23 June 2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): diclofenac (systemic formulations)

Procedure No. PSUSA/00001048/202109

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Adacium Rapid 50mg Tablets	not available	PL 20046/0078	FOCUS PHARMACEUTICALS LIMITED	XI
Akis 25 mg solución inyectable	HU/H/0796/001	83367	IBSA FARMACEUTICI ITALIA	ES
Akis 25 mg solución inyectable	HU/H/0796/001	83367	IBSA FARMACEUTICI ITALIA	ES
Akis 25 mg solución inyectable en jeringa precargada	HU/H/0796/004	83369	IBSA FARMACEUTICI ITALIA	ES
Akis 25 mg solución inyectable en jeringa precargada	HU/H/0796/004	83369	IBSA FARMACEUTICI ITALIA	ES
AKIS 25 mg/ml Solution for Injection	HU/H/0796/001	PL 21039/0040	IBSA FARMACEUTICI ITALIA	XI
AKIS 25 mg/ml Solution for Injection	HU/H/0796/001	PL 21039/0040	IBSA FARMACEUTICI ITALIA	XI
AKIS 25 mg/ml solution for injection in prefilled syringe	HU/H/0796/004	PL 21039/0043	IBSA FARMACEUTICI ITALIA	XI
AKIS 25 mg/ml solution for injection in prefilled syringe	HU/H/0796/004	PL 21039/0043	IBSA FARMACEUTICI ITALIA	XI
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528198	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528200	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528212	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528198	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528200	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528212	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile in siringa	HU/H/0796/004	040528287	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
preriempita				
AKIS 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528299	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528301	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528287	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528299	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528301	IBSA FARMACEUTICI ITALIA	IT
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 2 7	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 3 4	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 4 1	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 2 7	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 3 4	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 4 1	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 2 6	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 3 3	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 4 0	LABORATOIRES GENEVRIER	FR

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AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 2 6	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 3 3	LABORATOIRES GENEVRIER	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 4 0	LABORATOIRES GENEVRIER	FR
Akis 50 mg solución inyectable	HU/H/0796/002	83368	IBSA FARMACEUTICI ITALIA	ES
Akis 50 mg solución inyectable	HU/H/0796/002	83368	IBSA FARMACEUTICI ITALIA	ES
Akis 50 mg solución inyectable en jeringa precargada	HU/H/0796/005	83370	IBSA FARMACEUTICI ITALIA	ES
Akis 50 mg solución inyectable en jeringa precargada	HU/H/0796/005	83370	IBSA FARMACEUTICI ITALIA	ES
AKIS 50 mg/ml Solution for Injection	HU/H/0796/002	PL 21039/0041	IBSA FARMACEUTICI ITALIA	XI
AKIS 50 mg/ml Solution for Injection	HU/H/0796/002	PL 21039/0041	IBSA FARMACEUTICI ITALIA	XI
AKIS 50 mg/ml solution for injection in prefilled syringe	HU/H/0796/005	PL 21039/0044	IBSA FARMACEUTICI ITALIA	XI
AKIS 50 mg/ml solution for injection in prefilled syringe	HU/H/0796/005	PL 21039/0044	IBSA FARMACEUTICI ITALIA	XI
AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528224	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528236	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528248	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528224	IBSA FARMACEUTICI ITALIA	IT

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AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528236	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528248	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528313	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528325	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528337	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528313	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528325	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528337	IBSA FARMACEUTICI ITALIA	IT
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 5 8	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 6 5	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 7 2	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 5 8	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 6 5	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 7 2	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 5 7	LABORATOIRES GENEVRIER	FR

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AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 8 8	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 6 4	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 5 7	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 8 8	LABORATOIRES GENEVRIER	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 6 4	LABORATOIRES GENEVRIER	FR
Akis 75 mg solución inyectable	HU/H/0796/003	83371	IBSA FARMACEUTICI ITALIA	ES
Akis 75 mg solución inyectable	HU/H/0796/003	83371	IBSA FARMACEUTICI ITALIA	ES
Akis 75 mg solución inyectable en jeringa precargada	HU/H/0796/006	83372	IBSA FARMACEUTICI ITALIA	ES
Akis 75 mg solución inyectable en jeringa precargada	HU/H/0796/006	83372	IBSA FARMACEUTICI ITALIA	ES
AKIS 75 mg/ml Solution for Injection	HU/H/0796/003	PL 21039/0042	IBSA FARMACEUTICI ITALIA	XI
AKIS 75 mg/ml Solution for Injection	HU/H/0796/003	PL 21039/0042	IBSA FARMACEUTICI ITALIA	XI
AKIS 75 mg/ml solution for injection in prefilled syringe	HU/H/0796/006	PL 21039/0045	IBSA FARMACEUTICI ITALIA	XI
AKIS 75 mg/ml solution for injection in prefilled syringe	HU/H/0796/006	PL 21039/0045	IBSA FARMACEUTICI ITALIA	XI
AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528251	IBSA FARMACEUTICI ITALIA	IT

