9 February 2017
EMA/89247/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ibuprofen / pseudoephedrine

Procedure no.: PSUSA/00001711/201607

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Larofen Plus 200mg/30mg, comprimate filmate	not available	119/2007/01	LAROPHARM SRL	RO
MODAFEN 200 MG/30 MG COMPRIMATE FILMATE	not available	480/2007/02	ZENTIVA, K.S.	RO
MODAFEN 200 MG/30 MG COMPRIMATE FILMATE	not available	480/2007/03	ZENTIVA, K.S.	RO
MODAFEN 200 MG/30 MG COMPRIMATE FILMATE	not available	480/2007/04	ZENTIVA, K.S.	RO
Modafen 200 mg/30 mg comprimate filmate	not available	480/2007/01	ZENTIVA, K.S.	RO
Olytabs 200 mg/30 mg Filmtabletten	DE/H/4183/001/DC	93268.00.00	JOHNSON & JOHNSON GMBH	DE
Olytabs 200 mg/30 mg Filmtabletten	DE/H/4183/001/DC	93268.00.00	JOHNSON & JOHNSON GMBH	DE
Olytabs 200 mg/30 mg Filmtabletten	DE/H/4183/001/DC	93268.00.00	JOHNSON & JOHNSON GMBH	DE
Olytabs 200 mg/30 mg Filmtabletten	DE/H/4183/001/DC	93268.00.00	JOHNSON & JOHNSON GMBH	DE
Sudafed Extra, 200 mg + 30 mg, tabletki powlekane	DE/H/4183/001/DC	23235	MCNEIL PRODUCTS LIMITED	PL
Sudafed Extra, 200 mg + 30 mg, tabletki powlekane	DE/H/4183/001/DC	23235	MCNEIL PRODUCTS LIMITED	PL
Sudafed Extra, 200 mg + 30 mg, tabletki powlekane	DE/H/4183/001/DC	23235	MCNEIL PRODUCTS LIMITED	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
Sudafed Extra, 200 mg + 30 mg, tabletki powlekane	DE/H/4183/001/DC	23235	MCNEIL PRODUCTS LIMITED	PL
Олитабс 200 mg/30 mg филмирани таблетки	DE/H/4183/001	20160014	MCNEIL PRODUCTS LIMITED	BG
Olytabs 200 mg/30 mg comprimate filmate	DE/H/4183/001	8529/2016/03	MCNEIL PRODUCTS LIMITED	RO
Olytabs 200 mg/30 mg comprimate filmate	DE/H/4183/001	8529/2016/01	MCNEIL PRODUCTS LIMITED	RO
Olytabs 200 mg/30 mg comprimate filmate	DE/H/4183/001	8529/2016/04	MCNEIL PRODUCTS LIMITED	RO
Olytabs 200 mg/30 mg comprimate filmate	DE/H/4183/001	8529/2016/02	MCNEIL PRODUCTS LIMITED	RO
Olytabs 200 mg/30 mg filmsko obložene tablete	DE/H/4183/001	H/16/02163/004	MCNEIL PRODUCTS LIMITED	SI
Olytabs 200 mg/30 mg filmom obložene tablete	DE/H/4183/001	HR-H-962261553	JOHNSON & JOHNSON S.E.D.O.O.	HR
Olytabs 200 mg/30 mg filmom obložene tablete	DE/H/4183/001	HR-H-962261553	JOHNSON & JOHNSON S.E.D.O.O.	HR
Olytabs 200 mg/30 mg filmom obložene tablete	DE/H/4183/001	HR-H-962261553	JOHNSON & JOHNSON S.E.D.O.O.	HR
Olytabs 200 mg/30 mg filmsko obložene tablete	DE/H/4183/001	H/16/02163/002	MCNEIL PRODUCTS LIMITED	SI
Olytabs 200 mg/30 mg filmom obložene tablete	DE/H/4183/001	HR-H-962261553	JOHNSON & JOHNSON S.E.D.O.O.	HR
Olytabs 200 mg/30 mg filmsko obložene tablete	DE/H/4183/001	H/16/02163/001	MCNEIL PRODUCTS LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Olytabs 200 mg/30 mg filmsko obložene tablete	DE/H/4183/001	H/16/02163/003	MCNEIL PRODUCTS LIMITED	SI
Олитабс 200 mg/30 mg филмирани таблетки	DE/H/4183/001	20160014	MCNEIL PRODUCTS LIMITED	BG
Олитабс 200 mg/30 mg филмирани таблетки	DE/H/4183/001	20160014	MCNEIL PRODUCTS LIMITED	BG
Олитабс 200 mg/30 mg филмирани таблетки	DE/H/4183/001	20160014	MCNEIL PRODUCTS LIMITED	BG
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg filmomhulde tabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg Filmtabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg comprimés pelliculés	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg filmomhulde tabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg comprimés pelliculés	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg Filmtabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg comprimés pelliculés	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg filmomhulde tabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg Filmtabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg Filmtabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg comprimés pelliculés	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Ibuprofen/Pseudoephedrine Hydrochloride Diapharm 200 mg/30 mg filmomhulde tabletten	DE/H/4183/001	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	DE/H/4183/001/DC	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	DE/H/4183/001/DC	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	DE/H/4183/001/DC	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg/30mg Film-coated Tablets	DE/H/4183/001	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg/30mg Film-coated Tablets	DE/H/4183/001	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	DE/H/4183/001	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	DE/H/4183/001	PA 823/68/1	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Metafen ZATOKI, 200 mg + 30 mg, tabletki	not available	11005	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
MODAFEN filmom obalené tablety	not available	07/0263/01-S	ZENTIVA, K.S.	SK
MODAFEN filmom obalené tablety	not available	07/0263/01-S	ZENTIVA, K.S.	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
MODAFEN filmom obalené tablety	not available	07/0263/01-S	ZENTIVA, K.S.	SK
Acatar Zatoki, 200 mg + 30 mg, tabletki powlekane	not available	11339	US PHARMACIA SP. Z O.O.	PL
IBUPROM ZATOKI, 200 mg + 30 mg, tabletki powlekane	not available	9650	US PHARMACIA SP. Z O.O.	PL
Laboratoria PolfaŁódź ZATOKI, 200 mg + 30 mg, tabletki	not available	21816	BIO PROFIL POLSKA SP Z OO	PL
EXTRALGIN COLD 200 mg/30 mg film-coated tablets Ibuprofen/Pseudoephedrine hydrochloride	not available	20100639	ZENTIVA, K.S.	BG
EXTRALGIN COLD 200 mg/30 mg film-coated tablets Ibuprofen/Pseudoephedrine hydrochloride	not available	20100639	ZENTIVA, K.S.	BG
EXTRALGIN COLD 200 mg/30 mg film-coated tablets Ibuprofen/Pseudoephedrine hydrochloride	not available	20100639	ZENTIVA, K.S.	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
EXTRALGIN COLD 200 mg/30 mg film-coated tablets Ibuprofen/Pseudoephedrine hydrochloride	not available	20100639	ZENTIVA, K.S.	BG
Ибупром Синус 200 mg/30 mg обвити таблетки	not available	20080222	US PHARMACIA SP. Z O.O.	BG
IbuSinex 200 mg/30 mg Filmtabletten	DE/H/4187/001	93272.00.00	DIAPHARM GMBH & CO. KG	DE
Ibuprofen/Pseudoephedrinhydrochlorid Diapharm 200 mg/30 mg Filmtabletten	DE/H/4186/001	93271.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
Momenxsin 200 mg/30 mg compresse rivestite con film	DE/H/4186/001	043682018	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momenxsin 200 mg/30 mg compresse rivestite con film	DE/H/4186/001	043682020	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momenxsin 200 mg/30 mg compresse rivestite con film	DE/H/4186/001	043682032	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Momenxsin 200 mg/30 mg compresse rivestite con film	DE/H/4186/001	043682044	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ибупром Синус 200 mg/30 mg обвити таблетки	not available	20080222	US PHARMACIA SP. Z O.O.	BG
ratioGrippal® 200 mg/30 mg Filmtabletten	not available	92698.00.00	RATIOPHARM GMBH	DE
Fidiprofen flu 200 mg+ 30 mg tablete	not available	UP/I-530-09/11-01/363	FIDIFARM D.O.O.	HR
Sudafed Sinus Pressure & Pain Tablets	not available	PL 15513/0126	MCNEIL PRODUCTS LIMITED	UK
Sudafed Sinus Pressure & Pain Tablets	not available	PL 15513/0126	MCNEIL PRODUCTS LIMITED	UK
Sudafed Sinus Pressure & Pain Tablets	not available	PL 15513/0126	MCNEIL PRODUCTS LIMITED	UK
Sudafed Sinus Pressure & Pain Tablets	not available	PL 15513/0126	MCNEIL PRODUCTS LIMITED	UK
Modafen potahované tablety	not available	07/136/98-C	ZENTIVA, K.S.	CZ
Modafen potahované tablety	not available	07/136/98-C	ZENTIVA, K.S.	CZ
Modafen potahované tablety	not available	07/136/98-C	ZENTIVA, K.S.	CZ
