



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

25 May 2016
EMA/475009/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance(s): diclofenac (systemic formulations)

Procedure No.: PSUSA/00001048/201509



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		PL 20046/0078	Focus Pharmaceuticals Limited	United Kingdom
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	021855 & 021856 & 021857	Ibsa Farmaceutici Italia	Cyprus
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	021858 & 021859 & 021860	Ibsa Farmaceutici Italia	Cyprus
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	34009 300 350 4 7 & 34009 300 350 5 4 & 34009 300 350 6 1 & 34009 300 350 8 5 & 34009 300 350 9 2 & 34009 300 351 0 8 & 34009 300 351 1 5 & 34009 300 351 2 2 & 34009 550 124 5 0	Laboratoires Genevrier	France
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	34009 300 351 4 6 & 34009 300 351 5 3 & 34009 300 351 6 0 & 34009 300 351 7 7 & 34009 300 351 8 4 & 34009 300 351 9 1 & 34009 300 352 0 7 & 34009 300 352 1 4 & 34009 550 124 6 7	Laboratoires Genevrier	France
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	83946.00.00 & 83947.00.00 & 83948.00.00	Ibsa Farmaceutici Italia	Germany
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	83949.00.00 & 83950.00.00 & 83951.00.00	Ibsa Farmaceutici Italia	Germany
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	OGYI-T-22385/01 & OGYI-T-22385/02 & OGYI-T-22385/03 & OGYI-T-22385/04 & OGYI-T-22385/05 & OGYI-T-22385/06 & OGYI-T-22385/07 & OGYI-T-22385/08 & OGYI-T-22385/09	Ibsa Pharma Kft	Hungary
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	OGYI-T-22385/10 & OGYI-T-22385/11 & OGYI-T-22385/12 & OGYI-T-22385/13 & OGYI-T-22385/14 & OGYI-T-22385/15 & OGYI-T-22385/16 & OGYI-T-22385/17 & OGYI-T-22385/18	Ibsa Pharma Kft	Hungary
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	040528010 & 040528022 & 040528034 & 040528046 & 040528059 & 040528061 & 040528073 & 040528085 & 040528097	Ibsa Farmaceutici Italia	Italy

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Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	040528109 & 040528111 & 040528123 & 040528135 & 040528147 & 040528150 & 040528162 & 040528174 & 040528186	Ibsa Farmaceutici Italia	Italy
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	21017 & 21018 & 21019	Ibsa Farmaceutici Italia	Poland
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	21020 & 21021 & 21022	Ibsa Farmaceutici Italia	Poland
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	29/0188/13-S & 29/0189/13-S & 29/0190/13-S	Ibsa Slovakia S.R.O.	Slovakia
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	29/0185/13-S & 29/0186/13-S & 29/0187/13-S	Ibsa Slovakia S.R.O.	Slovakia
Akis	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	PL 21039/0018 & PL 21039/0019 & PL 21039/0020	Ibsa Farmaceutici Italia	United Kingdom
Akis	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	PL 21039/0021 & PL 21039/0023 & PL 21039/0024	Ibsa Farmaceutici Italia	United Kingdom
Akisflam		036058016 & 036058028 & 036058030	Ibsa Farmaceutici Italia S.R.L.	Italy
Catafast		088/00303	Novartis Pharmaceuticals Uk Limited	Malta
Cataflam		BE147402	Novartis Pharma N.V. (Art57)	Belgium
Cataflam		BE353972	Novartis Pharma N.V. (Art57)	Belgium
Cataflam		BE353972	Novartis Pharma N.V. (Art57)	Belgium
Cataflam		18492	Novartis Pharmaceuticals Uk Limited	Cyprus
Cataflam		18493	Novartis Pharmaceuticals Uk Limited	Cyprus
Cataflam		060994	Novartis Finland Oy (Art57)	Estonia

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Cataflam		OGYI-T-5573/02	Novartis Hungária Kft. Pharma (Art57)	Hungary
Cataflam		OGYI-T-5573/03	Novartis Hungária Kft. Pharma (Art57)	Hungary
Cataflam		PA 13/88/2	Novartis Pharmaceuticals Uk Limited	Ireland
Cataflam		99-0168	Novartis Finland Oy (Art57)	Latvia
Cataflam		LT/1/94/0948/002	Novartis Finland Oy (Art57)	Lithuania
Cataflam		0010/01/10/7175	Novartis Pharma N.V. (Art57)	Luxembourg
Cataflam		0010/01/10/7175	Novartis Pharma N.V. (Art57)	Luxembourg
Cataflam		088/00301	Novartis Pharmaceuticals Uk Limited	Malta
Cataflam		088/00302	Novartis Pharmaceuticals Uk Limited	Malta
Cataflam		RVG 13245	Novartis Pharma B.V. (Art57)	Netherlands
Cataflam		05-3486	Novartis Norge As (Art57)	Norway
Cataflam		8287	Novartis Norge As (Art57)	Norway
Cataflam		99-7997	Novartis Norge As	Norway
Cataflam		R/3707	Novartis Poland Sp. Z O. O. (Art57)	Poland
Cataflam		4566196 & 4566790 & 9716522 & 9716530	Laboratorio Normal-Produtos Farmaceuticos, Lda	Portugal
Cataflam Dispersible		BE172514	Novartis Pharma N.V. (Art57)	Belgium
Cataflam Dispersible		BE172514	Novartis Pharma N.V. (Art57)	Belgium
Cataflam Dolo		OGYI-T-5573/04 & OGYI-T-5573/05 & OGYI-T-5573/06	Novartis Hungária Kft. Consumer Health	Hungary
Cataflam-V		OGYI-T-5573/01	Novartis Hungária Kft. Pharma (Art57)	Hungary
Deflamat		AIC 028534016 & AIC 028534028	Daiichi Sankyo Italia S.P.A	Italy

