/2343/001	21712	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα	DE/H/2343/001	021712	RECKITT BENCKISER HELLAS HEALTHCARE SA	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
(πορτοκάλι)				
Nurofen για Παιδιά 4 % Πόσιμο Εναιώρημα (φράουλα)	DE/H/2343/002	021713	RECKITT BENCKISER HELLAS HEALTHCARE SA	CY
Nurofen, 200 mg kaetud tabletid	not available	218498	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen, 200 mg ravimplaastrid	DE/H/5067/001	987219	RECKITT BENCKISER (POLAND) S.A.	EE
Nurofen, 200 mg, tabletki powlekane	not available	R/7068	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen, 50 mg/g, żel	not available	10062	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860014	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860026	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860038	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860040	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860053	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860115	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860065	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860077	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860089	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860127	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg Capsule Molli	NL/H/4313/001	041860091	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofencaps 400 mg	NL/H/4313/001	041860103	RECKITT BENCKISER	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
Capsule Molli			HEALTHCARE (ITALIA) SPA	
NUROFENCAPS 400 mg, capsule molle	NL/H/4313/001	NL38369	RECKITT BENCKISER HEALTHCARE FRANCE	FR
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 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 221 335 5 3	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 221 337 8 2	RECKITT BENCKISER HEALTHCARE FRANCE	FR
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 221 339 0 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247056	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247070	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247082	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
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247106	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247132	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247120	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247068	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247094	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247157	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247169	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247031	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247144	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247029	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247017	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247043	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
Nurofenkid Febbre e Dolore 100 mg, capsule molli masticabili	DE/H/6014/001	044247118	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFENPLAST 200 mg, emplâtre médicamenteux	DE/H/5067/001	34009 301 708 5 4	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFENPLAST 200 mg, emplâtre médicamenteux	DE/H/5067/001	34009 301 708 6 1	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofenplast 200mg, emplâtre médicamenteux	DE/H/5067/001	NL45945	RECKITT BENCKISER HEALTHCARE FRANCE	FR
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/1/19/4400/004	RECKITT BENCKISER (POLAND) S.A.	LT
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
Pathofen 100 mg, zachte kauwcapsules	NL/H/4843/001	RVG 123962	PATHEON SOFTGELS 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
Perdofemina 400 mg, comprimés pelliculés	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, comprimés pelliculés	not available	BE228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
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
Perdofemina 400 mg, filmomhulde tabletten	not available	BE228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, filmomhulde tabletten	not available	BE228453	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdofemina 400 mg, filmomhulde tabletten	not available	BE228453	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, filmomhulde tabletten	not available	BE 228453	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, comprimés pelliculés	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
Perdophen 200 mg, filmomhulde tabletten	not available	BE196637	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Perdophen 200 mg,	not available	BE196637	JOHNSON & JOHNSON	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
filmomhulde tabletten			CONSUMER N.V./S.A.	