Modafen potahované tablety	not available	07/136/98-C	ZENTIVA, K.S.	CZ
Ibum Zatoki, 200 mg + 30 mg, tabletki powlekane	not available	21376	HASCO-LEK	PL
IBUM GRIP, 200 mg + 30 mg, tabletki powlekane	not available	10924	HASCO-LEK	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
Nurofen Sinus and Pain Film-Coated Tablets Ibuprofen 200mg Pseudoephedrine Hydrochloride 30mg	not available	PA 979/65/1	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Sinus and Pain Film-Coated Tablets Ibuprofen 200mg Pseudoephedrine Hydrochloride 30mg	not available	PA 979/65/1	RECKITT BENCKISER IRELAND LTD.	IE
Nurofen Cold and Flu 200 mg/30 mg filmtabletta	not available	OGYI-T-6797/02	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen Sinus Pressure & Headache Relief 200mg/30mg Tablets	not available	PL 00063/0718	RECKITT BENCKISER HEALTHCARE (UK) LIMITED	UK
Nurofen Raceala si Gripa 200 mg/30 mg comprimate filmate	not available	4144/2011/02	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	RO
Нурофен Стопколд 200 mg/30 mg филмирани таблетки (Nurofen Stopcold 200 mg/30 mg film-coated tablets)	not available	9800356	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	BG
Nurofen Cold and Flu	not available	19671	RECKITT BENCKISER HELLAS CHEMICALS ABEE	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen StopGrip potahované tablety	not available	07/612/96-C	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	CZ
RHINUREFLEX, comprimé pelliculé	not available	338 856-5	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN RHUME, comprimé pelliculé	not available	343 625-8	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen Cold and Flu 200 mg/30mg filmtabletta	not available	OGYI-T-6797/01	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	HU
Nurofen Cold & Flu Film-coated Tablets, Ibuprofen 200mg, Pseduoephedrine Hydrochloride 30mg	not available	PA 979/33/1	RECKITT BENCKISER IRELAND LTD.	IE
NUROFEN INFLUENZA E RAFFREDDORE 200 mg + 30 mg Compresse Rivestite	not available	034246013	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	IT
NUROFEN INFLUENZA E RAFFREDDORE 200 mg + 30 mg Compresse Rivestite	not available	034246025	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	IT
Nurofen Antigrip 200 mg/30 mg apvalkotās tabletes	not available	04-0197	RECKITT BENCKISER (POLAND) S.A.	LV
Nurofen Cold & Flu	not available	MA 190/00404	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen Zatoki, 200 mg + 30 mg, tabletki powlekane	not available	7787	RECKITT BENCKISER (POLAND) S.A.	PL
Nurofen Răceală şi Gripă 200 mg/30 mg comprimate filmate	not available	4144/2011/01	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	RO
NUROFEN STOPGRIP	not available	07/0662/96-S	RECKITT BENCKISER HEALTHCARE INTERNATIONAL LIMITED	SK
Nurofen Cold & Flu	not available	PL 00063/0375	RECKITT BENCKISER HEALTHCARE (UK) LIMITED	UK
NUROFEN COLD & FLU	not available	44616/10	RECKITT BENCKISER HELLAS CHEMICALS ABEE	GR
Nurofen Cold and Flu 200 mg + 30 mg filmom obložene tablete	not available	UP/I-530-09/11-02/20	RECKITT BENCKISER (CROATIA) D.O.O.	HR
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/4352/001	91698	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	GR
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE BOEHRINGER INGELHEIM 200 mg/30 mg comprimés pelliculés	UK/H/4352/001	2014040030	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	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
Ibuprofen/Pseudoephedrine hydrochloride Boehringer Ingelheim 200 mg/30 mg Filmtabletten	UK/H/4352/001	2014040030	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	LU
Ibuprofen/Pseudoephedrine hydrochloride Boehringer Ingelheim 200 mg/30 mg filmomhulde tabletten	UK/H/4352/001	BE445164	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	BE