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Deflamat		AIC 028534030	Daiichi Sankyo Italia S.P.A	Italy
Diclac		OGYI-T-4000/05 & OGYI-T-4000/06	Sandoz Hungária Kft	Hungary
Diclac		OGYI-T-4000/07 & OGYI-T-4000/08	Sandoz Hungária Kft	Hungary
Diclac	NOT APPLICABLE	OGYI-T-4000/03 & OGYI-T-4000/04	Sandoz Hungária Kft	Hungary
Diclac		PA 711/9/3	Rowex Ltd	Ireland
Diclac	99-0827-003	99-0827	Sandoz Pharmaceuticals D.D.	Latvia
Diclac	NOT APPLICABLE	29/0104/93-S	Sandoz Pharmaceuticals D.D.	Slovakia
Diclac Duo		9577 & 9578	Sandoz Gmbh	Poland
Diclo 75 mg		PA 281/106/1	Pinewood Laboratories Ltd.	Ireland
Dicloident		4881389;	Laboratórios Atral, S.A.	Portugal
Diclo-Divido		13847.01.00	Actavis Deutschland Gmbh & Co. Kg	Germany
Diclo-Divido Long		13847.00.00	Actavis Deutschland Gmbh & Co. Kg	Germany
Diclofenac		97-0174	Glaxosmithkline Latvia Sia	Latvia
Diclofenac		97-0175	Glaxosmithkline Latvia Sia	Latvia
Diclofenac Akut 1a Pharma		1-24778	1a Pharma Gmbh	Austria
Diclofenac Alfa Wassermann		033612033	Alfa Wassermann Spa	Italy
Diclofenac Alfa Wassermann		033612058	Alfa Wassermann Spa	Italy
Diclofenac Alfa Wassermann		033612096	Alfa Wassermann Spa	Italy
Diclofenac Crescent Pharma		PL 20416/0219	Crescent Pharma Limited	United Kingdom
Diclofenac Crescent Pharma		PL 20416/0220	Crescent Pharma Limited	United Kingdom
Diclofenac Hikma		5200.00.01	Hikma Farmaceutica (Portugal), S.A.	Germany
Diclofenac Mylan		NL 16785 & NL 16786	Mylan S.A.S	France
Diclofenac Nch		70999.00.00	Novartis Consumer Health Gmbh	Germany

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Diclofenac Potassium		PA 30/41/1	Novartis Consumer Health (Uk) Limited	Ireland
Diclofenac Potassium 50mg Tablets & Adacium™ Rapid 50mg Tablets		PL 20046/0078	Focus Pharmaceuticals Limited	United Kingdom
Diclofenac Potassium Actavis	UK/N/0001	PL 30306/0255	Actavis Group Ptc Ehf.	United Kingdom
Diclofenac Potassium Actavis	UK/N/0001	PL 30306/0256	Actavis Group Ptc Ehf.	United Kingdom
Diclofenac Potassium Focus Pharmaceuticals		PL 20046/0078	Focus Pharmaceuticals Limited	United Kingdom
Diclofenac Ratiopharm		31404.00.00	Ratiopharm Gmbh	Germany
Diclofenac Sandoz Ltd		PL 04416/0361 & PL 04416/0362	Sandoz Ltd	United Kingdom
Diclofenac Sodium Injection 75 mg in 3 ml		PL 12762/0421	Mercury Pharma International Limited	United Kingdom
Diclofenac Sodium Injection 75 mg in 3 ml		PL 12762/0092	Mercury Pharmaceuticals Limited	United Kingdom
Diclofenac Sodium Ratiopharm		4369.00.00	Ratiopharm Gmbh	Germany
Diclofenac Sodium Tablets		PL 12762/0420	Mercury Pharma International Limited	United Kingdom
Diclofenac Sodium Tablets		PL 12762/0421	Mercury Pharma International Limited	United Kingdom
Diclofenac Zentiva		74529.00.00	Winthrop Arzneimittel Gmbh	Germany
Diclofenac-Kalium Al		74528.00.00	Aliud Pharma Gmbh	Germany
Diclofenac-Kalium Stada		74527.00.00	Stadapharm Gmbh	Germany
Diclofenacnatrium Sandoz	RVG 25586	RVG 25586	Sandoz B.V.	Netherlands