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AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528263	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528275	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528251	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528263	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528275	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528349	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528352	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528364	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528349	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528352	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528364	IBSA FARMACEUTICI ITALIA	IT
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 865 8 9	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 865 9 6	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 866 0 2	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 865 8 9	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution	HU/H/0796/003	34009 301 865 9 6	LABORATOIRES GENEVRIER	FR

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injectable				
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 866 0 2	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 866 9 5	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 0 1	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 1 8	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 866 9 5	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 0 1	LABORATOIRES GENEVRIER	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 1 8	LABORATOIRES GENEVRIER	FR
AKIS Plus 25 mg/ml injekčný roztok	HU/H/0796/001	29/0328/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 25 mg/ml injekčný roztok	HU/H/0796/001	29/0328/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 25 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/004	29/0331/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 25 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/004	29/0331/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok	HU/H/0796/002	29/0329/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok	HU/H/0796/002	29/0329/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok naplnený	HU/H/0796/005	29/0332/17-S	IBSA SLOVAKIA S.R.O.	SK

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v injekčnej striekačke				
AKIS Plus 50 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/005	29/0332/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok	HU/H/0796/003	29/0330/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok	HU/H/0796/003	29/0330/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/006	29/0333/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/006	29/0333/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS, 25 mg/ml, roztwór do wstrzykiwań	HU/H/0796/001	24766	IBSA FARMACEUTICI ITALIA	PL
AKIS, 25 mg/ml, roztwór do wstrzykiwań	HU/H/0796/001	24766	IBSA FARMACEUTICI ITALIA	PL
AKIS, 25 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/004	24769	IBSA FARMACEUTICI ITALIA	PL
AKIS, 25 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/004	24769	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań	HU/H/0796/002	24767	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań	HU/H/0796/002	24767	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/005	24770	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/005	24770	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań	HU/H/0796/003	24768	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór	HU/H/0796/003	24768	IBSA FARMACEUTICI ITALIA	PL

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do wstrzykiwań				
AKIS, 75 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/006	24771	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/006	24771	IBSA FARMACEUTICI ITALIA	PL
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
CATAFLAM 50 mg dosepulver til mikstur, oppløsning	not available	05-3486	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
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 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50, 50 mg, tabletki powlekane	not available	R/3707	NOVARTIS POLAND SP. Z O. O.	PL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50mg coated tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
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
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/09	GLAXOSMITHKLINE-CONSUMER KFT.	HU

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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
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
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	HU
DEALGIC 100 mg capsule rigide a rilascio prolungato	not available	028943025	FARMA GROUP S.R.L.	IT
DEALGIC 75 mg capsule rigide a rilascio prolungato	not available	028943013	FARMA GROUP S.R.L.	IT
DICLAC 150 DUO, 150 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9578	SANDOZ GMBH	PL
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
DICLAC 75 DUO, 75 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9577	SANDOZ GMBH	PL
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	not available	OGYI-T-4000/03	SANDOZ HUNGÁRIA KFT	HU

