11 February 2021
EMA/96159/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ibuprofen / pseudoephedrine

Procedure no.: PSUSA/00001711/202007

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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
Metafen ZATOKI, 200 mg + 30 mg, tabletki	not available	11005	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
MODAFEN EXTRA GRIP, 200 mg + 30 mg, tabletki powlekane	not available	9603	SANOFI-AVENTIS SP Z.O.O.	PL
MODAFEN EXTRA GRIP, 200 mg + 30 mg, tabletki powlekane	not available	9603	SANOFI-AVENTIS SP Z.O.O.	PL
Ibuprofen and Pseudoephedrine hydrochloride Farmalider 200 mg/ 30 mg/10 ml oral suspension	not available	PL 35667/0020	FARMALIDER, S.A.	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/01	SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/02	SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/03	SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/04	SOLACIUM PHARMA S.R.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
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/05	SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/01	S.C. SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/02	S.C. SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/03	S.C. SOLACIUM PHARMA S.R.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
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/05	S.C. SOLACIUM PHARMA S.R.L.	RO
Ibusinus 200 mg/30 mg comprimate filmate	not available	3080/2010/04	S.C. SOLACIUM PHARMA S.R.L.	RO
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE SANOFI 200 mg/30 mg comprimés pelliculés	DE/H/5826/001	0735942	SANOFI BELGIUM	LU
Ibuprofen/Pseudoephedrine hydrochloride Sanofi 200 mg/30 mg Filmtabletten	DE/H/5826/001	BE445164	SANOFI BELGIUM	BE
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE SANOFI 200 mg/30 mg comprimés pelliculés	DE/H/5826/001	BE445164	SANOFI 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
Ibuprofen/Pseudoephedrine hydrochloride Sanofi 200 mg/30 mg filmomhulde tabletten	DE/H/5826/001	BE445164	SANOFI BELGIUM	BE
Bisolpront 200 mg + 30 mg comprimidos revestidos por película	DE/H/5826/001	5666110	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Bisolpront 200 mg + 30 mg comprimidos revestidos por película	DE/H/5826/001	5666102	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
ZERINOACTIV 200 mg/30 mg compresse rivestite con film	DE/H/5826/001	041218013	SANOFI S.P.A	IT
ZERINOACTIV 200 mg/30 mg compresse rivestite con film	DE/H/5826/001	041218025	SANOFI 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
Мукогрип 200 mg/30 mg филмирани таблетки	DE/H/5826/001	20120408	SANOFI BULGARIA EOOD	BG
Bisolfren 200 mg/30 mg comprimidos recubiertos con película	DE/H/5826/001	76389	SANOFI-AVENTIS, S.A.	ES
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5826/001	91698	SANOFI-AVENTIS AEBE	GR
Lasynac 200mg/30 mg film coated tablets	DE/H/5826/001	PL 04425/0722	AVENTIS PHARMA LTD	UK
Lasynac 200mg/30 mg film coated tablets	DE/H/5826/001	PL 04425/0722	AVENTIS PHARMA LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Мукогрип 200 mg/30 mg филмирани таблетки	DE/H/5826/001	20120408	SANOFI BULGARIA EOOD	BG
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5826/001	21691	SANOFI-AVENTIS AEBE	CY
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5826/001	21691	SANOFI-AVENTIS AEBE	CY
Bisolfren 200 mg/30 mg comprimidos recubiertos con película	DE/H/5826/001	76389	SANOFI-AVENTIS, S.A.	ES
Mucocold 200mg/30mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/5826/001	91698	SANOFI-AVENTIS AEBE	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
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE SANOFI 200 mg/30 mg comprimés pelliculés	DE/H/5826/001	BE445164	SANOFI BELGIUM	BE