Ibuprofen/Pseudoephedrine hydrochloride Boehringer Ingelheim 200 mg/30 mg Filmtabletten	UK/H/4352/001	BE445164	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	BE
Ibuprofen/Pseudoephedrine hydrochloride Boehringer Ingelheim 200 mg/30 mg filmomhulde tabletten	UK/H/4352/001	BE445164	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	BE
Bisolpront 200 mg + 30 mg comprimidos revestidos por película	UK/H/4352/001	5666110	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	PT
Bisolpront 200 mg + 30 mg comprimidos revestidos por película	UK/H/4352/001	5666102	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	PT
BoxaGrippal 200 mg / 30 mg Filmtabletten	UK/H/4352/001	85528.00.00	BOEHRINGER INGELHEIM INTERNATIONAL 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
Мукогрип 200 mg/30 mg филмирани таблетки	UK/H/4352/001	20120408	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	BG
Lasynac 200mg/30mg film coated tablets	UK/H/4352/001/DC	PL 14598/0091	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	UK
Bisolfren 200 mg/30 mg comprimidos recubiertos con película	UK/H/4352/001	76389	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	ES
BoxaGrippal™ 200 mg/30 mg - Filmtabletten	UK/H/4352/001	1-31668	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	AT
Mucogrip 200mg/30mg filmtabletta	UK/H/4352/001	OGYI-T-22371/01	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	HU
Mucogrip 200mg/30mg filmtabletta	UK/H/4352/001	OGYI-T-22371/02	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	HU
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/4352/001	21691	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	CY
IBUPROFENE E PSEUDOEFEDRINA BOEHRINGER INGELHEIM 200 mg/30 mg compresse rivestite con film	UK/H/4352/001	041218013	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	IT
IBUPROFENE E PSEUDOEFEDRINA BOEHRINGER INGELHEIM 200 mg/30 mg compresse rivestite con film	UK/H/4352/001	041218025	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE BOEHRINGER INGELHEIM 200 mg/30 mg comprimés pelliculés	UK/H/4352/001	BE445164	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	BE
Dolormin Grippal 200 mg/30 mg Filmtabletten	DE/H/4182/001/DC	93267.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin Grippal 200 mg/30 mg Filmtabletten	DE/H/4182/001/DC	93267.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin Grippal 200 mg/30 mg Filmtabletten	DE/H/4182/001/DC	93267.00.00	JOHNSON & JOHNSON GMBH	DE
Dolormin Grippal 200 mg/30 mg Filmtabletten	DE/H/4182/001/DC	93267.00.00	JOHNSON & JOHNSON GMBH	DE
Ibuprofen/Pseudoephedrine Hydrochloride 200mg/30mg film coated tablets	DE/H/4182/001	PA823/067/001	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Ibuprofen/Pseudoephedrine Hydrochloride 200mg/30mg film coated tablets	DE/H/4182/001	PA823/067/001	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
Ibuprofen/Pseudoephedrine Hydrochloride 200mg/30mg film coated tablets	DE/H/4182/001	PA0823/067/001	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibuprofen/Pseudoephedrine Hydrochloride 200mg/30mg film coated tablets	DE/H/4182/001	PA823/067/001	MCNEIL HEALTHCARE (IRELAND) LIMITED	IE
IbuHEXAL Grippal 200 mg/30 mg Filmtabletten	DE/H/4184/001	93269.00.00	HEXAL AG	DE
Ibutren Flu 200 mg/30 mg filmom obložene tablete	DE/H/4184/001	HR-H-099217339	SANDOZ D.O.O.	HR
Dolorflu akut 200 mg/30 mg - Filmtabletten	DE/H/4184/001	136793	SANDOZ GMBH	AT
ILOXEN 200 mg/30 mg apvalkotās tabletes	DE/H/4185/001	16-0009	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LV
GRIPOMED 200 mg/30 mg Filmtabletten	DE/H/4185/001	93270.00.00	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	DE