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tabletta				
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/04	SANDOZ HUNGÁRIA KFT	HU
Diclac 75mg Prolonged-release Tablets	not available	PA0711/009/003	ROWEX LTD	IE
Diclo-Divido long: Hartkapseln, retardiert Wirkstoff: Diclofenac-Natrium 100 mg	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
Diclofenac akut 1A Pharma 50 mg - lösbare Tabletten	not available	1-24778	1A PHARMA GMBH	AT
DICLOFENAC MYLAN 50 mg, comprimé gastro-résistant	not available	NL 16786	MYLAN S.A.S	FR
Diclofenac potassium 12.5mg tablets	not available	PL 36687/0214	TORRENT PHARMA (UK) LTD.	XI
Diclofenac Potassium 25 mg Tablets	not available	PL 20075/0682	ACCORD HEALTHCARE LIMITED	XI
Diclofenac Potassium 50 mg Tablets	not available	PL 20046/0078	FOCUS PHARMACEUTICALS LIMITED	XI
Diclofenac Potassium 50 mg Tablets	not available	PL 20075/0683	ACCORD HEALTHCARE LIMITED	XI
Diclofenac Tablets 25 mg	not available	PL 04416/0361	SANDOZ LTD	XI
Diclofenac Tablets BP 25mg	not available	PL 20416/0219	CRESCENT PHARMA LIMITED	XI
Diclofenac Tablets BP 25mg	not available	PL 20416/0219	CRESCENT PHARMA LIMITED	XI
Diclofenac Tablets BP 50mg	not available	PL 20416/0220	CRESCENT PHARMA LIMITED	XI
Diclofenac Tablets BP 50mg	not available	PL 20416/0220	CRESCENT PHARMA LIMITED	XI
Diclofenac Zentiva 25 mg	not available	74529.00.00	WINTHROP ARZNEIMITTEL	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Filmtabletten			GMBH	
Diclofenacnatrium Apotex 100 mg	not available	RVG 56779	APOTEX EUROPE B.V.	NL
Diclofenacnatrium Apotex 25 mg	not available	RVG 57384	APOTEX EUROPE B.V.	NL
Diclofenacnatrium Apotex 50 mg	not available	RVG 57385	APOTEX EUROPE B.V.	NL
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
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
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	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
Labs 75 mg Retardtabletten				
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
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	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
Retardtabletten				
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
Diclomax Retard.	not available	PL 27827/0005	GALEN LIMITED	XI
Diclo-ratiopharm® bei Schmerzen und Fieber 25 mg Filmtabletten	not available	74527.00.00	RATIOPHARM GMBH	DE
DICLOREUM 100 MG COMPRESSE A RILASCIO PROLUNGATO	not available	024515088	ALFASIGMA S.P.A.	IT
Dicloreum 150 mg capsule cu eliberare prelungită	not available	8293/2015/01	ALFASIGMA S.P.A.	RO
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 50 MG GRANULATO PER SOLUZIONE ORALE	not available	024515138	ALFASIGMA S.P.A.	IT
DICLOREUM 75 MG/3 ML SOLUZIONE INIETTABILE PER USO INTRAMUSCOLARE	not available	024515076	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618015	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25	not available	028618041	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
mg granulato per soluzione orale				
Diclovol 75mg SR Tablets	not available	PL 04416/0645	SANDOZ LTD	XI
Dolotren 100 mg supositorios	not available	57.203	FAES FARMA, S.A.	ES
Dolotren 46,5 mg comprimidos dispersables	not available	60.068	FAES FARMA, S.A.	ES
Dolotren 50 mg comprimidos gastrorresistentes	not available	57.202	FAES FARMA, S.A.	ES
Dolotren 75 mg solución inyectable	not available	57.204	FAES FARMA, S.A.	ES
Dolotren retard 100 mg cápsulas duras de liberación prolongada	not available	58.435	FAES FARMA, S.A.	ES
Dolo-Voltarén 46, 5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 615 1 9	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 641 2 1	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 616 8 7	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 615 1 9	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 641 2 1	IBSA PHARMA SAS	FR
FLECTOR 25 mg,	not available	34009 352 616 8 7	IBSA PHARMA SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
granulés pour solution buvable en sachet-dose				
Flector 50 mg granule pro perorální roztok v sáčku	not available	29/514/99-C	IBSA SLOVAKIA S.R.O.	CZ
Flector 50 mg granule pro perorální roztok v sáčku	not available	29/514/99-C	IBSA SLOVAKIA S.R.O.	CZ
FLECTOR 50 mg, comprimé	not available	34009 301 460 6 4	LABORATOIRES GENEVRIER	FR
FLECTOR 50 mg, comprimé	not available	34009 301 460 6 4	LABORATOIRES GENEVRIER	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 617 4 8	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 642 9 9	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 617 4 8	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 642 9 9	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose.	not available	34009 352 618 0 9	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose.	not available	34009 352 618 0 9	IBSA PHARMA SAS	FR
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/01	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/02	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/03	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5	CZ/H/0928/001	OGYI-T-23250/04	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
mg lágy kapszula				
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/01	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/02	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/03	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/04	IBSA PHARMA KFT	HU
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617037	IBSA FARMACEUTICI ITALIA	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617052	IBSA FARMACEUTICI ITALIA	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617037	IBSA FARMACEUTICI ITALIA	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617052	IBSA FARMACEUTICI ITALIA	IT
FLECTOR EP 12,5 mg mäkké kapsuly	CZ/H/0928/001	29/0249/17-S	IBSA SLOVAKIA S.R.O.	SK
FLECTOR EP 12,5 mg mäkké kapsuly	CZ/H/0928/001	29/0249/17-S	IBSA SLOVAKIA S.R.O.	SK
Flector EP Rapid 50 mg granulát	not available	29/0906/96-S	IBSA SLOVAKIA S.R.O.	SK
Flector EP Rapid 50 mg granulát	not available	29/0906/96-S	IBSA SLOVAKIA S.R.O.	SK
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 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