Ibuprofen/Pseudoephedrine hydrochloride Sanofi 200 mg/30 mg filmomhulde tabletten	DE/H/5826/001	BE445164	SANOFI BELGIUM	BE
BoxaGrippal® Erkältungstabletten 200 mg / 30 mg Filmtabletten	DE/H/5826/001	85528.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal® Erkältungstabletten 200 mg / 30 mg Filmtabletten	DE/H/5826/001	85528.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal® 200 mg/30 mg - Filmtabletten	DE/H/5826/001	1-31668	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	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
BoxaGrippal® 200 mg/30 mg - Filmtabletten	DE/H/5826/001	1-31668	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	AT
Ibuprofen/Pseudoephedrine hydrochloride Sanofi 200 mg/30 mg Filmtabletten	DE/H/5826/001	BE445164	SANOFI BELGIUM	BE
IBUPROFEN/PSEUDOEPHEDRINE HYDROCHLORIDE SANOFI 200 mg/30 mg comprimés pelliculés	DE/H/5826/001	0735956	SANOFI BELGIUM	LU
Larofen Plus 200mg/30mg comprimate filmate	not available	9424/2016/01	LAROPHARM SRL	RO
GRIPPAL COMPLEX DoppelherzPharma 200 mg/30 mg Filmtabletten	DE/H/4187/001	93272.00.00	DOPPELHERZ PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ibucomp 200 mg/30 mg Filmtabletten	AT/H/0614/001	137621	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
SpaltGrippal 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	FR
Laboratoria Polfałódź ZATOKI, 200 mg + 30 mg, tabletki	not available	21816	LABOLATORIA POLFA ŁÓDŹ 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
Boots Cold & Flu Relief with Ibuprofen.	not available	PL 00014/0600	THE BOOTS COMPANY PLC	UK
MODAFEN 200 mg/30 mg filmom obalené tablety	not available	07/0263/01-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
MODAFEN 200 mg/30 mg filmom obalené tablety	not available	07/0263/01-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
MODAFEN 200 mg/30 mg filmom obalené tablety	not available	07/0263/01-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
ILOXEN 200 mg/30 mg apvalkotās tabletes	DE/H/4185/001	16-0009	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
GRIPOMED 200 mg/30 mg Filmtabletten	DE/H/4185/001	93270.00.00	BAUSCH HEALTH IRELAND LIMITED	DE
Febrilek 200 mg / 30 mg õhukese polümeerikattega tabletid	DE/H/4185/001	899915	BAUSCH HEALTH IRELAND LIMITED	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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ILOXEN 200 mg/30 mg plėvele dengtos tabletės	DE/H/4185/001	LT/1/16/3881/004	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Gripomed 200 mg/30 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/4185/001	3090001	BAUSCH HEALTH IRELAND LIMITED	GR
Advil Cold & Flu Coated Tablets Ibuprofen 200 mg Pseudoephedrine Hydrochloride 30 mg	not available	MA 969/00201	PFIZER CONSUMER HEALTHCARE LTD.	MT
Rhinorelief 200 mg/30 mg tablete	HR-H-550772325	HR-H-550772325	JADRAN-GALENSKI LABORATORIJ D.D.	HR
Ibuprofen and Pseudoephedrine hydrochloride Farmalider 200 mg/ 30 mg/10 ml oral suspension	not available	PL 35667/0020	FARMALIDER, S.A.	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rhinorelief 200 mg/30 mg tablete	HR-H-550772325	HR-H-550772325	JADRAN-GALENSKI LABORATORIJ D.D.	HR
Frenaxsin 200 mg/30 mg comprimidos recubiertos con película.	not available	84970	JOHNSON & JOHNSON S.A.	ES
Frenaxsin 200 mg/30 mg comprimidos recubiertos con película.	not available	84970	JOHNSON & JOHNSON S.A.	ES
Ibuprofeno/Pseudoefedrina Nutra Essential 20 mg/ml + 3 mg/ml suspensión oral	not available	84977	NUTRA ESSENTIAL OTC, S.L.	ES
Acatar Zatoki, 200 mg + 30 mg, tabletki powlekane	not available	11339	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
RobiCold Cold & Flu Relief 200 mg / 30 mg Soft Capsules	IE/H/0642/001	PL 00165/0388	PFIZER CONSUMER HEALTHCARE LTD.	UK
Advil Cold & Flu Relief 200 mg / 30 mg Soft Capsules	IE/H/0642/001	PA 822/164/002	PFIZER HEALTHCARE IRELAND	IE