Febrilek 200 mg / 30 mg õhukese polümeerikattega tabletid	DE/H/4185/001	899915	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
ILOXEN 200 mg/30 mg plėvele dengtos tabletės	DE/H/4185/001	LT/1/16/3881/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
ILOXEN 200 mg/30 mg plėvele dengtos tabletės	DE/H/4185/001	LT/1/16/3881/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
ILOXEN 200 mg/30 mg plėvele dengtos tabletės	DE/H/4185/001	LT/1/16/3881/003	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
ILOXEN 200 mg/30 mg plėvele dengtos tabletės	DE/H/4185/001	LT/1/16/3881/004	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Efedoxin, 200 mg + 30 mg, tabletki powlekane	DE/H/4185/001	23302	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
BoxaGrippal® Erkältungssaft 200 mg/10 ml + 30 mg/10 ml Suspension zum Einnehmen	UK/H/5545/001	90668.00.00	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	DE
Ibuprofen/Pseudoephedrine hydrochloride 100mg/15mg per 5 ml Oral suspension	UK/H/5545/001	PL 14598/0103	BOEHRINGER INGELHEIM INTERNATIONAL GMBH	UK
Acatar Zatoki, 200 mg + 30 mg, tabletki powlekane	not available	11339	US PHARMACIA SP. Z O.O.	PL
IBUPROM ZATOKI, 200 mg + 30 mg, tabletki powlekane	not available	9650	US PHARMACIA SP. Z O.O.	PL
Advil Cold 200 mg/30 mg bevont tabletta	IE/H/0420/001	OGYI-T-22705/02	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Advil Cold 200 mg/30 mg bevont tabletta	IE/H/0420/001	OGYI-T-22705/01	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	HU
Ibuprofen/Pseudoephedrine HCl Pfizer 200 mg/30 mg comprimés enrobés	IE/H/0420/001	BE463680	PFIZER B.V.	BE
Robicold 200 mg/30 mg obalené tablety	IE/H/0420/001	07/040/15-C	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RobiCold Sinus Relief 200 mg, 30 mg Tablets	IE/H/0420/001	PL 00165/0391	PFIZER CONSUMER HEALTHCARE LTD.	UK
Ibuprofen/Pseudoephedrine HCl Pfizer 200 mg/30 mg omhulde tabletten	IE/H/0420/001	BE463680	PFIZER B.V.	BE
Ibuprofen/Pseudoephedrine HCl Pfizer 200mg/30 mg überzogene Tabletten	IE/H/0420/001	BE463680	PFIZER B.V.	BE
Robicold 200 mg/30 mg obalené tablety	IE/H/0420/001	07/0264/14-S	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	SK
Advil Zatoki, 200 mg + 30 mg, tabletki powlekane	IE/H/0420/001	22369	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	PL
RobiCold Cold & Flu Relief 200 mg / 30 mg Soft Capsules	UK/H/5670/001	PL 00165/0388	PFIZER CONSUMER HEALTHCARE LTD.	UK
Advil Sinus Relief Soft Capsules Ibuprofen 200 mg Pseudoephedrine hydrochloride 30 mg	UK/H/5670/001	PA 0822/164/002	PFIZER HEALTHCARE IRELAND	IE
Robicold Rapid 200 mg/30 mg mäkké kapsuly	UK/H/5670/001	07/0235/16-S	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	SK
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/01	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/02	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/08	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/14	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/12	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/03	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/10	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/06	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/04	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/16	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/05	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/13	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/07	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/11	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus Relief Soft Capsules Ibuprofen 200 mg Pseudoephedrine hydrochloride 30 mg	UK/H/5670/001	MA969/00202	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	MT
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/09	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
