30 May 2024
EMA/302227/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): diclofenac (topical formulations)

Procedure No. PSUSA/00010342/202309

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Algoplast-ratiopharm 140 mg gyógyszeres tapasz	DE/H/1479/001	OGYI-T-20780/01	RATIOPHARM GMBH	HU
Algoplast-ratiopharm 140 mg gyógyszeres tapasz	DE/H/1479/001	OGYI-T-20780/02	RATIOPHARM GMBH	HU
Algoplast-ratiopharm 140 mg gyógyszeres tapasz	DE/H/1479/001	OGYI-T-20780/04	RATIOPHARM GMBH	HU
Algoplast-ratiopharm 140 mg gyógyszeres tapasz	DE/H/1479/001	OGYI-T-20780/03	RATIOPHARM GMBH	HU
Amgesic Muscle & Back Pain Relief 1% Gel	not available	PL 19255/0021	AMDEEPCHA LIMITED	XI
Amgesic Pain Relief 1% Gel	not available	PL 19255/0022	AMDEEPCHA LIMITED	XI
ANTALCALM 140 mg, emplâtre médicamenteux	DE/H/2677/001	34009 276 207 9 9	PIERRE FABRE MEDICAMENT	FR
ANTALCALM 140 mg, emplâtre médicamenteux	DE/H/2677/001	34009 276 205 6 0	PIERRE FABRE MEDICAMENT	FR
ANTALCALM 140 mg, emplâtre médicamenteux	DE/H/2677/001	34009 276 206 2 1	PIERRE FABRE MEDICAMENT	FR
Artrimove 23,2 mg/g gel	not available	5779731	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Artrimove 23,2 mg/g gel	not available	5779756	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Artrimove 23,2 mg/g gel	not available	5779749	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
BrexiAdvance 40 mg/ml spray cutaneo, soluzione	SI/H/0234/001	049582012	NUTRA ESSENTIAL OTC, S.L.	IT
Calminemed 140 mg	BE/H/0244/001	044275016	IBSA FARMACEUTICI ITALIA	IT

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cerotto medicato				
Calminemed 140 mg cerotto medicato	BE/H/0244/001	044275028	IBSA FARMACEUTICI ITALIA	IT
Calminemed 140 mg cerotto medicato	BE/H/0244/001	044275042	IBSA FARMACEUTICI ITALIA	IT
Calminemed 140 mg cerotto medicato	BE/H/0244/001	044275030	IBSA FARMACEUTICI ITALIA	IT
Cinfadol Diclofenaco 11,6 mg/g gel	not available	74.347	LABORATORIOS CINFA, S.A.	ES
Cinfadol SPRAY 39,2 mg/ml solución para pulverización cutánea	not available	85218	LABORATORIOS CINFA, S.A.	ES
DENACLOF Οφθαλμικές σταγόνες, διάλυμα 0,1% (W/V)	not available	9752/8-2-2012	LABORATOIRES THEA	GR
DENACLOF Οφθαλμικές σταγόνες, διάλυμα 0,1% w/v (1 δόση)	not available	104543/22-10-2020	LABORATOIRES THEA	GR
Dicipan, 10 mg/g, gel	not available	5633227	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Dicipan, 10 mg/g, gel	not available	5633235	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Dicipan, 10 mg/g, gel	not available	5633243	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Diclac 20 mg/g gel	HU/H/0745/001	HR-H-618429305	SANDOZ D.O.O.	HR
Diclac 23,2 mg/g gelis	HU/H/0745/001	LT/1/23/5177/002	SANDOZ PHARMACEUTICALS D.D.	LT
Diclac 23,2 mg/g gelis	HU/H/0745/001	LT/1/23/5177/004	SANDOZ PHARMACEUTICALS D.D.	LT
Diclac 23,2 mg/g gelis	HU/H/0745/001	LT/1/23/5177/001	SANDOZ PHARMACEUTICALS D.D.	LT
Diclac 23,2 mg/g gelis	HU/H/0745/001	LT/1/23/5177/003	SANDOZ PHARMACEUTICALS D.D.	LT
Diclac 23,2 mg/g gels	HU/H/0745/001	23-0084	SANDOZ PHARMACEUTICALS D.D.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/01	SANDOZ HUNGÁRIA KFT	HU
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/02	SANDOZ HUNGÁRIA KFT	HU
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/03	SANDOZ HUNGÁRIA KFT	HU
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/04	SANDOZ HUNGÁRIA KFT	HU
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/05	SANDOZ HUNGÁRIA KFT	HU
Diclac Dolo 50 mg/g gél	not available	OGYI-T-23354/06	SANDOZ HUNGÁRIA KFT	HU
Diclac Long 20 mg/g gél	HU/H/0745/001	OGYI-T-23354/07	SANDOZ HUNGÁRIA KFT	HU
Diclac Long 20 mg/g gél	HU/H/0745/001	OGYI-T-23354/08	SANDOZ HUNGÁRIA KFT	HU
Diclac Long 20 mg/g gél	HU/H/0745/001	OGYI-T-23354/09	SANDOZ HUNGÁRIA KFT	HU
Diclac Long 20 mg/g gél	HU/H/0745/001	OGYI-T-23354/10	SANDOZ HUNGÁRIA KFT	HU
Diclo beta Schmerzgel	not available	2200471.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Dicloabak 1 mg/ml Augentropfen, Lösung	FR/H/0296/001	BE291225	LABORATOIRES THEA	BE
Dicloabak 1 mg/ml Augentropfen, Lösung	FR/H/0296/001	2007060067	LABORATOIRES THEA	LU
DICLOABAK 1 mg/ml colírio, solução	FR/H/0296/001	5021373	LABORATOIRES THEA	PT
Dicloabak 1 mg/ml collyre en solution	FR/H/0296/001	BE291225	LABORATOIRES THEA	BE
Dicloabak 1 mg/ml collyre en solution	FR/H/0296/001	2007060067	LABORATOIRES THEA	LU
DICLOABAK 1 mg/ml ocní kapky, roztok	FR/H/0296/001	64/110/11-C	LABORATOIRES THEA	CZ
Dicloabak 1 mg/ml oogdruppels, oplossing	FR/H/0296/001	BE291225	LABORATOIRES THEA	BE
Dicloabak 1 mg/ml oogdruppels, oplossing	FR/H/0296/001	RVG 34473	LABORATOIRES THEA	NL
Dicloabak 1 mg/ml silmätipat, liuos	FR/H/0296/001	29132	LABORATOIRES THEA	FI
Dicloabak 1 mg/ml, ögondroppar, lösning	FR/H/0296/001	29132	LABORATOIRES THEA	FI
Dicloabak 1 mg/ml, ögondroppar, lösning	FR/H/0296/001	44958	LABORATOIRES THEA	SE
DICLOABAK 1 mg/ml, οφθαλμικές σταγόνες, διάλυμα	FR/H/0296/001	46241/14/17-05-2019	LABORATOIRES THEA	GR
DICLOABAK 1mg/ml,	FR/H/0296/001	69475	LABORATOIRES THEA	ES