Advil Sinus Relief 200 mg / 30 mg Soft Capsules	IE/H/0642/001	MA 969/00202	PFIZER HEALTHCARE IRELAND	MT
Advil Cold Rapid 200 mg/30 mg lágy kapszula	IE/H/0642/001	OGYI-T-22705/03	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Cold Rapid 200 mg/30 mg lágy kapszula	IE/H/0642/001	OGYI-T-22705/04	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Cold Rapid 200 mg/30 mg lágy kapszula	IE/H/0642/001	OGYI-T-22705/06	GLAXOSMITHKLINE-CONSUMER 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
Advil Cold Rapid 200 mg/30 mg lágy kapszula	IE/H/0642/001	OGYI-T-22705/05	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Robicold Rapid 200 mg/30 mg mäkké kapsuly	IE/H/0642/001	07/0235/16-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.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
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/06	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/05	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/07	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/09	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/08	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/12	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.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
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/15	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/11	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/14	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/10	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/13	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Advil Răceală și Gripă 200 mg/30 mg capsule moi	IE/H/0642/001	8760/2016/16	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.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
RobiCold Sinus Relief 200 mg, 30 mg Tablets	IE/H/0420/001	PL 00165/0391	PFIZER CONSUMER HEALTHCARE LTD.	UK
Ibuprofen/Pseudoephedrine HCl Pfizer 200mg/30 mg überzogene Tabletten	IE/H/0420/001	BE463680	PFIZER B.V.	BE
Advil Zatoki, 200 mg + 30 mg, tabletki powlekane	IE/H/0420/001	22369	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	PL
Ibuprofen/Pseudoephedrine HCl Pfizer 200 mg/30 mg comprimés enrobés	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
Advil Cold & Flu Coated Tablets Ibuprofen 200 mg Pseudoephedrine Hydrochloride 30 mg	IE/H/0420/001	PA 0822/164/001	PFIZER HEALTHCARE IRELAND	IE
Advil Cold 200 mg/30 mg bevont tabletta	IE/H/0420/001	OGYI-T-22705/02	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Advil Cold 200 mg/30 mg bevont tabletta	IE/H/0420/001	OGYI-T-22705/01	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Robicold 200 mg/30 mg obalené tablety	IE/H/0420/001	07/040/15-C	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC 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
Robicold 200 mg/30 mg obalené tablety	IE/H/0420/001	07/0264/14-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Modafen 200 mg/30 mg comprimate filmate	not available	480/2007/02	SANOFI ROMANIA SRL	RO
Modafen 200 mg/30 mg comprimate filmate	not available	480/2007/03	SANOFI ROMANIA SRL	RO
Modafen 200 mg/30 mg comprimate filmate	not available	480/2007/04	SANOFI ROMANIA SRL	RO
Modafen 200 mg/30 mg comprimate filmate	not available	480/2007/01	SANOFI ROMANIA SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
IBUVALEN FLU 200 mg/30 mg comprimate filmate	not available	9937/2017/01	POLISANO PHARMACEUTICALS S.A.	RO
BoxaGrippal forte Erkältungstabletten 400 mg/60 mg Filmtabletten	DE/H/5826/002	97699.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal forte Erkältungstabletten 400 mg/60 mg Filmtabletten	DE/H/5826/002	97699.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal forte Erkältungstabletten 400 mg/60 mg Filmtabletten	DE/H/5826/002	97699.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal® forte 400 mg/60 mg - Filmtabletten	DE/H/5826/002	138002	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	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