Advil Sinus și Răceală 200 mg/30 mg capsule moi	UK/H/5670/001	8760/2016/15	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	RO
SpaltGrippal 200 mg/30 mg Weichkapseln	UK/H/5671/001	94021.00.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
Ibuprofen/Pseudoephedrine HCl Pfizer 200 mg / 30 mg Comprimés enrobé	IE/H/0420/001	2015040073	PFIZER B.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Spalt Effekt 30mg/200 mg überzogene Tabletten	FR/H/0238/001	62958.00.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
RHINADVIL RHUME IBUPROFENE/PSEUDOEPHEDRINE, comprimé enrobé	FR/H/0238/001	334 084-8	PFIZER SANTE FAMILIALE SAS	FR
Advil Cold & Flu Coated Tablets Ibuprofen 200 mg Pseudoephedrine Hydrochloride 30 mg	IE/H/0420/001	PA 822/164/1	PFIZER HEALTHCARE IRELAND	IE
Advil Cold & Flu Coated Tablets Ibuprofen 200 mg Pseudoephedrine Hydrochloride 30 mg	not available	MA 969/00201	PFIZER CONSUMER HEALTHCARE LTD.	MT
Boots Cold & Flu Relief with Ibuprofen	not available	00014/0600	THE BOOTS COMPANY PLC	UK
Ibuprofen/Pseudoephedrinhydrochlorid Krewel 200 mg/30 mg Filmtabletten	DE/H/4188/001	93273.00.00	KREWEL MEUSELBACH GMBH	DE
Grippecton 200 mg/30 mg potahované tablety	DE/H/4188/001	07/155/16-C	KREWEL MEUSELBACH GMBH	CZ
Grippecton 200 mg/30 mg filmom obalené tablety	DE/H/4188/001	07/0143/16-S	KREWEL MEUSELBACH GMBH	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
Ibuprofene e pseudoefedrina Wick Pharma 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499032	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	IT
Ibuprofene e pseudoefedrina Wick Pharma 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499044	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	IT
WICK DuoGrippal 200 mg/30 mg Filmtabletten	IT/H/0331/001	88707.00.00	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	DE
Gripaduo 200 mg/30 mg comprimidos recubiertos con película	IT-H-0331-001-DC	78831	LABORATORIOS VICKS, S.L.	ES
Ibuprofen/Pszeudoefedrin Wick 200 mg/30 mg filmtabletta	IT/H0331/001	OGYI-T-22836/01	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	HU
Ibuprofen/Pszeudoefedrin Wick 200mg/30mg filmtabletta	IT/H0331/001/DC	OGYI-T-22836/02	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	HU
Ibuprofen/Pszeudoefedrin Wick 200mg/30mg filmtabletta	IT/H0331/001/DC	OGYI-T-22836/03	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	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
Ibuprofene e pseudoefedrina Wick Pharma 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499057	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	IT
Infex Zatoki	IT-H-0331-001-DC	22111	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
WICK DayMed Duo 200mg/30mg Filmtabletten	IT/H/0331/001	135488	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
TEDOLFEN 200mg/30mg comprimate filmate	IT/H/0331/001	6744/2014/01-04	TEVA PHARMACEUTICALS S.R.L	RO
ALGOFLEX COLD 200 mg/30 mg filmtabletta	not available	OGYI-T-22831/03	SANOFI-AVENTIS ZRT	HU
ALGOFLEX COLD 200 mg/30 mg filmtabletta	not available	OGYI-T-22831/01	SANOFI-AVENTIS ZRT	HU
ALGOFLEX COLD 200 mg/30 mg filmtabletta	not available	OGYI-T-22831/02	SANOFI-AVENTIS ZRT	HU
MODAFEN, 200 mg + 30 mg, tabletki powlekane	not available	9603	ZENTIVA, K.S.	PL
MODAFEN, 200 mg + 30 mg, tabletki powlekane	not available	9603	ZENTIVA, K.S.	PL
MODAFEN, 200 mg + 30 mg, tabletki powlekane	not available	9603	ZENTIVA, K.S.	PL
RHINATHIOL COLD 200 MG/30 MG FILMTABLETTA	not available	OGYI-T-9161/02	SANOFI-AVENTIS ZRT	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RHINATHIOL COLD 200 MG/30 MG FILMTABLETTA	not available	OGYI-T-9161/01	SANOFI-AVENTIS ZRT	HU
RHINATHIOL COLD 200 MG/30 MG FILMTABLETTA	not available	OGYI-T-9161/03	SANOFI-AVENTIS ZRT	HU