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Dicloin	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	29/234/13-C & 29/235/13-C & 29/236/13-C	Ibsa Slovakia S.R.O.	Czech Republic
Dicloin	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	29/237/13-C & 29/238/13-C & 29/239/13-C	Ibsa Slovakia S.R.O.	Czech Republic
Dicloin	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	75814/2-10-2013 & 75815/2-10-2013 & 75816/2-10-2013	Ibsa Farmaceutici Italia	Greece
Dicloin	UK/H/3585/001 & UK/H/3585/002 & UK/H/3585/003	75814/2-10-2013 & 75815/2-10-2013 & 75816/2-10-2013	Ibsa Farmaceutici Italia	Greece
Dicloin	UK/H/3585/004 & UK/H/3585/005 & UK/H/3585/006	75817/2-10-2013 & 75818/2-10-2013 & 75819/2-10-2013	Ibsa Farmaceutici Italia	Greece
Diclomax Retard		PL 27827/0005	Galen Limited	United Kingdom
Diclon	12735-1986	12735	Sandoz A/S	Denmark
Diclon	NL/H/0173/002	31064	Sandoz Gmbh	Denmark
Diclon	NL/H/0173/003 & NL/H/0173/004	31065 & 31066	Sandoz Gmbh	Denmark
Diclopax	DK/H/0582/001	33076	Actavis Group Ptc Ehf.	Denmark
Dicloral		032085019	Athena Pharma Italia Srl	Italy
Dicloral		032085033	Athena Pharma Italia Srl	Italy
Dicloream		29/126/94-A/C	Alfa Wassermann Spa	Czech Republic
Dicloream		29/126/94-B/C	Alfa Wassermann Spa	Czech Republic
Dicloream		29/844/92-B/C	Alfa Wassermann Spa	Czech Republic
Dicloream		024515049	Alfa Wassermann Spa	Italy
Dicloream		024515052 & 024515064	Alfa Wassermann Spa	Italy
Dicloream		024515088	Alfa Wassermann Spa	Italy
Dicloream		024515114	Alfa Wassermann Spa	Italy

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
Diclorem		024515138	Alfa Wassermann Spa	Italy
Diclorem		8293/2006/01	Alfa Wassermann Spa	Romania
Diclorem		29/0336/94-S	Alfa Wassermann Spa	Slovakia
Diclorem		29/0668/92-C/S	Alfa Wassermann Spa	Slovakia
Diclorem		29/0844/92-C/S	Alfa Wassermann Spa	Slovakia
Diclorem Dolore		028618015 & 028618041	Alfa Wassermann Spa	Italy
Diclorem Dolore		028618027 & 028618039	Alfa Wassermann Spa	Italy
Diclorem Iniettabile		024515076	Alfa Wassermann Spa	Italy
Diclorem Retard		29/127/94-C	Alfa Wassermann Spa	Czech Republic
Diclorem Retard		29/0338/94-S	Alfa Wassermann Spa	Slovakia
Diclovol	PL 04416/0645	PL 04416/0645	Sandoz Ltd	United Kingdom
Dicuno	SE/H/791/002/DC	29/496/11-C, 29/497/11-C	Vitabalans Oy	Czech Republic
Dicuno	SE/H/791/002/DC	45574, 45575	Vitabalans Oy	Denmark
Dicuno	SE/H/791/002/DC	743711, 743811	Vitabalans Oy	Estonia
Dicuno	SE/H/791/002/DC	27849, 27850	Vitabalans Oy	Finland
Dicuno	SE/H/791/002/DC	79584.00.00, 79585.00.00	Vitabalans Oy	Germany
Dicuno	SE/H/791/002/DC	11-0212, 11-0213	Vitabalans Oy	Latvia
Dicuno	SE/H/791/002/DC	LT/1/11/2667/001-008	Vitabalans Oy	Lithuania
Dicuno	SE/H/791/002/DC	09-6972, 09-6973	Vitabalans Oy	Norway
Dicuno	SE/H/791/001/DC	19642	Vitabalans Oy	Poland
Dicuno	SE/H/791/002/DC	19643	Vitabalans Oy	Poland
Dicuno	SE/H/791/002/DC	29/0242/11-S, 29/0243/11-S	Vitabalans Oy	Slovakia
Dicuno	SE/H/791/002/DC	5363-I-2352 - 2357/11	Vitabalans Oy	Slovenia
Dicuno	SE/H/791/002/DC	42951, 42952	Vitabalans Oy	Sweden

