29 September 2022
EMA/806370/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): ketoprofen (all formulations except topical)

Procedure No. PSUSA/00001809/202201

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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Alrheumunâ, 50 mg	not available	6169911.00.00	TEOFARMA S.R.L.	DE
ARTROSILENE 160 mg supposte	not available	024022030	DOMPÉ FARMACEUTICI S.P.A.	IT
ARTROSILENE 160 mg/2 ml soluzione iniettabile	not available	024022170	DOMPÉ FARMACEUTICI S.P.A.	IT
ARTROSILENE 320 mg capsule rigide a rilascio prolungato	not available	024022129	DOMPÉ FARMACEUTICI S.P.A.	IT
BI PROFENID LP 100 mg, comprimé sécable à libération prolongée	not available	34009 399 803 9 3	SANOFI-AVENTIS FRANCE	FR
BI PROFENID LP 100 mg, comprimé sécable à libération prolongée	not available	34009 399 804 5 4	SANOFI-AVENTIS FRANCE	FR
BI PROFENID LP 100 mg, comprimé sécable à libération prolongée	not available	34009 399 802 2 5	SANOFI-AVENTIS FRANCE	FR
Bi-Profenid, 150 mg, tabletki o zmodyfikowanym uwalnianiu	not available	4121	SANOFI-AVENTIS FRANCE	PL
Bi-Profenid, 150 mg, tabletki o zmodyfikowanym uwalnianiu	not available	4121	SANOFI-AVENTIS FRANCE	PL
FASTUM®	not available	54.445	LABORATORIOS MENARINI, S.A.	ES
FLEXEN 100 mg polvere e solvente per soluzione iniettabile per uso endovenoso	not available	023401096	ITALFARMACO S.P.A	IT
FLEXEN 100 mg polvere e solvente per soluzione iniettabile per uso intramuscolare.	not available	023401108	ITALFARMACO 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
FLEXEN 100 mg supposte	not available	023401058	ITALFARMACO S.P.A	IT
FLEXEN 200 mg capsule rigide a rilascio prolungato	not available	023401110	ITALFARMACO S.P.A	IT
FLEXEN 50 mg capsule molli	not available	023401021	ITALFARMACO S.P.A	IT
Ibifen 100 mg capsule rigide	not available	024994081	IBI G. LORENZINI SPA	IT
Ibifen 100 mg/2,5 ml soluzione iniettabile per uso intramuscolare	not available	024994182	IBI G. LORENZINI SPA	IT
Ibifen 100 mg/2,5 ml soluzione iniettabile per uso intramuscolare	not available	024994232	IBI G. LORENZINI SPA	IT
Ibifen 100 mg/5 ml soluzione iniettabile per uso endovenoso	not available	024994194	IBI G. LORENZINI SPA	IT
Ibifen 100 mg/5 ml soluzione iniettabile per uso endovenoso	not available	024994244	IBI G. LORENZINI SPA	IT
Ibifen 200 mg compresse a rilascio prolungato	not available	024994168	IBI G. LORENZINI SPA	IT
Ibifen 25 mg/ml gocce orali soluzione.	not available	024994220	IBI G. LORENZINI SPA	IT
Ibifen 50 mg capsule rigide	not available	024994117	IBI G. LORENZINI SPA	IT
Ibifen 50 mg granulato effervescente	not available	024994170	IBI G. LORENZINI SPA	IT
KETIFEXIN 25 mg granule pentru soluţie oralã în plic	HR/H/0179/001	11353/2019/01	S.C. SANDOZ S.R.L.	RO
KETIFEXIN 25 mg granule pentru soluţie oralã în plic	HR/H/0179/001	11353/2019/02	S.C. SANDOZ S.R.L.	RO
KETIFEXIN 50 mg granule pentru soluţie	HR/H/0179/002	11354/2019/01	S.C. SANDOZ 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
oralã în plic				
KETIFEXIN 50 mg granule pentru soluţie oralã în plic	HR/H/0179/002	11354/2019/02	S.C. SANDOZ S.R.L.	RO
KETIFEXIN 50 mg granule pentru soluţie oralã în plic	HR/H/0179/002	11354/2019/03	S.C. SANDOZ S.R.L.	RO
Ketoflix, 50 mg, granulat do sporządzania roztworu doustnego, w saszetce	not available	26412	BIOFARM SP. Z O.O.	PL
Ketonal Akut Rapid 25 mg granule za oralnu otopinu	HR/H/0179/001	HR-H-136057643	SANDOZ D.O.O.	HR
Ketonal Akut Rapid 50 mg granule za oralnu otopinu u vrecici	HR/H/0179/002	HR-H-327547843	SANDOZ D.O.O.	HR
Ketonal Sprint, 25 mg, granulat do sporządzania roztworu doustnego, w saszetce	HR/H/0179/001	24985	SANDOZ GMBH	PL
Ketonal Sprint, 50 mg, granulat do sporządzania roztworu doustnego, w saszetce	HR/H/0179/002	24986	SANDOZ GMBH	PL
Ketoprofen 25mg Granules	IE/H/0959/001	PL 02855/0305	DOMPÉ FARMACEUTICI S.P.A.	XI
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 580 376 1 0	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 580 378 4 9	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 580 379 0 0	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg,	FR/H/0469/001	34009 419 166 9 4	CARELIDE	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