BoxaGrippal® forte 400 mg/60 mg - Filmtabletten	DE/H/5826/002	138002	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	AT
BoxaGrippal® forte 400 mg/60 mg - Filmtabletten	DE/H/5826/002	138002	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	AT
BoxaGrippal® Erkältungssaft 200 mg/10 ml + 30 mg/10 ml Suspension zum Einnehmen	DE/H/5827/001	90668.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
BoxaGrippal® Erkältungssaft 200 mg/10 ml + 30 mg/10 ml Suspension zum Einnehmen	DE/H/5827/001	90668.00.00	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	DE
IBUPROM ZATOKI, 200 mg + 30 mg, tabletki powlekane	not available	9650	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
Sinuphene 200 mg/30 mg filmomhulde tabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg Filmtabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg filmomhulde tabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	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
Sinuphene 200 mg/30 mg Filmtabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg filmomhulde tabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg Filmtabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg Filmtabletten	IE/H/1134/01/DC	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
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Sinuphene 200 mg/30 mg filmomhulde tabletten	IE/H/1134/01/DC	BE484782	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	IE/H/1134/01/DC	PA 330/56/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	IE/H/1134/01/DC	PA 330/56/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	IE/H/1134/01/DC	PA 330/56/1	JOHNSON & JOHNSON (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
Non-Drowsy Sudapro Head Cold 200mg / 30mg Film-coated Tablets	IE/H/1134/01/DC	PA 330/56/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
ACTISINU 200 mg/30 mg, compresse rivestite con film	IE/H/1134/01/DC	043681042	JOHNSON & JOHNSON S.P.A.	IT
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	2016110302	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	2016110302	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	2016110302	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sinuphene 200 mg/30 mg comprimés pelliculés	IE/H/1134/01/DC	2016110302	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Sudafed Sinus Pressure & Pain 200mg/30mg film-coated tablets	IE/H/1134/01/DC	PL 15513/0396	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Sudafed Sinus Pressure & Pain 200mg/30mg film-coated tablets	IE/H/1134/01/DC	PL 15513/0396	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Sudafed Sinus Pressure & Pain 200mg/30mg film-coated tablets	IE/H/1134/01/DC	PL 15513/0396	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK
Sudafed Sinus Pressure & Pain 200mg/30mg film-coated tablets	IE/H/1134/01/DC	PL 15513/0396	MCNEIL PRODUCTS LIMITED (MAIDENHEAD)	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Non-Drowsy Sudapro Head Cold 200mg/30mg film-coated tablets	IE/H/1134/01/DC	MA 1315/00701	JOHNSON & JOHNSON (IRELAND) LTD.	MT
Non-Drowsy Sudapro Head Cold 200mg/30mg film-coated tablets	IE/H/1134/01/DC	MA 1315/00701	JOHNSON & JOHNSON (IRELAND) LTD.	MT
Non-Drowsy Sudapro Head Cold 200mg/30mg film-coated tablets	IE/H/1134/01/DC	MA 1315/00701	JOHNSON & JOHNSON (IRELAND) LTD.	MT
Non-Drowsy Sudapro Head Cold 200mg/30mg film-coated tablets	IE/H/1134/01/DC	MA 1315/00701	JOHNSON & JOHNSON (IRELAND) LTD.	MT
Cegrinaso 200 mg + 30 mg comprimidos revestidos por película	IE/H/1134/001	5779525	JOHNSON & JOHNSON 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
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