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Dixol		OGYI-T-20 733/01	Angelini Pharma österreich Gmbh	Hungary
Dixol Forte		OGYI-T-20733/02	Angelini Pharma österreich Gmbh	Hungary
Dolo-Voltarén		61440	Novartis Farmacéutica S.A. (Art57)	Spain
Econac 100mg Suppositories		PL 10972/0069	Mercury Pharma Group Limited	United Kingdom
Econac injection 75 mg / 3 ml		PL 10972/0070	Mercury Pharma Group Limited	United Kingdom
Econac SR 75mg Tablets		PL 12762/0100	Mercury Pharmaceuticals Limited	United Kingdom
Econac XL 100mg Tablets		PL 12762/0101	Mercury Pharmaceuticals Limited	United Kingdom
Eminocs	NL/H/0800/001	038049019 & 038049021 & 038049033 & 038049045	Alfa Wassermann Spa	Italy
Eminocs	NL/H/0800/001	RVG 31532	Alfa Wassermann Spa	Netherlands
EVINOPON CUT.SOL 1,5% W/W		11054/13-2-2012	BROS E.Π.E.	Greece
Flameril		4637096 & 4637195 & 9548818 & 9548826	Laboratorio Normal-Produtos Farmaceuticos, Lda	Portugal
Flameril		4637484 & 8548909	Laboratorio Normal-Produtos Farmaceuticos, Lda	Portugal
Flameril		9549006	Laboratorio Normal-Produtos Farmaceuticos, Lda	Portugal
Flameril Retard		4498291 & 4637393 & 9594002	Laboratorio Normal-Produtos Farmaceuticos, Lda	Portugal
Flector		34009 352 615 1 9 & 34009 352 616 8 7 & 34009 352 617 4 8 & 34009 352 618 0 9	Laboratoires Genevrier	France
Flector		34009 352 641 2 1 & 34009 352 642 9 9	Laboratoires Genevrier	France

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 Dolore		028617037	Ibsa Farmaceutici Italia	Italy
Flector Ep Rapid		29/514/99-C	Ibsa Slovakia S.R.O.	Czech Republic
Flector Ep Rapid		29/0906/96-S	Ibsa Slovakia S.R.O.	Slovakia
Flector Rapid		OGYI-T-05033/03 & OGYI-T-05033/04	Ibsa Pharma Kft	Hungary
Flogofenac		025536020	A. Menarini - Industrie Farmaceutiche Riunite - S.R.L.	Italy
Glimbax		14588	Angelini Pharma Polska Sp. Z O.O.	Poland
Glimbax		7416/2006/01	Angelini Pharma österreich GmbH	Romania
Glimbax		95/0200/04-S	Angelini Pharma österreich GmbH	Slovakia
Glimbax		95/0378/13-S	Angelini Pharma österreich GmbH	Slovakia
Indicam		036972014	Altergon Italia S.R.L.	Italy
Indicam		036972026 & 036972038	Altergon Italia S.R.L.	Italy
Inforce		036973028 & 036973030	Altergon Italia S.R.L.	Italy
Inforce		036973079 & 036973081 & 036973093 & 036973131 & 036973143 & 036973156	Altergon Italia S.R.L.	Italy
Itamifast	IT/H/352/001-002	041736012 & 041736024 & 041736036 & 041736048	Fidia Farmaceutici S.P.A	Italy
Kappadi	SE/H/1162/001 & SE/H/1162/002	041735010 & 041735022 & 041735034 & 041735046	Alfa Wassermann Spa	Italy
Kappadi	SE/H/1162/001 & SE/H/1162/002	43386 & 43387	Alfa Wassermann Spa	Sweden
Klófen-L	910127	910127	Actavis Hf.	Iceland
Modifenac		18609	Actavis Group Hf.	Denmark
Modifenac	IS/N/0001	980422	Actavis Group Hf.	Iceland
Modifenac®	96-3618	96-3618	Actavis Group Hf.	Norway
Motifene		BE 176671	Daiichi Sankyo Belgium S.A	Belgium
Motifene		0235011	Daiichi Sankyo Belgium S.A	Luxembourg

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Motifene		0345531	Daiichi Sankyo Belgium S.A	Luxembourg
Motifene		PL 08265/0003	Daiichi Sankyo Uk Ltd	United Kingdom
Novapirina		024951028	Novartis Consumer Health S.P.A.	Italy
Otriflu		PA 30/42/1	Novartis Consumer Health (Uk) Limited	Ireland
Otriflu		RVG 29462	Novartis Consumer Health B.V.	Netherlands
Otriflu		3765583 & 3765682 & 5047220 & 5047238	Glaxosmithkline Consumer Healthcare	Portugal
Rheumatac Retard 75		PL 20072/0219	Amdipharm UK Limited	United Kingdom
Rhumalgan Cr		PL 04416/0243	Sandoz Ltd	United Kingdom
Rhumalgan Cr		PL 4416/0242	Sandoz Ltd	United Kingdom
Rhumalgan Sr		PL 04416/0737	Sandoz Ltd	United Kingdom
Rhumalgan XI		PL 04416/0738	Sandoz Ltd	United Kingdom
Slofenac Sr		PL 34976/0007 & PL 34976/0008	Ri Pharma Ltd	United Kingdom
Traulen		033420.023	Op Pharma S.R.L.	Italy
Traulen		033420.047	Op Pharma S.R.L.	Italy
Volsaid		PL 8829/0045 & PL 8829/0046	Chiesi Limited	United Kingdom
Voltadvance		035500014 & 035500026	Novartis Consumer Health S.P.A.	Italy
Voltaren		1-15554	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		1-15868 & 1-16308	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		1-15915	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		1-16506	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		1-20832	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		1-31725	Novartis Pharma Gmbh (Art57)	Austria
Voltaren		BE095916	Novartis Pharma N.V. (Art57)	Belgium