solution pour perfusion				
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 419 165 2 6	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 580 377 8 8	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 419 162 3 6	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001	34009 419 161 7 5	CARELIDE	FR
KETOPROFENE CARELIDE 100 mg, solution pour perfusion	FR/H/0469/001/DC	1211/11060032	CARELIDE	LU
KETOPROFENE MEDISOL 100 mg/4 ml, solution à diluer pour perfusion	not available	NL42621	MEDISOL	FR
KETOPROFENE RANBAXY 100 mg, comprimé pelliculé	not available	34009 378 482 9 9	RANBAXY PHARMACIE GENERIQUES	FR
KETOPROFENE RANBAXY 50 mg, gélule	not available	34009 385 246 5 9	RANBAXY PHARMACIE GENERIQUES	FR
KETOPROFENE ZENTIVA LP 100 mg, comprimé sécable à libération prolongée	not available	34009 347 191 2 7	ZENTIVA FRANCE	FR
KETOPROFENE ZENTIVA LP 100 mg, comprimé sécable à libération prolongée	not available	34009 347 189 8 4	ZENTIVA FRANCE	FR
KETOPROFENE ZENTIVA LP 100 mg, comprimé sécable à libération prolongée	not available	34009 347 190 6 6	ZENTIVA FRANCE	FR
OKi 160 mg supposte	not available	028511057	DOMPÉ FARMACEUTICI 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
OKi 160 mg/2 ml soluzione iniettabile per uso intramuscolare	not available	028511158	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi 80 mg granulato per soluzione orale	not available	028511095	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi 80 mg granulato per soluzione orale	not available	028511222	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi 80 mg granulato per soluzione orale	not available	28511210	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi 80 mg/ml gocce orali, soluzione	not available	028511145	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414092	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414104	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414066	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414080	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414041	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414054	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414039	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414015	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414130	DOMPÉ FARMACEUTICI 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
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414027	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414116	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414128	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414078	DOMPÉ FARMACEUTICI S.P.A.	IT
OKi dolore e febbre 25 mg compresse effervescenti	NL/H/4359/001	048414142	DOMPÉ FARMACEUTICI S.P.A.	IT
Okiact 25 mg Coated Granules in Sachet	DE/H/5268/01/DC	56847	DOMPÉ FARMACEUTICI S.P.A.	SE
okiact 25 mg comprimidos revestidos	NL/H/3583/001-002/DC	5791702	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg comprimidos revestidos	NL/H/3583/001-002/DC	5791652	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg comprimidos revestidos	NL/H/3583/001-002/DC	5791660	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg comprimidos revestidos	NL/H/3583/001-002/DC	5791710	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg comprimidos revestidos	NL/H/3583/001-002/DC	5791678	DOMPÉ FARMACEUTICI S.P.A.	PT
Okiact 25 mg filmdragerad tablett	DE/H/5300/001	56848	DOMPÉ FARMACEUTICI S.P.A.	SE
okiact 25 mg granulado revestido em saqueta	NL/H/3583/001-002/DC	5791736	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg granulado revestido em saqueta	NL/H/3583/001-002/DC	5791728	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg granulado revestido em saqueta	NL/H/3583/001-002/DC	5791769	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg granulado revestido em saqueta	NL/H/3583/001-002/DC	5791751	DOMPÉ FARMACEUTICI S.P.A.	PT
okiact 25 mg granulado	NL/H/3583/001-	5791744	DOMPÉ FARMACEUTICI	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
revestido em saqueta	002/DC		S.P.A.	