RHINATHIOL COLD 200 MG/30 MG FILMTABLETTA	not available	OGYI-T-9161/02	SANOFI-AVENTIS ZRT	HU
Modafen 200 mg/30 mg potahované tablety	not available	07/136/98-C	SANOFI-AVENTIS SRO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Modafen 200 mg/30 mg potahované tablety	not available	07/136/98-C	SANOFI-AVENTIS SRO	CZ
Modafen 200 mg/30 mg potahované tablety	not available	07/136/98-C	SANOFI-AVENTIS SRO	CZ
Modafen 200 mg/30 mg potahované tablety	not available	07/136/98-C	SANOFI-AVENTIS SRO	CZ
Ibu - 1 A Pharma Grippal 200 mg/30 mg Filmtabletten	DE/H/4184/001	93269.00.00	1 A PHARMA GMBH	DE
Dolorflu akut 200 mg/30 mg - Filmtabletten	DE/H/4184/001	136793	SANDOZ 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
Daikol 200 mg/30 mg plėvele dengtos tabletės	not available	LT/1/16/3982/001	SIA INGEN PHARMA	LT
Vicks Flu Action 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499032	PROCTER & GAMBLE S.R.L	IT
Vicks Flu Action 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499044	PROCTER & GAMBLE S.R.L	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	78.831	LABORATORIOS VICKS, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 200 mg/30 mg filmtabletta	IT/H/0331/001	OGYI-T-22836/02	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	HU
Ibuprofen/Pszeudoefedrin Wick 200 mg/30 mg filmtabletta	IT/H/0331/001	OGYI-T-22836/03	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	HU
Vicks Flu Action 200 mg/30 mg compresse rivestite con film	IT/H/0331/001	042499057	PROCTER & GAMBLE S.R.L	IT
Ibuprofen + Pseudoefedryna 123ratio, 200 mg + 30 mg, tabletki powlekane	IT/H/0331/001	22111	TEVA PHARMACEUTICALS POLSKA 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
TEDOLFEN 200mg/30mg comprimate filmate	IT/H/0331/001	6744/2014/01	TEVA PHARMACEUTICALS S.R.L	RO
WICK DayMed Duo 200 mg/30 mg Filmtabletten	IT/H/0331/001	135488	WICK PHARMA ZWEIGNIEDERLASSUNG DER PROCTER & GAMBLE GMBH	AT
TEDOLFEN 200mg/30mg comprimate filmate	IT/H/0331/001	6744/2014/02	TEVA PHARMACEUTICALS S.R.L	RO
TEDOLFEN 200mg/30mg comprimate filmate	IT/H/0331/001	6744/2014/03	TEVA PHARMACEUTICALS S.R.L	RO
Ibuseu, 200 mg/30 mg filmdragerade tabletter	SE/H/1892/001	52535	ORIFARM GENERICS A/S	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
APSELAN PLUS, 200 mg + 30 mg, tabletki powlekane	not available	25027	POLFARMEX S.A.	PL
Ibuprofen/ Pseudoephedrine hydrochloride Pfizer 200mg/30mg Capsules	DE/H/5902/001	PL 00165/0389	PFIZER CONSUMER HEALTHCARE LTD.	UK
SpaltGrippal 200 mg/30 mg Weichkapseln	DE/H/5902/001	94021.00.00	PFIZER CONSUMER HEALTHCARE GMBH	DE
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 228 7 7	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 228 6 0	PFIZER SANTE FAMILIALE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 228 8 4	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 228 9 1	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 0 7	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 1 4	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 2 1	PFIZER SANTE FAMILIALE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 3 8	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 4 5	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 5 2	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 6 9	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 8 3	PFIZER SANTE FAMILIALE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 229 9 0	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 230 0 3	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 230 1 0	PFIZER SANTE FAMILIALE	FR
RHINADVILCAPS RHUME IBUPROFENE/PSEUDOEPHEDRINE 200 mg /30 mg, capsule molle	DE/H/5902/001	34009 301 230 2 7	PFIZER SANTE FAMILIALE	FR