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Voltaren		BE109261	Novartis Pharma N.V. (Art57)	Belgium
Voltaren		BE113206	Novartis Pharma N.V. (Art57)	Belgium
Voltaren		BE121116	Novartis Pharma N.V. (Art57)	Belgium
Voltaren		BE235977	Novartis Consumer Health N.V	Belgium
Voltaren		18400	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18434	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18443	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18444	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18454	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18455	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18457	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		18459	Novartis Pharmaceuticals Uk Limited	Cyprus
Voltaren		29/186/80-C	Novartis, S.R.O. (Art57)	Czech Republic
Voltaren		29/294/91-C	Novartis, S.R.O. (Art57)	Czech Republic
Voltaren		09002 & 10109	Novartis Healthcare A/S (Art57)	Denmark
Voltaren		10282	Novartis Healthcare A/S (Art57)	Denmark
Voltaren		10651	Novartis Healthcare A/S (Art57)	Denmark
Voltaren		11452	Novartis Healthcare A/S (Art57)	Denmark
Voltaren		061694	Novartis Finland Oy (Art57)	Estonia
Voltaren		7442 & 7906	Novartis Finland Oy (Art57)	Finland
Voltaren		8090	Novartis Finland Oy (Art57)	Finland

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Voltaren		8629	Novartis Finland Oy (Art57)	Finland
Voltaren		14651.00.00	Novartis Pharma Gmbh (Art57)	Germany
Voltaren		14651.00.02	Novartis Pharma Gmbh (Art57)	Germany
Voltaren		2502.00.01	Novartis Pharma Gmbh (Art57)	Germany
Voltaren		520.00.00	Novartis Pharma Gmbh (Art57)	Germany
Voltaren		6164405.00.00	Novartis Pharma Gmbh (Art57)	Germany
Voltaren		122880101	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		122880201 & 122880202	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		122880301 & 122880801 & 122880802	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		122880401	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		122880501 & 122880601	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		122880901	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren		OGYI-T-5572/01 & OGYI-T-5572/02	Novartis Hungária Kft. Pharma (Art57)	Hungary
Voltaren		OGYI-T-5572/03 & OGYI-T-5572/04	Novartis Hungária Kft. Pharma (Art57)	Hungary
Voltaren		OGYI-T-5572/05	Novartis Hungária Kft. Pharma (Art57)	Hungary
Voltaren		OGYI-T-5572/06	Novartis Hungária Kft. Pharma (Art57)	Hungary
Voltaren		731593 & 822964	Novartis Healthcare A/S (Art57)	Iceland
Voltaren		843406	Novartis Healthcare A/S (Art57)	Iceland
Voltaren		IS/1/07/105/01	Novartis Healthcare A/S (Art57)	Iceland
Voltaren		IS/1/07/105/02	Novartis Healthcare A/S (Art57)	Iceland
Voltaren		023181011	Novartis Farma S.P.A. (Art57)	Italy
Voltaren		023181023	Novartis Farma S.P.A. (Art57)	Italy
Voltaren		023181035 & 023181074	Novartis Farma S.P.A. (Art57)	Italy

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		023181047	Novartis Farma S.P.A. (Art57)	Italy
Voltaren		023181086	Novartis Farma S.P.A. (Art57)	Italy
Voltaren		00-0517	Novartis Finland Oy (Art57)	Latvia
Voltaren		04-0291	Novartis Finland Oy	Latvia
Voltaren		12-0185	Novartis Finland Oy	Latvia
Voltaren		LT/1/94/0948/001	Novartis Finland Oy	Lithuania
Voltaren		LT/1/94/1300/005	Novartis Finland Oy (Art57)	Lithuania
Voltaren		0010/02/10/0205	Novartis Pharma N.V. (Art57)	Luxembourg
Voltaren		0218/02/10/0047	Novartis Consumer Health N.V	Luxembourg
Voltaren		2008049767	Novartis Pharma N.V. (Art57)	Luxembourg
Voltaren		088/04501 & 088/04502	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren		088/04503	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren		088/04505	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren		MA 088/04504	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren		MA 088/04506	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren		RVG 07003	Novartis Pharma B.V. (Art57)	Netherlands
Voltaren		RVG 07460	Novartis Pharma B.V. (Art57)	Netherlands
Voltaren		10-8136	Novartis Norge As	Norway
Voltaren		7377	Novartis Norge As (Art57)	Norway
Voltaren		7780 & 7781	Novartis Norge As (Art57)	Norway
Voltaren		7782 & 7783 & 7784	Novartis Norge As (Art57)	Norway
Voltaren		16009	Novartis Consumer Health Gmbh	Poland
Voltaren		17890	Novartis Consumer Health Gmbh	Poland