Okitask 25 mg apvalkotās granulas paciņā	NL/H/3583/001	19-0169	DOMPÉ FARMACEUTICI S.P.A.	LV
Okitask 25 mg apvalkotās tabletes	NL/H/3583/001	19-0170	DOMPÉ FARMACEUTICI S.P.A.	LV
Okitask 25 mg Befilmtes Granulat im Beutel	DE/H/5268/001	139014	DOMPÉ FARMACEUTICI S.P.A.	AT
Okitask 25 mg Befilmtes Granulat im Beutel	DE/H/5268/001/DC	2200136.00.00	DOMPÉ FARMACEUTICI S.P.A.	DE
Okitask 25 mg bruistabletten	NL/H/4359/001	RVG 122948	DOMPÉ FARMACEUTICI S.P.A.	NL
Okitask 25 mg Coated Granules in Sachet	IE/H/0959/001	PA 1186/024/001	DOMPÉ FARMACEUTICI S.P.A.	IE
Okitask 25 mg comprimate filmate	DE/H/5300/001	11609/2019/01	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg comprimate filmate	DE/H/5300/001	11609/2019/02	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg comprimate filmate	DE/H/5300/001	11609/2019/03	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg comprimate filmate	DE/H/5300/001	11609/2019/04	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg comprimate filmate	DE/H/5300/001	11609/2019/05	DOMPÉ FARMACEUTICI S.P.A.	RO
okitask 25 mg Effervescent tablet	NL/H/4359/01	36019	DOMPÉ FARMACEUTICI S.P.A.	FI
Okitask 25 mg Film-Coated Tablets	IE/H/0960/001	PA 1186/024/002	DOMPÉ FARMACEUTICI S.P.A.	IE
Okitask 25 mg filmom obalené tablety	DE/H/5300/001	29/0117/19-S	DOMPÉ FARMACEUTICI S.P.A.	SK
Okitask 25 mg filmomhulde tabletten	NL/H/3583/002	RVG 119317	DOMPÉ FARMACEUTICI S.P.A.	NL
Okitask 25 mg filmtabletta	NL/H/3583/002	OGYI-T-23279/04	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg filmtabletta	NL/H/3583/002	OGYI-T-23279/03	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg filmtablette	DE/H/5300/001/DC	2200521.00.00	DOMPÉ FARMACEUTICI S.P.A.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Okitask 25 mg Filmtabletten	DE/H/5300/001	139015	DOMPÉ FARMACEUTICI S.P.A.	AT
Okitask 25 mg granulat, drasjert i dosepose	DE/H/5268/001/DC	17-11858	DOMPÉ FARMACEUTICI S.P.A.	NO
Okitask 25 mg granulátum	NL/H/3583/001	OGYI-T-23279/02	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg granulátum	NL/H/3583/001	OGYI-T-23279/01	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg granule drajefiate în plic	DE/H/5268/001	11610/2019/01	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg granule drajefiate în plic	DE/H/5268/001	11610/2019/02	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg granule drajefiate în plic	DE/H/5268/001	11610/2019/04	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg granule drajefiate în plic	DE/H/5268/001	11610/2019/03	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg granule drajefiate în plic	DE/H/5268/001	11610/2019/05	DOMPÉ FARMACEUTICI S.P.A.	RO
Okitask 25 mg kalvopäällysteiset tabletit	NL/H/3583/002	34225	DOMPÉ FARMACEUTICI S.P.A.	FI
Okitask 25 mg obalené granule v sácku	DE/H/5268/001	29/350/17-C	DOMPÉ FARMACEUTICI S.P.A.	CZ
Okitask 25 mg obalené granuly vo vrecku	DE/H/5268/001	29/0116/19-S	DOMPÉ FARMACEUTICI S.P.A.	SK
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/04	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/06	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/11	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/02	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/07	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/09	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg	NL/H/4359/001	OGYI-T-23705/05	DOMPÉ FARMACEUTICI	HU

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pezsgotabletta			S.P.A.	