Product 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	HU/H/0796/001	OGYI-T-23318/01	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	HU/H/0796/001	OGYI-T-23318/02	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	HU/H/0796/001	OGYI-T-23318/03	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	HU/H/0796/001	OGYI-T-23318/01	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	HU/H/0796/001	OGYI-T-23318/02	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció	HU/H/0796/001	OGYI-T-23318/03	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/004	OGYI-T-23318/10	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/004	OGYI-T-23318/10	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/04	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/05	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/06	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/04	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/05	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció	HU/H/0796/002	OGYI-T-23318/06	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/005	OGYI-T-23318/11	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/005	OGYI-T-23318/11	IBSA PHARMA KFT	HU
Flector Rapiven 75	HU/H/0796/003	OGYI-T-23318/07	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
mg/ml oldatos injekció				
Flector Rapiven 75 mg/ml oldatos injekció	HU/H/0796/003	OGYI-T-23318/08	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	HU/H/0796/003	OGYI-T-23318/09	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	HU/H/0796/003	OGYI-T-23318/07	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	HU/H/0796/003	OGYI-T-23318/08	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció	HU/H/0796/003	OGYI-T-23318/09	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció eloretöltött fecskendoben	HU/H/0796/006	OGYI-T-23318/12	IBSA PHARMA KFT	HU
Flector Rapiven 75 mg/ml oldatos injekció eloretöltött fecskendoben	HU/H/0796/006	OGYI-T-23318/12	IBSA PHARMA KFT	HU
FLECTORFLAM 50 mg granulato per soluzione orale	not available	036058030	IBSA FARMACEUTICI ITALIA	IT
FLECTORFLAM 50 mg granulato per soluzione orale	not available	036058030	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608014	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608026	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608038	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608040	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608014	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608026	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608038	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
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608040	IBSA FARMACEUTICI ITALIA	IT
Flectorgo, 12,5 mg, kapsułki, miękkie	CZ/H/0928/001	25425	IBSA FARMACEUTICI ITALIA	PL
Flectorgo, 12,5 mg, kapsułki, miękkie	CZ/H/0928/001	25425	IBSA FARMACEUTICI ITALIA	PL
FLOGOFENAC 100 mg capsule rigide a rilascio prolungato	not available	025536020	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Itamifast 25 mg compresse rivestite con film	IT/H/0352/001	041736012	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	041736036	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 50 mg compresse rivestite con film	IT/H/0352/002	041736048	FIDIA FARMACEUTICI S.P.A	IT
Modifenac 75 mg hart hylki með breyttan losunarhraða	not available	980422	ACTAVIS GROUP PTC EHF.	IS
Modifenac 75 mg kapsler med modifisert frisetting, harde	not available	96-3618	ACTAVIS GROUP PTC EHF.	NO
Motifene 75 mg gélules à libération modifiée.	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
Motifene 75 mg gélules à libération modifiée.	not available	0235011	DAIICHI SANKYO BELGIUM S.A	LU
Motifene 75 mg, capsules met gereguleerde afgifte, hard	not available	BE 176671	DAIICHI SANKYO BELGIUM S.A	BE

