29 May 2019
EMA/430529/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: diclofenac (systemic formulations

Procedure no.: PSUSA/00001048/201809

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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
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/08	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/07	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/03	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/04	SANDOZ HUNGÁRIA KFT	HU
DICLOREUM 150 mg capsule rigide a rilascio prolungato	not available	024515114	ALFASIGMA S.P.A.	IT
DICLOREUM 50 MG COMPRESSE GASTRORESISTENTI	not available	024515049	ALFASIGMA S.P.A.	IT
DICLOREUM 100 MG COMPRESSE A RILASCIO PROLUNGATO	not available	024515088	ALFASIGMA S.P.A.	IT
DICLOREUM 75 MG/3 ML SOLUZIONE INIETTABILE PER USO INTRAMUSCOLARE	not available	024515076	ALFASIGMA S.P.A.	IT
Voltaren 25, 25 mg cápsulas moles	not available	5204466	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, 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
Voltaren 25, 25 mg cápsulas moles	not available	5204508	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 25, 25 mg cápsulas moles	not available	5204458	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 25, 25 mg cápsulas moles	not available	5204474	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren Dolo 12,5 mg lágy kapszula	not available	OGYI-T-5572/21	GLAXOSMITHKLINE-CONSUMER KFT.	HU
VOLTAREN DOLO 12,5 MG LÁGY KAPSZULA	not available	OGYI-T-5572/22	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Akti 12,5 mg plevele dengtos tabletes	not available	LT/1/94/0948/003	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 12,5 mg plėvele dengtos tabletės	not available	LT/1/94/0948/003	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 12,5 mg plėvele dengtos tabletės	not available	LT/1/94/0948/003	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Actigo Extra 25 mg obalené tablety	not available	29/0227/01-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/06	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/04	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/05	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/23	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/25	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/24	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren K 25 mg, omhulde tabletten	not available	RVG 13244	GLAXOSMITHKLINE CONSUMER HEALTHCARE B.V	NL
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS 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
Voltaren T 25 mg mjuka kapslar.	not available	41904	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
Voltaren T 12,5 mg filmdragerade tabletter	not available	16081	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
Voltaren T 12,5 mg mjuka kapslar.	not available	42233	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
NOVAPIRINA 25 mg compresse rivestite	not available	024951028	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.P.A.	IT
Diclofenac 100 mg supozitoriji	not available	97-0175	GLAXOSMITHKLINE LATVIA SIA	LV
Diclofenac 50 mg supozitoriji	not available	97-0174	GLAXOSMITHKLINE LATVIA SIA	LV
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/001	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/002	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/003	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING 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
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/004	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	SI
DICLOREUM 50 MG GRANULATO PER SOLUZIONE ORALE	not available	024515138	ALFASIGMA S.P.A.	IT
Voltaren Acti-Go 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren Akti, 12,5 mg õhukese polümeerikattega tabletid	not available	434104	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	EE
Voltaren Akti, 12,5 mg õhukese polümeerikattega tabletid	not available	434104	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	EE
Voltaren Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250302	NOVARTIS (HELLAS) S.A.C.I.	GR
Dolotren 100 mg supositorios	not available	57.203	FAES FARMA, S.A.	ES
DICLAC 150 DUO, 150 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9578	SANDOZ GMBH	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
DICLAC 75 DUO, 75 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9577	SANDOZ GMBH	PL
DICLOREUM 150 mg capsule rigide a rilascio prolungato	not available	024515114	ALFASIGMA S.P.A.	IT
DICLOREUM 50 MG COMPRESSE GASTRORESISTENTI	not available	024515049	ALFASIGMA S.P.A.	IT
DICLOREUM 100 MG COMPRESSE A RILASCIO PROLUNGATO	not available	024515088	ALFASIGMA S.P.A.	IT
DICLOREUM 75 MG/3 ML SOLUZIONE INIETTABILE PER USO INTRAMUSCOLARE	not available	024515076	ALFASIGMA S.P.A.	IT
DICLOFENAC MYLAN 50 mg, comprimé gastro-résistant	not available	NL 16786	MYLAN S.A.S	FR
LUASE 50 mg comprimidos gastrorresistentes	not available	57.205	ALFASIGMA ESPAÑA, S.L.	ES
Voltaren K 12,5 mg, filmomhulde tabletten	not available	RVG 20982	GLAXOSMITHKLINE CONSUMER HEALTHCARE B.V	NL
DICLOREUM DOLORE 25 mg compresse gastroresistenti	not available	028618039	ALFASIGMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DICLOREUM DOLORE 25 mg compresse gastroresistenti	not available	028618027	ALFASIGMA S.P.A.	IT
Dicloreum 150 mg capsule cu eliberare prelungită	not available	8293/2015/01	ALFASIGMA S.P.A.	RO
Dicloreumdol 25 mg compresse rivestite con film	IT/H/0483/001	041735022	ALFASIGMA S.P.A.	IT