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, comprimés pelliculés	not available	BE226904	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
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
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, 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, 200 mg Filmtabletten	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 Filmtabletten	not available	BE196637	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
Peribu 400 mg solución para perfusión.	PT/H/1732/001	84556	ALTAN PHARMACEUTICALS S.A.	ES
Phorpain 5 %w/w Gel	not available	PL 10972/0091	MERCURY PHARMA GROUP LTD	XI

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
Phorpain gel maximum strength	not available	PL 10972/0082	MERCURY PHARMA GROUP LTD	XI
Phorpain maximum strength 10% w/w Gel	not available	PL 10972/0090	MERCURY PHARMA GROUP LTD	XI
Poindol 200 mg granulado para solución oral	not available	76907	FARMALIDER, S.A.	ES
proff Schmerzcreme	not available	1-20032	DOLORGIET GMBH & CO KG	AT
proff Schmerzcreme	not available	0473/85/03/0258	DOLORGIET GMBH & CO KG	LU
proff Schmerzgel	not available	1192.00.03	DOLORGIET GMBH & CO KG	DE
Proff® Schmerzcreme	not available	1192.00.02	DOLORGIET GMBH & CO KG	DE
Proff® Schmerzcreme	not available	1192.00.02	DOLORGIET GMBH & CO KG	DE
Proff® Schmerzcreme	not available	1192.00.02	DOLORGIET GMBH & CO KG	DE
ratioDolor Ibuprofen 400 mg Filmtabletten	DE/H/2590/001	1-30782	TEVA B.V	AT
Sainsbury's Healthcare Long Lasting Pain Relief 200mg Modified Release Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Solibu 400 mg Infusionslösung	PT/H/1732/001	98473.00.00	ALTAN PHARMA LTD	DE
Solibu 400 mg Solução para perfusão	PT/H/1732/001	5747266	ALTAN PHARMACEUTICALS S.A.	PT
Solibu 400 mg Solução para perfusão	PT/H/1732/001	5747258	ALTAN PHARMACEUTICALS S.A.	PT
Solibu 600 mg Infusionslösung	PT/H/1732/002	2201636.00.00	ALTAN PHARMA LTD	DE
Solibu 600 mg Solução para perfusão	PT/H/1732/002	5747274	ALTAN PHARMACEUTICALS S.A.	PT
Solibu 600 mg Solução para perfusão	PT/H/1732/002	5747308	ALTAN PHARMACEUTICALS S.A.	PT
Solibu 600 mg solución para perfusión.	PT/H/1732/002	83680	ALTAN PHARMACEUTICALS S.A.	ES
Spalt Forte 400 mg Weichkapseln	not available	9032.01.00	PHARMASGP 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	PHARMASGP GMBH	DE
Spalt Migräne 400 mg Weichkapseln	not available	12732.00.00	PHARMASGP GMBH	DE
Spalt Mobil 400 mg Weichkapseln	not available	9632.01.00	PHARMASGP GMBH	DE
SPEDIFEN 200 mg, comprimé	not available	34009 360 811 0 9	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 812 7 7	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 813 3 8	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 815 6 7	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 811 0 9	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 812 7 7	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 813 3 8	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, comprimé	not available	34009 360 815 6 7	ZAMBON FRANCE S.A.	FR
SPEDIFEN 200 mg, suspension buvable en sachet, édulcorée à la saccharine sodique et au maltitol liquide	not available	34009 302 080 1 4	ZAMBON FRANCE S.A.	FR
Spedifen 400 mg filmtabletta	not available	OGYI-T-7162/05	ZAMBON S.P.A.	HU
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
SPEDIFEN 400 mg,	not available	34009 300 210 3 3	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
comprimé pelliculé				
SPEDIFEN 400 mg, comprimé pelliculé	not available	34009 362 516 6 3	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	34009 362 517 2 4	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	34009 362 518 9 2	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, comprimé pelliculé	not available	34009 382 851 5 4	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 510 5 3	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 510 5 3	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, suspension buvable en sachet, édulcorée à la saccharine sodique, au maltitol liquide et à la thaumatine	not available	34009 302 080 2 1	ZAMBON FRANCE S.A.	FR
SPEDIFEN 400 mg, suspension buvable en sachet, édulcorée à la saccharine sodique, au maltitol liquide et à la thaumatine	not available	34009 302 080 2 1	ZAMBON FRANCE S.A.	FR