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
Product full name (in authorisation country as per Art. 57 data)	MRP/DCP or CP Authorisation number (if not available then insert not available)	National Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Rhumalgan CR 100	not available	PL 04416/0243	SANDOZ LTD	XI
Rhumalgan CR 75	not available	PL 04416/0242	SANDOZ LTD	XI
Rhumalgan SR 75 mg Modified Release Capsules	not available	PL 04416/0737	SANDOZ LTD	XI
Rhumalgan XL 100 mg modified-release capsules.	not available	PL 04416/0738	SANDOZ LTD	XI
SAFIDOL 25 mg compresse rivestite con film	IT/H/0483/001	041735010	SAFI MEDICAL CARE S.R.L	IT
SAFIDOL 25 mg compresse rivestite con film	IT/H/0483/001	041735022	SAFI MEDICAL CARE S.R.L	IT
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
Sooneral or.sol 50 mg/ 5ml	not available	88209/15/07-03-2018	NASSINGTON LTD	GR
TRAULEN 100 mg	not available	033420.023	OP PHARMA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse a rilascio prolungato				
TRAULEN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	033420.047	OP PHARMA S.R.L.	IT
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 25 mg - Filmtabletten	not available	1-15554	NOVARTIS PHARMA GMBH	AT
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 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
Voltaren 100 mg peräpuikko	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 100 mg suppositoires	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
Voltaren 100 mg Zäpfchen	not available	BE109261	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 zetpillen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
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
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
Voltaren 25 mg comprimés gastro-résistants	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg enterotabletter	not available	7442	NOVARTIS FINLAND OY	FI
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg	not available	09737	NOVARTIS SVERIGE AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
enterotabletter				
Voltaren 25 mg enterotabletti	not available	7442	NOVARTIS FINLAND OY	FI
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
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
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25, 25 mg cápsulas moles	not available	5204466	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 25, 25 mg cápsulas moles	not available	5204466	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E HIGIENE, LDA	PT
Voltaren 25, 25 mg, cápsulas moles	not available	5204508	GLAXOSMITHKLINE CONSUMER HEALTHCARE, PRODUTOS PARA A SAÚDE E	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
			HIGIENE, LDA	
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 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 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
VOLTAREN 50 enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
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 comprimés gastro-résistants	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg	not available	4636197	NOVARTIS FARMA -	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos gastrorresistentes			PRODUTOS FARMACÊUTICOS S.A.	
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
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
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	09738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletti	not available	7906	NOVARTIS FINLAND OY	FI
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg maagsapresistente tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg	not available	122880202	NOVARTIS (HELLAS)	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
γαστροανθεκτικά δισκία			S.A.C.I.	
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
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 compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 75 mg δισκία επικαλυμμένα με υμένιο βραδείας αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία επικαλυμμένα με υμένιο βραδείας αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/ 3 ml oplossing voor injectie	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml	not available	4636593	NOVARTIS FARMA -	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
solução injetável			PRODUTOS FARMACÊUTICOS S.A.	
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solution injectable	not available	BE113206	NOVARTIS PHARMA N.V.	BE
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
Voltaren Acti Forte, 25 mg, tabletki powlekane	not available	16489	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren Acti, 12,5 mg, tabletki powlekane	not available	9462	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren Actigo Extra 25 mg obalené tablety	not available	29/0227/01-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Voltaren ActiGo Extra obalené tablety	not available	29/549/00-C	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	CZ
Voltaren Akti 12,5 mg apvalkotās tabletes	not available	04-0291	GLAXOSMITHKLINE DUNGARVAN LIMITED	LV
Voltaren Akti 12,5 mg ohukese polumeerikattega tabletid N20	not available	434104	GLAXOSMITHKLINE DUNGARVAN LIMITED	EE
Voltaren Akti 12,5 mg plėvele dengtos tabletės	not available	LT/1/94/0948/003	GLAXOSMITHKLINE DUNGARVAN 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 Akti 12,5 mg plėvele dengtos tabletės PRD1798846	not available	LT/1/94/0948/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti 12,5 mg plėvele dengtos tabletės PRD350247	not available	LT/1/94/0948/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti 25 mg mīkstās kapsulas	not available	12-0185	GLAXOSMITHKLINE DUNGARVAN LIMITED	LV
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/005	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/004	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės PRD1801418	not available	LT/1/94/0948/005	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti 25 mg minkštosios kapsulės PRD353399	not available	LT/1/94/0948/004	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Akti, 12,5 mg õhukese polümeerikattega tabletid	not available	434104	GLAXOSMITHKLINE DUNGARVAN LIMITED	EE
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/001	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/002	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/004	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Dolo 12,5 mg Filmtabletten	not available	51218.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
VOLTAREN DOLO 12,5 MG LÁGY KAPSZULA PRD1744411	not available	OGYI-T-5572/21	GLAXOSMITHKLINE-CONSUMER KFT.	HU
VOLTAREN DOLO 12,5	not available	OGYI-T-5572/22	GLAXOSMITHKLINE-	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MG LÁGY KAPSZULA PRD1772476			CONSUMER KFT.	
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 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
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
Voltaren Dolo 25 mg überzogene Tablette	not available	64958.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Dolo 25 mg überzogene Tablette	not available	43151.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Dolo Liquid 25 mg Weichkapsel	not available	82502.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Express Forte, 25 mg, kapsułki miękkie	not available	17890	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren Express, 12,5 mg, kapsułki, miękkie	not available	16009	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	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 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren K 12,5 mg, filmomhulde tabletten	not available	RVG 20982	GLAXOSMITHKLINE CONSUMER HEALTHCARE B.V.	NL
Voltaren K 25 mg, omhulde tabletten	not available	RVG 13244	GLAXOSMITHKLINE CONSUMER HEALTHCARE B.V.	NL
Voltaren Rapid 25 mg dragerade tabletter	not available	10202	NOVARTIS FINLAND OY	FI
Voltaren Rapid 25 mg mäkké kapsuly	not available	29/0709/10-S	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	SK
Voltaren Rapid 25 mg měkké tobolky	not available	29/173/11-C	GLAXOSMITHKLINE CONSUMER HEALTHCARE CZECH REPUBLIC S.R.O.	CZ
Voltaren Rapid 25 mg tabletti, päällystetty	not available	10202	NOVARTIS FINLAND OY	FI
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
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg Obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg	not available	10203	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
tabletti, päällystetty				
VOLTAREN RAPID 50 mg TABLETY obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 mg comprimés à libération prolongée	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletti	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte	not available	BE122071	NOVARTIS PHARMA N.V.	BE
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
Voltaren Retard 75 mg comprimés à libération prolongée	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren Retard 75 mg	not available	10919	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
depottabletter				
Voltaren Retard 75 mg depottabletti	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg Retardtabletten	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 SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
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 T 12,5 mg filmdragerade tabletter	not available	16081	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
Voltaren T 25 mg mjuka kapslar.	not available	41904	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	SE
Voltaren T 25 mg, dragerade tabletter	not available	12427	GLAXOSMITHKLINE CONSUMER HEALTHCARE APS	SE
Voltaren T 50 mg, dragerade tabletter	not available	11242	NOVARTIS SVERIGE AB	SE
Voltaren, 25 mg/1 ml, roztwór do wstrzykiwan lub do sporzadzania roztworu do infuzji	not available	R/1205	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, tabletki dojelitowe	not available	R/1204	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
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
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
VOLTAREN® 100 mg suppositories	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 12,5 mg svecke	not available	H/94/01506/001	NOVARTIS PHARMA GMBH	SI
Voltaren® 25 mg svecke	not available	H/94/01506/002	NOVARTIS PHARMA GMBH	SI
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
VOLTAREN® 50 mg suppositories	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
VOLTAREN® D	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
Voltaren® dispers -	not available	1-20832	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
lösliche Tabletten				
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® K Migräne 50 mg Überzogene Tabletten	not available	43151.01.00	NOVARTIS PHARMA GMBH	DE
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren® Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
VOLTAREN®12.5 mg suppositories	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTARENE 100 mg, suppositoire	not available	3400932214341	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