Nurofen Sinus and Pain Film-Coated Tablets Ibuprofen 200mg Pseudoephedrine Hydrochloride 30mg	not available	PA 979/65/1	RECKITT BENCKISER IRELAND LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nurofen 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/30mg filmtabletta	not available	OGYI-T-6797/02	RECKITT BENCKISER (HUNGARY) KFT	HU
Nurofen Sinus Relief	not available	PL 00063/0375	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Sinus Pressure & Headache Relief 200mg/30mg Tablets	not available	PL 00063/0718	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
Nurofen Răceală şi Gripă 200 mg/30 mg comprimate filmate	not available	4144/2011/02	RECKITT BENCKISER (ROMANIA) SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RHINUREFLEX, comprimé pelliculé	not available	338 856-5 OR 34009 338 856 5 6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN RHUME, comprimé pelliculé	not available	343 625-8 OR 34009 343 625 8 3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Нурофен Стопколд 200 mg/30 mg филмирани таблетки (Nurofen Stopcold 200 mg/30 mg film-coated tablets)	not available	9800356	RECKITT BENCKISER (ROMANIA) SRL	BG
NUROFEN® Cold and Flu	not available	19671	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
Nurofen StopGrip 200 mg/30 mg potahované tablety	not available	07/612/96-C	RECKITT BENCKISER (CZECH REPUBLIC) SPOL. S.R.O	CZ
RHINUREFLEX, comprimé pelliculé	not available	34009 338 856 5 6	RECKITT BENCKISER HEALTHCARE FRANCE	FR
NUROFEN RHUME, comprimé pelliculé	not available	34009 343 625 8 3	RECKITT BENCKISER HEALTHCARE FRANCE	FR
Nurofen Cold and Flu 200 mg/30mg filmtabletta	not available	OGYI-T-6797/01	RECKITT BENCKISER (HUNGARY) KFT	HU
Nurofen Cold & Flu Film-coated Tablets, Ibuprofen 200mg, Pseduoephedrine Hydrochloride 30mg	not available	PA 979/33/1	RECKITT BENCKISER IRELAND LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NUROFEN INFLUENZA E RAFFREDDORE 200 mg + 30 mg Compresse Rivestite	not available	034246013	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
NUROFEN INFLUENZA E RAFFREDDORE 200 mg + 30 mg Compresse Rivestite	not available	034246025	RECKITT BENCKISER HEALTHCARE (ITALIA) SPA	IT
Nurofen Antigrip 200 mg/30 mg apvalkotās tabletes	not available	04-0197	RECKITT BENCKISER (POLAND) S.A.	LV
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 (ROMANIA) SRL	RO
NUROFEN STOPGRIP	not available	07/0662/96-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 Cold & Flu	not available	PL 00063/0375	RECKITT BENCKISER HEALTHCARE (UK) LTD	UK
NUROFEN Cold and Flu	not available	44616/10/31-05-2011	RECKITT BENCKISER HELLAS HEALTHCARE SA	GR
Nurofen Cold and Flu 200 mg + 30 mg filmom obložene tablete	not available	HR-H-756646540	RECKITT BENCKISER (CROATIA) D.O.O.	HR
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
Ibum Zatoki Max, 400 mg + 60 mg, tabletki powlekane	not available	23760	HASCO-LEK	PL
ZATOTABS, 400 mg + 60 mg, tabletki powlekane	not available	24893	HASCO-LEK	PL
Pharmafren 200 mg / 30 mg cápsulas duras	not available	83605	LABORATORIOS CINFA, S.A.	ES
Ibuprofen/Pseudoephedrin ehydrochlorid 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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Grippecton 200 mg/30 mg filmom obalené tablety	DE/H/4188/001	07/0143/16-S	KREWEL MEUSELBACH GMBH	SK
Ibuprofen/Pseudoephedrin ehydrochlorid Angelini 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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Ibuprofen/Pseudoephedrinehydrochlorid Angelini 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
Ibuprofen/Pseudoephedrinehydrochlorid Angelini 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
Ibuprofen/Pseudoephedrinehydrochlorid Angelini 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
ratioGrippal® 200 mg/30 mg Filmtabletten	not available	92698.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ибупром Синус 200 mg/30 mg обвити таблетки	not available	20080222	US PHARMACIA SP. Z O.O.	BG