Dicloreumdol 25 mg compresse rivestite con film	IT/H/0483/001	041735010	ALFASIGMA S.P.A.	IT
Otriflu 12.5mg Film-coated Tablets	not available	PA 678/137/001	GLAXOSMITHKLINE CONSUMER HEALTHCARE (IRELAND) LTD	IE
Dolotren retard 100 mg cápsulas duras de liberación prolongada	not available	58.435	FAES FARMA, S.A.	ES
Diclofenac Zentiva 25 mg Filmtabletten	not available	74529.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Diclofenac Zentiva 25 mg Filmtabletten	not available	74529.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	381 408-0 OU 34009 381 408 0 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	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
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	381 409-7 OU 34009 381 409 7 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	355 324-8 OU 34009 355 324 8 0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	381 410-5 OU 34009 381 410 5 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	359 411-2 OU 34009 359 411 2 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENDOLO 12,5 mg, comprimé enrobé	not available	34009 355 325 4 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
AKIS 25 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/004	040528186	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/004	040528174	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/004	040528162	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/005	040528135	IBSA FARMACEUTICI ITALIA	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
AKIS 50 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/005	040528147	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/005	040528150	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/006	040528123	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/006	040528111	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/1 ml soluzione iniettabile in siringa preriempita	UK/H/3585/006	040528109	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	UK/H/3585/003	040528073	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	UK/H/3585/003	040528085	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	UK/H/3585/003	040528097	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	UK/H/3585/002	040528061	IBSA FARMACEUTICI ITALIA	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
AKIS 50 mg/ml soluzione iniettabile	UK/H/3585/002	040528059	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	UK/H/3585/002	040528046	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	UK/H/3585/001	040528010	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	UK/H/3585/001	040528022	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	UK/H/3585/001	040528034	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml Solution for Injection in prefilled syringe	UK/H/3585/004	PL 21039/0021	IBSA FARMACEUTICI ITALIA	UK
AKIS 50 mg/ml Solution for Injection in prefilled syringe	UK/H/3585/005	PL 21039/0023	IBSA FARMACEUTICI ITALIA	UK
AKIS 75 mg/ml Solution for Injection in prefilled syringe	UK/H/3585/006	PL 21039/0024	IBSA FARMACEUTICI ITALIA	UK
AKIS 25 mg/ml Solution for Injection	UK/H/3585/001	PL 21039/0018	IBSA FARMACEUTICI ITALIA	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
AKIS 50 mg/ml Solution for Injection	UK/H/3585/002	PL 21039/0019	IBSA FARMACEUTICI ITALIA	UK
AKIS 75 mg/ml Solution for Injection	UK/H/3585/003	PL 21039/0020	IBSA FARMACEUTICI ITALIA	UK
AKIS 25 mg/ml injekčný roztok	UK/H/3585/001	29/0188/13-S	IBSA SLOVAKIA S.R.O.	SK
AKIS 50 mg/ml injekcný roztok	UK/H/3585/002	29/0189/13-S	IBSA SLOVAKIA S.R.O.	SK
AKIS 75 mg/ml injekcný roztok	UK/H/3585/003	29/0190/13-S	IBSA SLOVAKIA S.R.O.	SK
Flector Rapid 25 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/004	OGYI-T-22385/10	IBSA PHARMA KFT	HU
Flector Rapid 25 mg/ml oldatos injekció eloretöltött fecskendoben	UK/H/3585/004	OGYI-T-22385/11	IBSA PHARMA KFT	HU
Flector Rapid 25 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/004	OGYI-T-22385/12	IBSA PHARMA KFT	HU
Flector Rapid 50 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/005	OGYI-T-22385/13	IBSA PHARMA 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
Flector Rapid 50 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/005	OGYI-T-22385/14	IBSA PHARMA KFT	HU
Flector Rapid 50 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/005	OGYI-T-22385/15	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/006	OGYI-T-22385/16	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/006	OGYI-T-22385/17	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/006	OGYI-T-22385/18	IBSA PHARMA KFT	HU
Flector Rapid 25 mg/ml oldatos injekció	UK/H/3585/001	OGYI-T-22385/01	IBSA PHARMA KFT	HU
Flector Rapid 25 mg/ml oldatos injekció	UK/H/3585/001	OGYI-T-22385/02	IBSA PHARMA KFT	HU
Flector Rapid 25 mg/ml oldatos injekció	UK/H/3585/001	OGYI-T-22385/03	IBSA PHARMA KFT	HU
Flector Rapid 50 mg/ml oldatos injekció	UK/H/3585/002	OGYI-T-22385/04	IBSA PHARMA 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
Flector Rapid 50 mg/ml oldatos injekció	UK/H/3585/002	OGYI-T-22385/05	IBSA PHARMA KFT	HU
Flector Rapid 50 mg/ml oldatos injekció	UK/H/3585/002	OGYI-T-22385/06	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció	UK/H/3585/003	OGYI-T-22385/07	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció	UK/H/3585/003	OGYI-T-22385/08	IBSA PHARMA KFT	HU
Flector Rapid 75 mg/ml oldatos injekció	UK/H/3585/003	OGYI-T-22385/09	IBSA PHARMA KFT	HU