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/01	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/03	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/08	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/10	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/13	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/14	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg pezsgotabletta	NL/H/4359/001	OGYI-T-23705/12	DOMPÉ FARMACEUTICI S.P.A.	HU
Okitask 25 mg potahované tablety	DE/H/5300/001	29/351/17-C	DOMPÉ FARMACEUTICI S.P.A.	CZ
Okitask 25 mg rakeet	NL/H/3583/001	33753	DOMPÉ FARMACEUTICI S.P.A.	FI
Okitask 25 mg tablett, filmdrasjert	DE/H/5300/001	17-11857	DOMPÉ FARMACEUTICI S.P.A.	NO
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 1 5	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 2 2	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 3 9	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 4 6	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 5 3	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 6 0	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 7 7	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 990 8 4	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg,	NL/H/4359/01	34009 301 990 9 1	DOMPÉ FARMACEUTICI	FR

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comprimé effervescent			S.P.A.	
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 991 0 7	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 991 1 4	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé effervescent	NL/H/4359/01	34009 301 991 2 1	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé pelliculé	NL/H/3583/001	34009 301 933 9 6	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé pelliculé	NL/H/3583/001	34009 301 933 8 9	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé pelliculé	NL/H/3583/001-002	34009 301 933 0 2	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé pelliculé	NL/H/3583/001-002	34009 301 933 3 3	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, comprimé pelliculé	NL/H/3583/001-002	34009 301 933 1 9	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25 mg, granulés enrobés en sachet	NL/H/3583/001	34009 301 933 2 7	DOMPÉ FARMACEUTICI S.P.A.	FR
Okitask 25mg omhuld granulaat	NL/H/3583/001	RVG 118238	DOMPÉ FARMACEUTICI S.P.A.	NL
Okitask granulat powlekany, 25 mg	NL/H/3583/001-002/DC	24139	DOMPÉ FARMACEUTICI S.P.A.	PL
Okitask tabletki powlekane, 25 mg	NL/H/3583/001-002/DC	24202	DOMPÉ FARMACEUTICI S.P.A.	PL
Okitask, 25 mg graanulid	IE/H/0959/001	982319	DOMPÉ FARMACEUTICI S.P.A.	EE
Okitask, 25 mg õhukese polümeerikattega tabletid	IE/H/0960/001	982219	DOMPÉ FARMACEUTICI S.P.A.	EE
Orudis 100 mg depotkapsler	not available	7904	SANOFI-AVENTIS NORGE AS	NO
Orudis 100 mg hörð forðahylki	not available	930030	SANOFI-AVENTIS NORGE AS	IS
ORUDIS 100 mg kapsel, hård	not available	8644	SANOFI OY	FI
ORUDIS 100 mg kapsel, hård	not available	8644	SANOFI OY	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
ORUDIS 100 mg kapsel, hård	not available	8644	SANOFI OY	FI
Orudis 100 mg kapsel, hård	not available	10058	SANOFI AB	SE
ORUDIS 100 mg kapseli, kova	not available	8644	SANOFI OY	FI
ORUDIS 100 mg kapseli, kova	not available	8644	SANOFI OY	FI
ORUDIS 100 mg kapseli, kova	not available	8644	SANOFI OY	FI