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		R/1203	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren		R/1204	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren		R/1205	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren		R/2769	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren		R/4897	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren		2134492 & 2134591 & 2134690 & 4636494	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren		4636098 & 4636197 & 9427815 & 9427831	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren		4636593 & 8446906	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren		4726394 & 4726493 & 9427856 & 9427864	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren		9447037	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren		4269/2004/01	Novartis Pharma GmbH (Art57)	Romania
Voltaren		4270/2004/01	Novartis Pharma GmbH (Art57)	Romania
Voltaren		4271/2004/01	Novartis Pharma GmbH (Art57)	Romania
Voltaren		4715/2004/01 & 4716/2004/01	Novartis Pharma GmbH (Art57)	Romania
Voltaren		29/0098/91-S	Novartis, S.R.O. (Art57)	Slovakia
Voltaren		29/0184/80-C/S	Novartis, S.R.O. (Art57)	Slovakia
Voltaren		29/0186/80-C/S	Novartis, S.R.O. (Art57)	Slovakia
Voltaren		29/0227/01-S	Novartis S.R.O. - Consumer Health Division	Slovakia
Voltaren		29/0247/80-C/S & 29/1029/92-S	Novartis, S.R.O. (Art57)	Slovakia
Voltaren		29/0294/91-C/S	Novartis, S.R.O. (Art57)	Slovakia

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		29/0492/07-S	Novartis, S.R.O. (Art57)	Slovakia
Voltaren		29/0508/08-S	Novartis S.R.O. - Consumer Health Division	Slovakia
Voltaren		29/0709/10-S	Novartis S.R.O. - Consumer Health Division	Slovakia
Voltaren		H/09/01954/001 & H/09/01954/002	Novartis Consumer Health Gmbh	Slovenia
Voltaren		55.004	Novartis Farmacéutica S.A. (Art57)	Spain
Voltaren		55005	Novartis Farmacéutica S.A. (Art57)	Spain
Voltaren		55010	Novartis Farmacéutica S.A. (Art57)	Spain
Voltaren		09737 & 09738	Novartis Sverige Ab (Art57)	Sweden
Voltaren		09818 & 09941 & 12840	Novartis Sverige Ab (Art57)	Sweden
Voltaren		09940	Novartis Sverige Ab (Art57)	Sweden
Voltaren		16081	Novartis Sverige Ab	Sweden
Voltaren		41904 & 42233	Novartis Sverige Ab	Sweden
Voltaren 12,5		3852589 & 3852688 & 5047246 & 5047253 & 5185525 & 5185533	Glaxosmithkline Consumer Healthcare	Portugal
Voltaren 12,5		5133525 & 5133533 & 5133541 & 5133558	Glaxosmithkline Consumer Healthcare	Portugal
Voltaren 25		5204458 & 5204466 & 5204474 & 5204508	Glaxosmithkline Consumer Healthcare	Portugal
Voltaren Acti		9462	Novartis Consumer Health Gmbh	Poland
Voltaren Acti Forte		16489	Novartis Consumer Health Gmbh	Poland
Voltaren Actigo		07/031/05-C	Novartis S.R.O. - Consumer Health Division	Czech Republic
Voltaren Acti-Go		251250101	Novartis (Hellas) S.A.C.I.	Greece
Voltaren Acti-Go		251250201	Novartis (Hellas) S.A.C.I.	Greece

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-Go		251250301 & 251250302	Novartis (Hellas) S.A.C.I.	Greece
Voltaren Actigo Extra		29/549/00-C	Novartis S.R.O. - Consumer Health Division	Czech Republic
Voltaren Akti		434104	Novartis Finland Oy	Estonia
Voltaren Akti		434104	Novartis Finland Oy	Estonia
Voltaren Akti		LT/1/94/0948/003	Novartis Finland Oy	Lithuania
Voltaren Akti		LT/1/94/0948/004	Novartis Finland Oy	Lithuania
Voltaren Akti		LT/1/94/0948/005	Novartis Finland Oy	Lithuania
Voltaren Akti		LT/1/94/0948/005	Novartis Finland Oy	Lithuania
Voltaren D		088/04507	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren Dispers		6164351.00.00	Novartis Pharma Gmbh (Art57)	Germany
Voltaren Dolo		18483	Novartis Healthcare A/S	Denmark
Voltaren Dolo		41943	Novartis Healthcare A/S	Denmark
Voltaren Dolo		43250	Novartis Healthcare A/S	Denmark
Voltaren Dolo		43151.00.00 & 64958.00.00	Novartis Consumer Health Gmbh	Germany
Voltaren Dolo		51218.00.00	Novartis Consumer Health Gmbh	Germany
Voltaren Dolo		OGYI-T-5572/08 & OGYI-T-5572/09	Novartis Hungária Kft. Consumer Health	Hungary
Voltaren Dolo		OGYI-T-5572/09	Novartis Hungária Kft. Consumer Health	Hungary
Voltaren Dolo		OGYI-T-5572/21 & OGYI-T-5572/22	Novartis Hungária Kft. Consumer Health	Hungary
Voltaren Dolo		OGYI-T-5572/23 & OGYI-T-5572/24 & OGYI-T-5572/25	Novartis Hungária Kft. Consumer Health	Hungary
Voltaren Dolo		OGYI-T-5572/26 & OGYI-T-5572/27 & OGYI-T-5572/28 & OGYI-T-5572/29 & OGYI-T-5572/30 & OGYI-T-5572/31	Novartis Hungária Kft. Consumer Health	Hungary
Voltaren Dolo		960229 & IS/1/11/037/02	Novartis Healthcare A/S	Iceland