AKIS, 25 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	UK/H/3585/010	24769	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	UK/H/3585/011	24770	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	UK/H/3585/012	24771	IBSA FARMACEUTICI ITALIA	PL
AKIS, 25 mg/ml, roztwór do wstrzykiwań	UK/H/3585/007	24766	IBSA FARMACEUTICI ITALIA	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
AKIS, 50 mg/ml, roztwór do wstrzykiwań	UK/H/3585/008	24767	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań	UK/H/3585/009	24768	IBSA FARMACEUTICI ITALIA	PL
FLECTORFLAM 50 mg granulato per soluzione orale	not available	036058030	IBSA FARMACEUTICI ITALIA	IT
DICLOIN 75 mg/ml ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	UK/H/3585/006	75819/2-10-2013	IBSA FARMACEUTICI ITALIA	GR
DICLOIN 50 mg/ml ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	UK/H/3585/005	75818/2-10-2013	IBSA FARMACEUTICI ITALIA	GR
DICLOIN 25 mg/ml ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	UK/H/3585/004	75817/2-10-2013	IBSA FARMACEUTICI ITALIA	GR
DICLOIN 25 mg/ml injekční roztok	UK/H/3585/001	29/234/13-C	IBSA SLOVAKIA S.R.O.	CZ
DICLOIN 50 mg/ml injekční roztok	UK/H/3585/002	29/235/13-C	IBSA SLOVAKIA S.R.O.	CZ
DICLOIN 75 mg/ml injekční roztok	UK/H/3585/003	29/236/13-C	IBSA SLOVAKIA 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
DICLOIN 25 mg/ml injekční roztok v předplněné injekční stříkačce	UK/H/3585/004	29/237/13-C	IBSA SLOVAKIA S.R.O.	CZ
DICLOIN 50 mg/ml injekční roztok v předplněné injekční stříkačce	UK/H/3585/005	29/238/13-C	IBSA SLOVAKIA S.R.O.	CZ
DICLOIN 75 mg/ml injekční roztok v předplněné injekční stříkačce	UK/H/3585/006	29/239/13-C	IBSA SLOVAKIA S.R.O.	CZ
AKIS 25 mg/ml injekcný roztok v naplnenej injekcnej striekacke	UK/H/3585/004	29/0185/13-S	IBSA SLOVAKIA S.R.O.	SK
AKIS 50 mg/ml injekcný roztok v naplnenej injekcnej striekacke	UK/H/3585/005	29/0186/13-S	IBSA SLOVAKIA S.R.O.	SK
AKIS 75 mg/ml injekcný roztok v naplnenej injekcnej striekacke	UK/H/3585/006	29/0187/13-S	IBSA SLOVAKIA S.R.O.	SK
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973079	ALTERGON ITALIA S.R.L.	IT
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973081	ALTERGON 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
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973093	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973131	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973143	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973156	ALTERGON ITALIA S.R.L.	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617037	IBSA FARMACEUTICI ITALIA	IT
DICLOIN 25 mg/ml ενέσιμο διάλυμα	UK/H/3585/001	75814/2-10-2013	IBSA FARMACEUTICI ITALIA	GR
DICLOIN 50 mg/ml ενέσιμο διάλυμα	UK/H/3585/002	75815/2-10-2013	IBSA FARMACEUTICI ITALIA	GR
DICLOIN 75 mg/ml ενέσιμο διάλυμα	UK/H/3585/003	75816/2-10-2013	IBSA FARMACEUTICI ITALIA	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
AKIS 50 mg/ml, solution injectable	UK/H/3585/002	34009 300 350 8 5	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	UK/H/3585/002	34009 300 350 9 2	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	UK/H/3585/002	34009 300 351 0 8	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	UK/H/3585/003	34009 300 351 1 5	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	UK/H/3585/003	34009 300 351 2 2	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	UK/H/3585/003	34009 550 124 5 0	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	UK/H/3585/001	34009 300 350 4 7	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	UK/H/3585/001	34009 300 350 5 4	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	UK/H/3585/001	34009 300 350 6 1	LABORATOIRES GENEVRIER	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
AKIS 75 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/006	34009 300 352 0 7	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/006	34009 300 352 1 4	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/006	34009 550 124 6 7	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/005	34009 300 351 7 7	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/005	34009 300 351 8 4	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/005	34009 300 351 9 1	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/004	34009 300 351 4 6	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/004	34009 300 351 5 3	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue pré-remplie	UK/H/3585/004	34009 300 351 6 0	LABORATOIRES GENEVRIER	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
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973042	ALTERGON ITALIA S.R.L.	IT
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973055	ALTERGON ITALIA S.R.L.	IT
INFORCE 25 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973067	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973105	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973117	ALTERGON ITALIA S.R.L.	IT
INFORCE 50 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973129	ALTERGON ITALIA S.R.L.	IT
Diclovex 25 mg Weichkapseln	not available	96490.00.00	IBSA FARMACEUTICI ITALIA	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
Diclovex 50 mg Weichkapseln	not available	96491.00.00	IBSA FARMACEUTICI ITALIA	DE
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617052	IBSA FARMACEUTICI ITALIA	IT
Flector RAPID 50 mg granulátum	not available	OGYI-T-05033/03	IBSA PHARMA KFT	HU
Flector RAPID 50 mg granulátum	not available	OGYI-T-05033/04	IBSA PHARMA KFT	HU
Flector EP Rapid 50 mg granulát	not available	29/0906/96-S	IBSA SLOVAKIA S.R.O.	SK