Orudis 100 mg solución inyectable	not available	55857	SANOFI-AVENTIS, S.A.	ES
ORUDIS 100 mg supposte	not available	023183041	SANOFI S.R.L.	IT
ORUDIS 100mg/2ml soluzione iniettabile per uso intramuscolare	not available	023183205	SANOFI S.R.L.	IT
ORUDIS 200 mg capsule rigide a rilascio prolungato	not available	023183193	SANOFI S.R.L.	IT
Orudis 200 mg depotkapsler	not available	7816	SANOFI-AVENTIS NORGE AS	NO
Orudis 200 mg depotkapsler	not available	7816	SANOFI-AVENTIS NORGE AS	NO
Orudis 50 mg cápsulas	not available	52097	SANOFI-AVENTIS, S.A.	ES
ORUDIS 50 mg capsule rigide	not available	023183027	SANOFI S.R.L.	IT
Orudis 50 mg kapsel, hård	not available	9927	SANOFI AB	SE
Orudis Retard 200 mg comprimidos de liberación prolongada	not available	57128	SANOFI-AVENTIS, S.A.	ES
Orudis Retard 200 mg depotkapsel, hård	not available	11583	SANOFI AB	SE
Orudis Retard 200 mg depotkapsel, hård	not available	11583	SANOFI AB	SE
Oruvail 100 mg Prolonged-Release	not available	PL 04425/0597	AVENTIS PHARMA 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
Capsules, Hard				
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 100 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0597	AVENTIS PHARMA LTD	XI
Oruvail 200 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0599	AVENTIS PHARMA LTD	XI
Oruvail 200 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0599	AVENTIS PHARMA LTD	XI
Oruvail 200 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0599	AVENTIS PHARMA LTD	XI
Oruvail 200 mg Prolonged-Release Capsules, Hard	not available	PL 04425/0599	AVENTIS PHARMA LTD	XI
Oruvail 200 mg	not available	PL 04425/0599	AVENTIS PHARMA 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
Prolonged-Release Capsules, Hard				
Oruvail 200mg Prolonged-Release Capsules, Hard	not available	PA 540/119/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Oruvail 200mg Prolonged-Release Capsules, Hard	not available	PA 540/119/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
PROFEMIGR 150 mg, comprimé sécable	not available	34009 325 414 9 2	SANOFI-AVENTIS FRANCE	FR
PROFEMIGR 150 mg, comprimé sécable	not available	34009 558 980 7 8	SANOFI-AVENTIS FRANCE	FR
PROFEMIGR 150 mg, comprimé sécable	not available	34009 362 745 5 6	SANOFI-AVENTIS FRANCE	FR
PROFEMIGR 150 mg, comprimé sécable	not available	34009 325 413 2 4	SANOFI-AVENTIS FRANCE	FR
PROFEMIGR 150 mg, comprimé sécable	not available	3400937522496	SANOFI-AVENTIS FRANCE	FR
Profenid 100 mg Ampullen	not available	1-19749	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 100 mg Ampullen	not available	1-19749	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 100 mg comprimate filmate	not available	6448/2014/01	SANOFI ROMANIA SRL	RO
Profenid 100 mg Kapseln	not available	1-19745	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 100 mg Kapseln	not available	1-19745	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 100 mg Kapseln	not available	1-19745	SANOFI-AVENTIS GMBH OSTERREICH	AT
PROFENID 100 mg supositórios	not available	9424101	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
PROFENID 100 mg, comprimé pelliculé	not available	34009 557 345 6 7	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 mg, comprimé pelliculé	not available	34009 333 020 6 1	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 MG, POUDRE POUR	not available	557 466-8	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SOLUTION INJECTABLE (IV) EN FLACON				