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		IS/1/11/037/01	Novartis Healthcare A/S	Iceland
Voltaren Dolo Liquid		51219.00.00	Novartis Consumer Health Gmbh	Germany
Voltaren Dolo Liquid		82502.00.00	Novartis Consumer Health Gmbh	Germany
Voltaren Fast		251250401 & 251250402 & 251250403	Novartis (Hellas) S.A.C.I. (Art57)	Greece
Voltaren K		RVG 13244	Novartis Consumer Health B.V.	Netherlands
Voltaren K		RVG 20982	Novartis Consumer Health B.V.	Netherlands
Voltaren Rapid		1-19098	Novartis Pharma Gmbh (Art57)	Austria
Voltaren Rapid		07/ 719/10-C	Novartis S.R.O. - Consumer Health Division	Czech Republic
Voltaren Rapid		29/098/91-S/C	Novartis, S.R.O. (Art57)	Czech Republic
Voltaren Rapid		29/173/11-C	Novartis S.R.O. - Consumer Health Division	Czech Republic
Voltaren Rapid		13633	Novartis Healthcare A/S (Art57)	Denmark
Voltaren Rapid		10202 & 10203	Novartis Finland Oy (Art57)	Finland
Voltaren Rapid		930159	Novartis Healthcare A/S (Art57)	Iceland
Voltaren Rapid		4587/2004/01	Novartis Pharma Gmbh (Art57)	Romania
Voltaren Resinat		17982.00.00	Novartis Pharma Gmbh (Art57)	Germany
Voltaren Resinat		0135/08049769	Novartis Pharma Gmbh (Art57)	Luxembourg
Voltaren Retard		1-16856	Novartis Pharma Gmbh (Art57)	Austria
Voltaren Retard		BE122071	Novartis Pharma N.V. (Art57)	Belgium
Voltaren Retard		BE165471	Novartis Pharma N.V. (Art57)	Belgium
Voltaren Retard		29/247/80-C	Novartis, S.R.O. (Art57)	Czech Republic
Voltaren Retard		10661 & 14222	Novartis Healthcare A/S (Art57)	Denmark
Voltaren Retard		061294	Novartis Finland Oy (Art57)	Estonia
Voltaren Retard		10919 & 9228	Novartis Finland Oy (Art57)	Finland

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		14651.00.01	Novartis Pharma Gmbh (Art57)	Germany
Voltaren Retard		94-0182	Novartis Finland Oy (Art57)	Latvia
Voltaren Retard		LT/1/94/1300/002	Novartis Finland Oy (Art57)	Lithuania
Voltaren Retard		2008049766	Novartis Pharma N.V. (Art57)	Luxembourg
Voltaren Retard		2008049766	Novartis Pharma N.V. (Art57)	Luxembourg
Voltaren Retard		MA 088/04510	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren Retard		RVG 15235	Novartis Pharma B.V. (Art57)	Netherlands
Voltaren Retard		9427823	Novartis Farma - Produtos Farmacêuticos S.A. Art57	Portugal
Voltaren Retard		4474/2004/01	Novartis Pharma Gmbh (Art57)	Romania
Voltaren Retard		56562 & 62024	Novartis Farmacéutica S.A. (Art57)	Spain
Voltaren Sr		MA 088/04509	Novartis Pharmaceuticals Uk Limited	Malta
Voltaren Sr		R/1207 & R/6637	Novartis Poland Sp. Z O. O. (Art57)	Poland
Voltaren T		11242 & 12427	Novartis Sverige Ab (Art57)	Sweden
Voltaren® K Migräne		43151.01.00	Novartis Pharma Gmbh (Art57)	Germany
Voltarendolo		355 324-8 & 355 325-4 & 359 411-2 OU 34009 359 411 2 1 & 381 408-0 OU 34009 381 408 0 4 & 381 409-7 OU 34009 381 409 7 2 & 381 410-5 OU 34009 381 410 5 4	Novartis Santé Familiale S.A.S. (Rue Louis Blériot)	France
Voltarene		3400931895299 & 3400931978718 & 3400933814458 & 3400933814519	Novartis Pharma S.A.S. (Art57)	France
Voltarene		3400932214341	Novartis Pharma S.A.S. (Art57)	France
Voltarene		3400932239535 & 3400932273591	Novartis Pharma S.A.S. (Art57)	France
Voltarene		3400932351176	Novartis Pharma S.A.S. (Art57)	France

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		3400932452224 & 3400955283614 & 3400955549444	Novartis Pharma S.A.S. (Art57)	France
Voltarene Lp		3400932460496 & 3400932460557 & 3400933070120 & 3400933070298	Novartis Pharma S.A.S. (Art57)	France
Voltarene Lp		3400933591960 & 3400933592042 & 3400933593452 & 3400933593513 & 3400934595615	Novartis Pharma S.A.S. (Art57)	France
Voltarol		PA 13/87/2	Novartis Pharmaceuticals Uk Limited	Ireland
Voltarol		PA 13/87/3	Novartis Pharmaceuticals Uk Limited	Ireland
Voltarol		09-1063	Novartis Norge As	Norway
Voltarol		09-6597	Novartis Norge As	Norway
Voltarol		99-7996	Novartis Norge As	Norway
Voltarol		PL 00030/0439	Novartis Consumer Health (Uk) Limited	United Kingdom
Voltarol		PL 00101/0466	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0467	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0470	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0472	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0473	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0474 & PL 00101/0475	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0476	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol		PL 00101/0477	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol Joint Pain 12.5mg Tablets		PL 00030/0073	Novartis Consumer Health (Uk)	United Kingdom