FLECTOR EP RAPID 50 mg granule pro perorální roztok v sáčku	not available	29/514/99-C	IBSA SLOVAKIA S.R.O.	CZ
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 615 1 9	LABORATOIRES GENEVRIER	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 641 2 1	LABORATOIRES GENEVRIER	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 616 8 7	LABORATOIRES GENEVRIER	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
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 617 4 8	LABORATOIRES GENEVRIER	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 642 9 9	LABORATOIRES GENEVRIER	FR
FLECTOR 50 mg, comprimé	not available	34009 301 460 6 4	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml Solution for Injection	UK/H/3585/007	PL 21039/0040	IBSA FARMACEUTICI ITALIA	UK
AKIS 50 mg/ml Solution for Injection	UK/H/3585/008	PL 21039/0041	IBSA FARMACEUTICI ITALIA	UK
AKIS 75 mg/ml Solution for Injection	UK/H/3585/009	PL 21039/0042	IBSA FARMACEUTICI ITALIA	UK
AKIS 25 mg/ml solution for injection in prefilled syringe	UK/H/3585/010	PL 21039/0043	IBSA FARMACEUTICI ITALIA	UK
AKIS 50 mg/ml solution for injection in prefilled syringe	UK/H/3585/011	PL 21039/0044	IBSA FARMACEUTICI ITALIA	UK
AKIS 75 mg/ml solution for injection in prefilled syringe	UK/H/3585/012	PL 21039/0045	IBSA FARMACEUTICI ITALIA	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
AKIS Plus 25 mg/ml injekčný roztok naplnený v injekčnej striekačke	UK/H/3585/010	29/0331/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 25 mg/ml injekčný roztok	UK/H/3585/007	29/0328/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok	UK/H/3585/008	29/0329/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok	UK/H/3585/009	29/0330/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok naplnený v injekčnej striekačke	UK/H/3585/011	29/0332/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok naplnený v injekčnej striekačke	UK/H/3585/012	29/0333/17-S	IBSA SLOVAKIA S.R.O.	SK
Flector Rapiven 25 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/010	OGYI-T-23318/10	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció előretöltött fecskendőben	UK/H/3585/011	OGYI-T-23318/11	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció eloretöltött fecskendoben	UK/H/3585/012	OGYI-T-23318/12	IBSA PHARMA 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
Flector Rapiven 25 mg/ml oldatos injekció	UK/H/3585/007	OGYI-T-23318/01	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	UK/H/3585/007	OGYI-T-23318/02	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	UK/H/3585/007	OGYI-T-23318/03	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	UK/H/3585/008	OGYI-T-23318/04	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	UK/H/3585/008	OGYI-T-23318/05	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	UK/H/3585/008	OGYI-T-23318/06	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	UK/H/3585/009	OGYI-T-23318/07	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	UK/H/3585/009	OGYI-T-23318/08	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	UK/H/3585/009	OGYI-T-23318/09	IBSA PHARMA 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
Dicloalgan 12,5 mg lágy kapszula	UK/H/6105/001	OGYI-T-23250/04	IBSA PHARMA KFT	HU
Dicloalgan 12,5 mg lágy kapszula	UK/H/6105/001	OGYI-T-23250/01	IBSA PHARMA KFT	HU
Dicloalgan 12,5 mg lágy kapszula	UK/H/6105/001	OGYI-T-23250/03	IBSA PHARMA KFT	HU
Dicloalgan 12,5 mg lágy kapszula	UK/H/6105/001	OGYI-T-23250/02	IBSA PHARMA KFT	HU
DAHILO 12,5 mg capsule molli	UK/H/6105/001	044608014	IBSA FARMACEUTICI ITALIA	IT
DAHILO 12,5 mg capsule molli	UK/H/6105/001	044608026	IBSA FARMACEUTICI ITALIA	IT
DAHILO 12,5 mg capsule molli	UK/H/6105/001	044608038	IBSA FARMACEUTICI ITALIA	IT
DAHILO 12,5 mg capsule molli	UK/H/6105/001	044608040	IBSA FARMACEUTICI ITALIA	IT
INFORCE 75 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973028	ALTERGON 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
INFORCE 75 mg/ml soluzione iniettabile per uso intramuscolare e sottocutaneo	not available	036973030	ALTERGON ITALIA S.R.L.	IT
Diclofenac Tablets BP 50mg	not available	PL 20416/0220	CRESCENT PHARMA LIMITED	UK
Diclofenac Tablets BP 25mg	not available	PL 20416/0219	CRESCENT PHARMA LIMITED	UK
FLOGOFENAC 100 mg capsule rigide a rilascio prolungato	not available	025536020	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
LUASE 50 mg comprimidos gastrorresistentes	not available	57.205	ALFASIGMA ESPAÑA, S.L.	ES
Diclofenac akut 1A Pharma 50 mg - lösbare Tabletten	not available	1-24778	1A PHARMA GMBH	AT
Voltaren T 25 mg mjuka kapslar.	not available	41904	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
Voltaren Rapid 25 mg mäkké kapsuly	not available	29/0709/10-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Diclofenac potassium NCH 12.5mg Film-coated Tablets	not available	PA 678/129/001	GLAXOSMITHKLINE CONSUMER HEALTHCARE (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
Diclac 150 mg retard tabletta	not available	OGYI-T-4000/05	SANDOZ HUNGÁRIA KFT	HU
Diclac 150 mg retard tabletta	not available	OGYI-T-4000/06	SANDOZ HUNGÁRIA KFT	HU
ВОЛТАРЕН ДОЛО 12.5 mg Филмирани таблетки	not available	20030609	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	BG
Diclofenac Potassium 25 mg Tablets	not available	PL 20075/0682	ACCORD HEALTHCARE LIMITED	UK
Diclofenac Potassium 50 mg Tablets	not available	PL 20075/0683	ACCORD HEALTHCARE LIMITED	UK
DICLOREUM 50 MG GRANULATO PER SOLUZIONE ORALE	not available	024515138	ALFASIGMA S.P.A.	IT