Spedifen 400, 400 mg, tabletki powlekane	not available	10868	ZAMBON S.P.A.	PL
Spedifen Rapid 400 mg granulátum	not available	OGYI-T-7162/01	ZAMBON S.P.A.	HU
Spedifen Rapid 400 mg granulátum	not available	OGYI-T-7162/02	ZAMBON S.P.A.	HU
Spedifen Rapid Forte 600 mg granulátum	not available	OGYI-T-7162/03	ZAMBON S.P.A.	HU
Spedifen Rapid Forte 600	not available	OGYI-T-7162/04	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
mg granulátum				
Spedifen Rapid Forte 600 mg granulátum	not available	OGYI-T-7162/03	ZAMBON S.P.A.	HU
Spedifen Rapid Forte 600 mg granulátum	not available	OGYI-T-7162/04	ZAMBON S.P.A.	HU
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
SPIDIDOL 400 mg compresse rivestite con film	not available	039600010	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
SPIDIDOL 400 mg granulato per soluzione orale gusto albicocca	not available	039600022	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL 400 mg granulato per soluzione orale gusto cola-limone	not available	039600034	ZAMBON ITALIA S.R.L.	IT
Spididol analgesico 200 mg compresse	not available	028710034	ZAMBON ITALIA S.R.L.	IT
Spididol analgesico 200 mg compresse effervescenti	not available	028710010	ZAMBON ITALIA S.R.L.	IT
SPIDIDOL ANALGESICO 200 mg granulato per	not available	028710022	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
soluzione orale				
SPIDIDOL ANALGESICO 200 mg/ml Gocce orali soluzione	not available	028710046	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457063	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457051	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457012	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457024	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457048	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 200 mg sospensione orale in bustina	HU/H/0828/001	044457036	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 400 mg sospensione orale in bustina	HU/H/0828/002	044457113	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 400 mg sospensione orale in bustina	HU/H/0828/002	044457125	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 400 mg sospensione orale in bustina	HU/H/0828/002	044457075	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 400 mg sospensione orale in bustina	HU/H/0828/002	044457087	ZAMBON ITALIA S.R.L.	IT
SPIDIDOLPOCKET 400 mg sospensione orale in	HU/H/0828/002	044457099	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
bustina				
SPIDIDOLPOCKET 400 mg sospensione orale in bustina	HU/H/0828/002	044457101	ZAMBON ITALIA S.R.L.	IT
Spidifen 200 mg comprimés	not available	BE386014	ZAMBON NV	BE
Spidifen 200 mg comprimés	not available	2011051165	ZAMBON NV	LU
Spidifen 200 mg granulaat voor drank	not available	BE157735	ZAMBON NV	BE
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 200 mg Granulat zur Herstellung einer Lösung zum Einnehmen	not available	BE157735	ZAMBON NV	BE
Spidifen 200 mg Granulat zur Herstellung einer Lösung zum Einnehmen	not available	2011051163	ZAMBON NV	LU
Spidifen 200 mg granulés pour solution buvable	not available	BE157735	ZAMBON NV	BE
Spidifen 200 mg granulés pour solution buvable	not available	2011051163	ZAMBON NV	LU
Spidifen 200 mg tablete	not available	HR-H-567134731	ZAMBON S.P.A.	HR
Spidifen 200 mg Tabletten	not available	BE386014	ZAMBON NV	BE
Spidifen 200 mg Tabletten	not available	BE386014	ZAMBON NV	BE
Spidifen 200 mg	not available	2011051165	ZAMBON 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
Tabletten				
SPIDIFEN 400 mg Compresse rivestite con film	not available	026916080	ZAMBON ITALIA S.R.L.	IT
Spidifen 400 mg comprimés pelliculés	NL/H/0341/001	BE244456	ZAMBON NV	BE
Spidifen 400 mg comprimés pelliculés	NL/H/0341/001	2011091290	ZAMBON NV	LU
Spidifen 400 mg filmom oblozene tablete	not available	HR-H-287190928	ZAMBON S.P.A.	HR
Spidifen 400 mg filmomhulde tabletten	NL/H/0341/001	BE244456	ZAMBON NV	BE
SPIDIFEN 400 mg Filmtabletten	NL/H/0341/001	BE244456	ZAMBON NV	BE
Spidifen 400 mg granulaat voor drank	not available	BE157805	ZAMBON NV	BE
Spidifen 400 mg Granulat zur Herstellung einer Lösung zum Einnehmen	not available	BE157805	ZAMBON NV	BE