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
			Limited	
Voltarol Pain-eze Extra Strength 25mg Tablets		PL 00030/0054	Novartis Consumer Health (Uk) Limited	United Kingdom
Voltarol Rapid		PL 00101/0481 & PL 00101/0482	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltarol Retard		PA 13/87/6	Novartis Pharmaceuticals Uk Limited	Ireland
Voltarol Retard		PA 13/87/7	Novartis Pharmaceuticals Uk Limited	Ireland
Voltarol Sr		PL 00101/0471	Novartis Pharmaceuticals Uk Limited	United Kingdom
Voltfast		20484	Novartis Finland Oy (Art57)	Finland
Voltfast		028945018	Novartis Farma S.P.A. (Art57)	Italy
Voltfast		028945020	Novartis Farma S.P.A. (Art57)	Italy
Voltfast		028945032	Novartis Farma S.P.A. (Art57)	Italy
Voltfast		5905/2013/01 & 5905/2013/02 & 5905/2013/03 & 5905/2013/04	Novartis Pharma GmbH (Art57)	Romania
Voltfast Sachets		PA 13/117/1	Novartis Pharmaceuticals Uk Limited	Ireland
Vóstar-S	DK/H/0582/001, DK/H/0582/002	IS/1/04/012/01, IS/1/04/012/02	Actavis Hf.	Iceland
Zeroflog		034373 011 & 034373 023	Valeas S.P.A.	Italy
Zeroflog		034373 035	Valeas S.P.A.	Italy
волтарен		20020736 & 20020737	Novartis Pharma GmbH (Art57)	Bulgaria
волтарен		20020738	Novartis Pharma GmbH (Art57)	Bulgaria
волтарен		20020739	Novartis Pharma GmbH (Art57)	Bulgaria
волтарен		20020741	Novartis Pharma GmbH (Art57)	Bulgaria
волтарен доло		20030609	Novartis Consumer Health GmbH	Bulgaria
волтарен доло		20100477	Novartis Consumer Health GmbH	Bulgaria

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 ретард		20020740	Novartis Pharma Gmbh (Art57)	Bulgaria
волтфаст		20060381	Novartis Pharma Gmbh (Art57)	Bulgaria

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
Tenoretic film-coated tablets		PL 17901/0049	Astrazeneca Uk Limited	GB
Tenoret		PL 17901/0048	Astrazeneca Uk Limited	GB
Atenolol Clortalidona Qualigen		56661	Qualigen, S.L.	ES
Atenololo Clortalidone Hexal	032805	032805018	Hexal S.P.A	IT
Atenololo Clortalidone Hexal	032805	032805020	Hexal S.P.A	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/001	033455015/M	Sandoz S.P.A.	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/001	033455027/M	Sandoz S.P.A.	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/001	033455054/M	Sandoz S.P.A.	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/002	033455039/M	Sandoz S.P.A.	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/002	033455041/M	Sandoz S.P.A.	IT
Atenololo/Clortalidone Sandoz	NL/H/0161/002	033455104/M	Sandoz S.P.A.	IT
Blokium-Diu		56.665	Almirall, S.A.	ES
Blokium-Diu		OGYI-T-02384	Almirall, S.A.	HU
Diube		024725032	Laboratorio Farmaceutico S.I.T. S.R.L.	IT
Diube		024725069	Laboratorio Farmaceutico S.I.T. S.R.L.	IT
Igroseles		024763056	Ucb Pharma Spa (Milan It)	IT
Igroseles		024763068	Ucb Pharma Spa (Milan It)	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
Atenolol Chlortalidone Sandoz		BE206376	SANDOZ N.V.	BE
Atenolol Chlortalidone Sandoz		BE206385	SANDOZ N.V.	BE
Atenolol comp. Sandoz® 50/12,5 mg		23824.00.00	SANDOZ	DE
Atehexal comp. mite		23826.00.00	HEXAL Ag	DE
Atehexal comp.		23826.01.00	HEXAL Ag	DE
Atenolol comp. Sandoz® 100/25 mg		39680.00.00	SANDOZ	DE
Atecor CT 50mg/12.5mg Film-coated		PA 711/20/1	ROWEX LTD	IE
Atecor CT 100mg/25mg Film-coated		PA 711/20/2	ROWEX LTD	IE
ATENOLOL/CHLOORTALIDON SANDOZ		RVG 15842	SANDOZ B.V.	NL
ATENOLOL/CHLOORTALIDON SANDOZ		RVG 15843	SANDOZ B.V.	NL
Atenolol/chloortalidon Sandoz 50/12,5		RVG 17035	SANDOZ B.V.	NL
Atenolol/chloortalidon Sandoz 100/25		RVG 17036	SANDOZ B.V.	NL