DICLOREUM 50, čípky	not available	29/126/94-A/C	ALFASIGMA S.P.A.	CZ
DICLOREUM 100, čípky	not available	29/126/94-B/C	ALFASIGMA S.P.A.	CZ
Voltaren Dolo 25 mg überzogene Tablette	not available	64958.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	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
Voltaren Dolo 25 mg überzogene Tablette	not available	43151.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Express, 12,5 mg, kapsułki, miękkie	not available	16009	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltarol 12,5 mg kapsel, myk	not available	09-6597	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
EMINOCS 50 mg/ml, drank	NL/H/0800/001	RVG 31532	ALFASIGMA S.P.A.	NL
EMINOCS 50 mg/ml soluzione orale	NL/H/0800/001	038049033	ALFASIGMA S.P.A.	IT
EMINOCS 50 mg/ml soluzione orale	NL/H/0800/001	038049019	ALFASIGMA S.P.A.	IT
EMINOCS 50 mg/ml soluzione orale	NL/H/0800/001	038049045	ALFASIGMA S.P.A.	IT
EMINOCS 50 mg/ml soluzione orale	NL/H/0800/001	038049021	ALFASIGMA S.P.A.	IT
Voltaren Express Forte, 25 mg, kapsułki miękkie	not available	17890	GLAXOSMITHKLINE CONSUMER HEALTHCARE 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
Voltaren Acti Forte, 25 mg, tabletki powlekane	not available	16489	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren Dolo Liquid 25 mg Weichkapsel	not available	82502.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
DICLOFENAC HIKMA 75 mg	not available	5200.00.01	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
DICLOFENAC Alfasigma 50 mg compresse	not available	033612033	ALFASIGMA S.P.A.	IT
DICLOFENAC Alfasigma 150 mg capsule a rilascio prolungato	not available	033612058	ALFASIGMA S.P.A.	IT
DICLOFENAC Alfasigma 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	033612096	ALFASIGMA S.P.A.	IT
Voltaren Acti, 12,5 mg, tabletki powlekane	not available	9462	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltarol 25 mg kapsler, myke	not available	10-8136	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
Voltaren Dolo 12,5 mg filmtabletta	not available	OGYI-T-5572/08	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
Voltaren Dolo 12,5 mg filmtabletta	not available	OGYI-T-5572/09	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltarol 12,5 mg tabletter, filmdrasjerte	not available	99-7996	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
Dolotren 50 mg comprimidos gastrorresistentes	not available	57.202	FAES FARMA, S.A.	ES
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS 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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS 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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS 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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS 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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Voltaren Akti 25 mg mīkstās kapsulas	not available	12-0185	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LV
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/09	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/10	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/11	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
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/12	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/13	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/14	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltarol 25 mg drasjerte tabletter	not available	09-1063	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618015	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618041	ALFASIGMA S.P.A.	IT
Voltaren Akti 12,5 mg apvalkotās tabletes	not available	04 – 0291	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LV
Cataflam 50 mg, überzogene Tabletten	not available	0010/01/10/7175	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg magensaftresistente Tabletten	not available	0010/02/10/0205	NOVARTIS PHARMA N.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
Voltaren Retard 100 mg Retardtabletten	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg Zäpfchen	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg suppositoires	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg comprimés à libération prolongée	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam Dispersible 46,5 mg, Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	BE172514	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg maagsapresistente tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS PHARMA GMBH	BG
Cataflam 50 mg lágy kapszula	not available	OGYI-T-5573/07	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Cataflam 50 mg lágy kapszula	not available	OGYI-T-5573/08	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Voltaren Rapid 50 mg cápsulas moles	not available	5668314	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Rapid 50 mg cápsulas moles	not available	5668322	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Acti-Go® Επικαλυμμένα Δισκία , 25mg	not available	251250101	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250302	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren Acti-Go 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	251250201	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTFAST 50 mg capsule molli	not available	028945044	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg capsule molli	not available	028945057	NOVARTIS FARMA S.P.A.	IT