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VOLTARENE 50 MG, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
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	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	3400933070298	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	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
Voltarol 12,5 mg kapsel, myk	not available	09-6597	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
Voltarol 12,5 mg tabletter, filmdrasjerte	not available	99-7996	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
Voltarol 25 mg drasjerte	not available	09-1063	GLAXOSMITHKLINE	NO

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tabletter			CONSUMER HEALTHCARE A/S	
Voltarol 25 mg kapsler, myke	not available	10-8136	GLAXOSMITHKLINE CONSUMER HEALTHCARE A/S	NO
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol Acti-Go 25mg Επικαλυμμένα Δισκία	not available	321460101	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SA	GR
VOLTAROL ACTI-GO, Επικαλυμμένα Δισκία 50 MG	not available	321460201	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SA	GR
VOLTAROL ACTI-GO, Επικαλυμμένο με λεπτό υμένιο δισκίο 12,5mg/tab	not available	321460301	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SA	GR
VOLTAROL ACTI-GO, Επικαλυμμένο με λεπτό υμένιο δισκίο 12,5mg/tab	not available	321460302	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SA	GR
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	PA 0896/034/003	NOVARTIS IRELAND LIMITED	IE
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories	not available	PL 00101/0472	NOVARTIS	XI

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12.5mg			PHARMACEUTICALS UK LIMITED	
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
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
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
ВОЛТАРЕН ДОЛО 12,5 mg Филмирани таблетки	not available	20030609	GLAXOSMITHKLINE DUNGARVAN LIMITED	BG
ВОЛТАРЕН ДОЛО 12,5 mg Филмирани таблетки	not available	20030609	GLAXOSMITHKLINE DUNGARVAN LIMITED	BG