SPIDIFEN 400 mg Granulato per soluzione orale aroma albicocca	not available	026916104	ZAMBON ITALIA S.R.L.	IT
Spidifen 400 mg granulés pour solution buvable	not available	BE157805	ZAMBON NV	BE
Spidifen 600 mg granulaat voor drank	not available	BE478853	ZAMBON NV	BE
SPIDIFEN 600 mg granulado para solução oral	not available	5912886	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 600 mg granulado para solução oral	not available	4618286	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN 600 mg granulado para solução	not available	4618385	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
oral				
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
Spidifen 600 mg Granulat zur Herstellung einer Lösung zum Einnehmen	not available	BE478853	ZAMBON NV	BE
Spidifen 600 mg Granulat zur Herstellung einer Lösung zum Einnehmen	not available	2016040085	ZAMBON NV	LU
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
SPIDIFEN 600 mg Granulato per soluzione orale aroma cola-limone	not available	026916167	ZAMBON ITALIA S.R.L.	IT
SPIDIFEN 600 mg granulato per soluzione orale aroma cola-limone	not available	026916142	ZAMBON ITALIA S.R.L.	IT
SPIDIFEN 600 mg Granulato per soluzione orale aroma menta-anice	not available	026916155	ZAMBON ITALIA S.R.L.	IT
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 600 mg granulés pour solution buvable	not available	2016040085	ZAMBON NV	LU
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 EF 400 mg granulado para solução oral	not available	4617981	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN EF 400 mg granulado para solução oral	not available	5912787	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
SPIDIFEN EF 400 mg granulado para solução oral	not available	4618088	ZAMBON - PRODUTOS FARMACÊUTICOS, LDA.	PT
Spidifen® 400 tablet, filmomhulde tabletten 400 mg	NL/H/0341/001	RVG 25385	ZAMBON NEDERLAND B.V.	NL
SPIFEN 200 mg, comprimé	not available	34009 360 816 2 8	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 817 9 6	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 818 5 7	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 819 1 8	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 816 2 8	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 817 9 6	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	34009 360 818 5 7	ZAMBON FRANCE S.A.	FR
SPIFEN 200 mg, comprimé	not available	34009 360 819 1 8	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 508 3 3	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 510 8 3	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 511 4 4	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 512 0 5	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 513 7 3	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, comprimé pelliculé	not available	34009 362 514 3 4	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 508 6 1	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 509 2 2	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 510 0 4	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 511 7 2	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 349 691 2 6	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 508 6 1	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution	not available	34009 347 509 2 2	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
buvable en sachet-dose				
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 510 0 4	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 347 511 7 2	ZAMBON FRANCE S.A.	FR
SPIFEN 400 mg, granulés pour solution buvable en sachet-dose	not available	34009 349 691 2 6	ZAMBON FRANCE S.A.	FR
Superdrug Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Tesco Health Ibuprofen 200mg Tablets	not available	PL 16028/0161	GALPHARM HEALTHCARE LIMITED	XI
Tesco Health Ibuprofen 400mg Tablets	not available	PL 16028/0040	GALPHARM HEALTHCARE LIMITED	XI
Tesco Health Ibuprofen 400mg Tablets	not available	PL 16028/0040	GALPHARM HEALTHCARE LIMITED	XI
Tesco Health Long Lasting Pain Relief, 200mg Modified Release Capsules, Hard	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Teva Ibuprofen 200mg Long lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