Voltaren® fast	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren® fast	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	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
Voltaren® fast	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
VOLTAREN®	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880101	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	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
VOLTAREN®	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880601	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN®	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTFAST 25 mg compresse rivestite	not available	028945018	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite	not available	028945020	NOVARTIS FARMA 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
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTFAST® 50 mg jauhe oraaliliuosta varten	not available	20484	NOVARTIS FINLAND OY	FI
Voltfast 50 mg pulver till oral lösning	not available	20484	NOVARTIS FINLAND OY	FI
Voltaren Rapid 25 mg dragerade tabletter	not available	10202	NOVARTIS FINLAND OY	FI
Voltaren Rapid 25 mg tabletti, päällystetty	not available	10202	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg tabletti, päällystetty	not available	10203	NOVARTIS FINLAND OY	FI
Dolo-Voltarén 46, 5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 100 mg peräpuikko	not available	8090	NOVARTIS FINLAND 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
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 25 mg enterotabletter	not available	7442	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren, 75 mg/3 ml süstelahus	not available	061694	SIA "NOVARTIS BALTICS"	EE
Voltaren 25 mg enterotabletti	not available	7442	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND 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
Voltaren 50 mg enterotabletti	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 75 mg/3 ml injekcinis/infuzinis tirpalas	not available	LT/1/94/1300/005	SIA "NOVARTIS BALTICS"	LT
Voltaren 75 mg/3 ml šķīdums injekcijām	not available	00-0517	SIA "NOVARTIS BALTICS"	LV
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg ilgstošās darbības apvalkotās tabletes	not available	94-0182	SIA "NOVARTIS BALTICS"	LV
Voltaren Retard 100 mg pailginto atpalaidavimo tabletės	not available	LT/1/94/1300/002	SIA "NOVARTIS BALTICS"	LT
Voltaren Retard 100 mg depottabletti	not available	9228	NOVARTIS FINLAND 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
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletti	not available	10919	NOVARTIS FINLAND OY	FI
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Cataflam 50, 50 mg, tabletki powlekane	not available	R/3707	NOVARTIS POLAND SP. Z O. O.	PL
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
VOLTAREN® K MIGRÄNE 50 MG Überzogene Tabletten	not available	43151.01.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium)	not available	17982.00.00	NOVARTIS 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
Hartkapseln				
ВОЛТФАСТ 50 MG ПРАХ ЗА ПЕРОРАЛЕН РАЗТВОР	not available	20060381	NOVARTIS PHARMA GMBH	BG
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS PHARMA GMBH	BG
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS PHARMA GMBH	BG
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Magensaftresistente Tabletten	not available	2502.00.01	NOVARTIS 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
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	BG
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren, 25 mg/1 ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
VOLTAREN 25 mg comprimate gastrorezistente	not available	4715/2004/01	NOVARTIS PHARMA GMBH	RO
Voltaren® 50 mg Magensaftresistente Tabletten	not available	14651.00.00	NOVARTIS PHARMA GMBH	DE
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS PHARMA GMBH	BG
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Voltaren, 50 mg, tabletki dojelitowe	not available	R/1204	NOVARTIS POLAND 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
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
VOLTAREN 50 mg comprimate gastrorezistente	not available	4716/2004/01	NOVARTIS PHARMA GMBH	RO
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Волтарен 75 mg/3 ml инжекционен разтвор	not available	20020741	NOVARTIS PHARMA GMBH	BG
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND 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
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE
CATAFLAM TABLETS 50MG	not available	18492	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Cataflam 1,5% σταγόνες	not available	18493	NOVARTIS PHARMACEUTICALS UK LIMITED	CY
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren T 50 mg, dragerade tabletter	not available	11242	NOVARTIS SVERIGE AB	SE
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA 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
Voltaren 100 mg suppositoires	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg zetpillen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren ® 25 mg - Filmtabletten	not available	1-15554	NOVARTIS PHARMA GMBH	AT
Voltaren 25 mg enterotabletter	not available	09737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren 50 mg comprimés gastro-résistants	not available	0010/02/10/0205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg comprimés gastro-résistants	not available	BE121116	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 50 mg enterotabletter	not available	09738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg maagsapresistente tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/ 3 ml oplossing voor injectie	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