PROFENID 100 MG, SUPPOSITOIRE	not available	34009 494 171 6 2	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 MG, SUPPOSITOIRE	not available	317 929-3	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 mg/2 ml solução injectável	not available	9424036	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
PROFENID 100 mg/2 ml, solution injectable (I.M.)	not available	34009 560 279 0 3	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 mg/2 ml, solution injectable (I.M.)	not available	34009 346 603 5 1	SANOFI-AVENTIS FRANCE	FR
PROFENID 100 mg/2 ml, solution injectable (I.M.)	not available	34009 342 921 2 5	SANOFI-AVENTIS FRANCE	FR
Profenid 50 mg Kapseln	not available	15.652	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 50 mg Kapseln	not available	15.652	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid 50 mg Konzentrat zur Infusionsbereitung	not available	1-20909	SANOFI-AVENTIS GMBH OSTERREICH	AT
PROFENID 50 mg, gélule	not available	34009 313 890 5 7	SANOFI-AVENTIS FRANCE	FR
PROFENID 50 mg, gélule	not available	34009 499 392 0 6	SANOFI-AVENTIS FRANCE	FR
PROFENID LP 200 mg, gélule à libération prolongée	not available	34009 331 583 3 0	SANOFI-AVENTIS FRANCE	FR
Profenid retard 200 mg Kapseln	not available	1-19744	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid retard 200 mg Kapseln	not available	1-19744	SANOFI-AVENTIS GMBH OSTERREICH	AT
Profenid, 100 mg, czopki	not available	R/0985	SANOFI-AVENTIS FRANCE	PL
Profenid, 100 mg, tabletki powlekane	not available	7607	SANOFI-AVENTIS FRANCE	PL
Rilies	not available	RVG 14148	NOVUM PHARMA BV	NL
Rilies	not available	RVG 14148	NOVUM PHARMA BV	NL
Rofenid Long Acting 200 mg, gélules à libération prolongée	not available	0197973	SANOFI 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
Rofenid Long Acting 200 mg, gélules à libération prolongée	not available	BE396487	SANOFI BELGIUM	BE
Rofenid Long Acting 200 mg, gélules à libération prolongée	not available	BE140095	SANOFI BELGIUM	BE
Rofenid Long Acting 200 mg, harde capsules met verlengde afgifte	not available	BE396487	SANOFI BELGIUM	BE
Rofenid Long Acting 200 mg, harde capsules met verlengde afgifte	not available	BE140095	SANOFI BELGIUM	BE
Rofenid Long Acting 200 mg, Hartkapseln, retardiert	not available	BE396487	SANOFI BELGIUM	BE
Rofenid Long Acting 200 mg, Hartkapseln, retardiert	not available	BE140095	SANOFI BELGIUM	BE
Rofenid Ready Mix i.m. 100 mg, Injektionslösung	not available	BE132063	SANOFI BELGIUM	BE
Rofenid Ready Mix I.M. 100 mg, oplossing voor injectie	not available	BE132063	SANOFI BELGIUM	BE
Rofenid Ready Mix I.M. 100 mg, solution injectable	not available	BE132063	SANOFI BELGIUM	BE
Rofenid Ready Mix I.M. 100 mg, solution injectable	not available	0100748	SANOFI BELGIUM	LU
TOPREC 1 MG/ML ENFANTS ET NOURISSONS, SIROP	not available	351 573-3	SANOFI-AVENTIS FRANCE	FR
TOPREC 25 mg, comprimé	not available	34009 561 167 1 3	SANOFI-AVENTIS FRANCE	FR
TOPREC 25 mg, comprimé	not available	34009 330 810 6 5	SANOFI-AVENTIS FRANCE	FR
Valket 200 Retard	not available	PL 11311/0460	TILLOMED LABORATORIES	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
			LTD	
БИ-ПРОФЕНИД 150 mg таблетки с изменено освобождаване	not available	9900097	SANOFI-AVENTIS GROUPE	BG
ПРОФЕНИД 100 mg, супозитории	not available	20030008	SANOFI-AVENTIS GROUPE	BG