The-Cooperative Ibuprofen 200mg Long Lasting Capsules	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	OPELLA HEALTHCARE AUSTRIA GMBH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	OPELLA HEALTHCARE AUSTRIA GMBH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	OPELLA HEALTHCARE AUSTRIA GMBH	AT
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	OPELLA HEALTHCARE AUSTRIA GMBH	AT
ThomaDuo 400 mg/100	DE/H/4270/001	137318	OPELLA HEALTHCARE	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
mg Filmtabletten			AUSTRIA GMBH	
ThomaDuo 400 mg/100 mg Filmtabletten	DE/H/4270/001	137318	OPELLA HEALTHCARE AUSTRIA GMBH	AT
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
Thomapyrin TENSION DUO 400 mg/100 mg Filmtabletten	DE/H/4270/001	93770.00.00	A. NATTERMANN & CIE. GMBH	DE
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
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
Wilko Long Lasting Pain Relief	not available	PL 16028/0120	GALPHARM HEALTHCARE LIMITED	XI
Winadol 200 mg	HU/H/0828/001	OGYI-T-23323/01	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
belsőleges szuszpenzió tasakban				
Winadol 200 mg belsőleges szuszpenzió tasakban	HU/H/0828/001	OGYI-T-23323/02	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 200 mg belsőleges szuszpenzió tasakban	HU/H/0828/001	OGYI-T-23323/03	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 400 mg belsoleges szuszpenzió tasakban	HU/H/0828/002	OGYI-T-23323/04	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 400 mg belsoleges szuszpenzió tasakban	HU/H/0828/002	OGYI-T-23323/05	NUTRA ESSENTIAL OTC, S.L.	HU
Winadol 400 mg belsoleges szuszpenzió tasakban	HU/H/0828/002	OGYI-T-23323/06	NUTRA ESSENTIAL OTC, S.L.	HU
Zenalgin® 800 mg Filmtabletten	SE/H/2058/003	140636	ZENTIVA, K.S.	AT
Zentiben 800 mg film-coated tablets	PT/H/2719/003/DC	TBD	BILLEV FARMACIJA VZHOD D.O.O.	PT
Zentidol 800 mg film-coated tablets	PT/H/2609/003/DC	TBD	BILLEV FARMACIJA VZHOD D.O.O.	PT
Nurofen for children 100 mg, μασώμενα καψάκια, μαλακά (Πορτοκάλι)	BE/H/0320/001	35066/15-4-2019	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
Nurofen for children 100 mg, μασώμενα καψάκια, μαλακά Πορτοκάλι	BE/H/0320/001	35066/28-3-2019	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
БРУФЕН РЕТАРД 800 mg таблетки с удължено освобождаване	SE/H/0912/005	20110338	MYLAN EOOD	BG
ИБУПРОМ РЕГУЛАР 200 mg филмирани таблетки	PL/H/0709/002	20230072	US PHARMACIA SP. Z O.O.	BG
ИБУПРОМ РЕГУЛАР 200	PL/H/0709/002	20230072	US PHARMACIA SP. Z O.O.	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
mg филмирани таблетки				
Ибупрофен Б. Браун 400 mg/100 ml инфузионен разтвор	ES/H/0390/001	20170169	B.BRAUN MELSUNGEN AG	BG
Ибупрофен Б. Браун 600 mg/100 ml инфузионен разтвор	ES/H/0392/001/DC	20170231	B.BRAUN MELSUNGEN AG	BG
Ибупрофен Нутра Есеншъл 100 mg перорална суспензия в саше	SI/H/0214/001/DC	20170314	NUTRA ESSENTIAL OTC, S.L.	BG
Ибупрофен Нутра Есеншъл 100 mg перорална суспензия в саше	SI/H/0214/002/DC	20170315	NUTRA ESSENTIAL OTC, S.L.	BG
Ибупрофен Нутра Есеншъл 400 mg перорална суспензия в саше	SI/H/0214/003	20170313	NUTRA ESSENTIAL OTC, S.L.	BG
МИГ джуниър 40 mg/ml перорална суспензия	DE/H/2598/002	20150332	BERLIN-CHEMIE AG	BG
МИГ за деца 20 mg/ml перорална суспензия	DE/H/2463/001	20110424	BERLIN-CHEMIE AG	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	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
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Ношпалгин 400 mg/ 100 mg филмирани таблетки	DE/H/5544/001/DC	20200090	OPELLA HEALTHCARE FRANCE SAS	BG
Нурофен 200 mg лечебен пластир	DE/H/5067/001	20180337	RECKITT BENCKISER (ROMANIA) SRL	BG
НУРОФЕН ЕКСПРЕС 200 mg меки капсули	DE/H/1482/001	20100224	RECKITT BENCKISER (ROMANIA) SRL	BG
Нурофен Експрес Форте 400 mg капсули, меки Nurofen Express Forte 400 mg capsules, soft	NL/H/4313/001	20120227	RECKITT BENCKISER (ROMANIA) SRL	BG
Нурофен за Юноши Ягода 200 mg/5 ml перорална суспензия	DE/H/2343/002	20150019	RECKITT BENCKISER (ROMANIA) SRL	BG
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
Нурофен за Юноши Портокал 100 mg меки капсули за дъвчене	PL/H/0602/001	20170065	RECKITT BENCKISER (ROMANIA) SRL	BG