Voltaren 75 mg/3 ml solution injectable	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren® dispers - lösliche Tabletten	not available	1-20832	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren Retard 100 mg comprimés à libération prolongée	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte	not available	BE122071	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Retard 75 mg comprimés à libération prolongée	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren suppositorier 100 mg	not available	09818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	09941	NOVARTIS SVERIGE AB	SE
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	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
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren® Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
VOLTARENE 100 mg, suppositoire	not available	3400932214341	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg ENFANT, suppositoire	not available	3400932239535	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg ENFANT, suppositoire	not available	3400932273591	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	3400931895299	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	3400931978718	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	3400933814458	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	3400933814519	NOVARTIS PHARMA S.A.S.	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
VOLTARENE 50 MG, comprimé enrobé gastro-résistant	not available	3400932351176	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE® 75 mg/3 ml, solution injectable	not available	3400932452224	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE® 75 mg/3 ml, solution injectable	not available	3400955283614	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE® 75 mg/3 ml, solution injectable	not available	3400955549444	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	3400933591960	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	3400933592042	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	3400933593452	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	3400933593513	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	3400934595615	NOVARTIS PHARMA S.A.S.	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
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	3400932460496	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	3400932460557	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	3400933070120	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	3400933070298	NOVARTIS PHARMA S.A.S.	FR
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Voltarol® Suppositories 12.5mg	not available	PL 00101/0472	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Rapid 50 mg Obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
VOLTAREN RAPID 50 mg TABLETY obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN injekční roztok	not available	29/186/80-C	NOVARTIS S.R.O.,	CZ
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
VOLTAREN 50 enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
VOLTAREN RETARD tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
CATAFLAM 50 mg dosepulver til mikstur, oppløsning	not available	05-3486	NOVARTIS NORGE AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cataflam 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren® 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS PHARMA GMBH	SI
Voltaren® 25 mg svečke	not available	H/94/01506/002	NOVARTIS PHARMA GMBH	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
Voltaren rapid 50 mg obložene tablete	not available	HR-H-090999073	NOVARTIS HRVATSKA D.O.O.	HR
VOLTAREN® D	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN®12.5 mg suppositories	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® 50 mg suppositories	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® 100 mg suppositories	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
Catafast® 50 mg powder for oral solution	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
CATAFLAM® 25 mg sugar-coated tablets	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM® 50 mg sugar-coated tablets	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® 25 mg gastro-resistant tablets	not available	MA1249/00704	NOVARTIS IRELAND 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
VOLTAREN® 50 mg gastro-resistant tablets	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® 75mg/3mL solution for injection	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN® SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
Cataflam 50 mg, comprimés enrobés	not available	0010/01/10/7175	NOVARTIS PHARMA N.V.	LU
Cataflam 50mg coated tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets	not available	PA0896/034/003	NOVARTIS 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
Voltaren Retard, 100 mg toimeainet prolongeeritult vabastavad tabletid	not available	061294	SIA "NOVARTIS BALTICS"	EE
Cataflam Dispersible 46,5 mg, comprimés dispersibles	not available	BE172514	NOVARTIS PHARMA N.V.	BE
Cataflam Dispersible 46,5 mg, dispergeerbare tabletten	not available	BE172514	NOVARTIS PHARMA N.V.	BE
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Diclofenac Potassium 50 mg Tablets	not available	PL 20046/0078	FOCUS PHARMACEUTICALS LIMITED	UK
Adacium Rapid 50mg Tablets	not available	PL 20046/0078	FOCUS PHARMACEUTICALS LIMITED	UK
Motifene 75 mg gélules à libération modifiée	not available	BE 176671	DAIICHI SANKYO BELGIUM S.A	BE
Motifene 75 mg, capsules met gereguleerde afgifte, hard	not available	BE 176671	DAIICHI SANKYO BELGIUM S.A	BE
Motifene 75 mg gélules à libération modifiée.	not available	0345531	DAIICHI SANKYO BELGIUM 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
Motifene 75 mg gélules à libération modifiée.	not available	0235011	DAIICHI SANKYO BELGIUM S.A	LU
Motifene 75 mg Capsules	not available	PL 08265/0003	DAIICHI SANKYO UK LTD	UK
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/005	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/005	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/004	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/004	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Acti-Go® Επικαλυμμένα Δισκία , 25mg	not available	251250101	NOVARTIS (HELLAS) S.A.C.I.	GR
Diclo-Divido long Hartkapseln, retardiert	not available	13847.00.00	PUREN PHARMA GMBH & CO. KG	DE
Diclo-Divido: Hartkapseln, retardiert Wirkstoff: Diclofenac-Natrium 75 mg	not available	13847.01.00	PUREN PHARMA GMBH & CO. KG	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
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618015	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618041	ALFASIGMA S.P.A.	IT
VOLTAREN ACTIGO EXTRA obalené tablety	not available	29/549/00-C	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	CZ
Itamifast 25 mg compresse rivestite con film	IT/H/0352/001	041736012	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 50 mg compresse rivestite con film	IT/H/0352/002	041736036	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 25 mg compresse rivestite con film	IT/H/0352/001	041736024	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 50 mg compresse rivestite con film	IT/H/0352/002	041736048	FIDIA FARMACEUTICI S.P.A	IT
Voltaren Rapid 25 mg měkké tobolky	not available	29/173/11-C	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	CZ
Dolotren 75 mg solución inyectable	not available	57.204	FAES FARMA, 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
DICLOREUM 50, čípky	not available	29/126/94-A/C	ALFASIGMA S.P.A.	CZ
DICLOREUM 100, čípky	not available	29/126/94-B/C	ALFASIGMA S.P.A.	CZ
ZEROFLOG 0,074 g/100 ml collutorio, 1 flacone	not available	034373 011	VALEAS S.P.A.	IT
ZEROFLOG 0,011 g/15 ml collutorio, 12 bustine	not available	034373 023	VALEAS S.P.A.	IT
ZEROFLOG 0,022 g/15 ml soluzione spray per mucosa orale	not available	034373 035	VALEAS S.P.A.	IT
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/30	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/29	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/26	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/31	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
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/28	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Dolo 25 mg lágy kapszula	not available	OGYI-T-5572/27	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Dolotren 46,5 mg comprimidos dispersables	not available	60.068	FAES FARMA, S.A.	ES
Diclomax Retard	not available	PL 27827/0005	GALEN LIMITED	UK
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5047246	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5047253	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5185533	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	3852589	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, 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
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	3852688	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5185525	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Rhumalgan SR 75 mg Modified Release Capsules	not available	PL 04416/0737	SANDOZ LTD	UK
Rhumalgan XL 100 mg modified-release capsules	not available	PL 04416/0738	SANDOZ LTD	UK
Diclofenac Tablets 25 mg	not available	PL 04416/0361	SANDOZ LTD	UK
® Diclovol 75mg SR Tablets	not available	PL 04416/0645	SANDOZ LTD	UK
Rhumalgan CR 75	not available	PL 4416/0242	SANDOZ LTD	UK
Rhumalgan CR 100	not available	PL 04416/0243	SANDOZ LTD	UK
Diclac 75mg Prolonged-release Tablets	not available	PA0711/009/003	ROWEX 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
Diclac Retard 100mg Prolonged-release Tablets	not available	PA0711/009/006	ROWEX LTD	IE
Voltaren Akti 25 mg dengtos tabletės	not available	LT/1/94/0948/001	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
Voltaren Akti 25 mg dengtos tabletės	not available	LT/1/94/0948/001	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	LT
DICLOREUM DOLORE 25 mg compresse gastroresistenti	not available	028618039	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25 mg compresse gastroresistenti	not available	028618027	ALFASIGMA S.P.A.	IT
Dicloreum 150 mg capsule cu eliberare prelungită	not available	8293/2015/01	ALFASIGMA S.P.A.	RO
Voltaren Dolo 12,5 mg Filmtabletten	not available	51218.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
DICLOFENAC Alfasigma 50 mg compresse	not available	033612033	ALFASIGMA S.P.A.	IT
DICLOFENAC Alfasigma 150 mg capsule a rilascio prolungato	not available	033612058	ALFASIGMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DICLOFENAC Alfasigma 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	033612096	ALFASIGMA S.P.A.	IT