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colirio en solución				
DICLOABAK, 1 mg/ml, krople do oczu, roztwór	FR/H/0296/001	14116	LABORATOIRES THEA	PL
DICLO-ADGC Schmerzgel forte 20 mg/g Gel	DE/H/7027/002	7003894.00.00	ZENTIVA PHARMA GMBH	DE
DicloAkut forte Schmerzgel 2 % Gel	CZ/H/0935/001	141622	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
DicloAkut Schmerzgel 1 % Gel	not available	139333	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
DICLOCULAR 0,1% collirio, soluzione	not available	028495012	OMNIVISION ITALIA S.R.L.	IT
DICLOCULAR 0,1% collirio, soluzione	not available	028495024	OMNIVISION ITALIA S.R.L.	IT
DICLODENT 0.074%, doseado a 0.74 mg/ml na forma de solução bucal	not available	4881389	LABORATÓRIOS ATRAL, S.A.	PT
DICLODENT 0.074%, doseado a 0.74 mg/ml na forma de solução bucal.	not available	5697321	LABORATÓRIOS ATRAL, S.A.	PT
Diclodolor 140 mg apósitos adhesivos medicamentosos.	DE/H/1479/001	70307	TEVA PHARMA S.L.U.,	ES
Diclofenac "Orifarm", gel	SE/H/1142/001	48080	ORIFARM GENERICS A/S	DK
Diclofenac 1% Gel	PT/H/2250/001	PL 19255/0019	AMDEEPCHA LIMITED	XI
Diclofenac 140 mg medicated plaster	not available	PA1104/005/001	IBSA FARMACEUTICI ITALIA	IE
Diclofenac 2.32% gel	not available	PL 17780/1169	ZENTIVA PHARMA UK LIMITED	XI
Diclofenac AbZ 30 mg/g Gel	DE/H/5138/001	99263.00.00	ABZ-PHARMA GMBH	DE
Diclofenac AbZ Schmerzgel 10 mg/g Gel	DE/H/5493/001	2201672.00.00	ABZ-PHARMA GMBH	DE
Diclofenac AbZ Schmerzgel 10 mg/g Gel	DE/H/6668/001	2206028.00.00	RATIOPHARM GMBH	DE
Diclofenac AbZ Schmerzgel 20 mg/g Gel	DE/H/5493/002	2201673.00.00	ABZ-PHARMA GMBH	DE

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Diclofenac AbZ Schmerzgel 20 mg/g Gel	DE/H/6668/002	2206029.00.00	RATIOPHARM GMBH	DE
Diclofenac acis 30 mg/g Gel	AT/H/0710/001	98529.00.00	ACIS ARZNEIMITTEL GMBH	DE
Diclofenac AL Schmerzgel forte 20 mg/g Gel	DE/H/7329/001	7007225.00.00	ALIUD PHARMA GMBH	DE
DICLOFENAC BGR CONSEIL 2 %, gel	not available	34009 302 762 2 8	BIOGARAN	FR
Diclofenac Bluepharma, 10 mg/g, gel	not available	5565031	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Diclofenac Dermapharm 3% Gel	AT/H/0710/001	137450	DERMAPHARM AG	AT
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml Augentropfen im Einzeldosisbehältnis	NL/H/2859/001	89756.00.00	DEVATIS GMBH	DE
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL

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Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac Devatis 1 mg/ml, oogdruppels, oplosssing, verpakking voor éénmalig gebruik	NL/H/2859/001	RVG 113364	DEVATIS GMBH	NL
Diclofenac diethylamine Teva 11,6 mg/g gel	DE/H/5493/001	19-12809	TEVA B.V	NO
Diclofenac diethylamine Teva 23,2 mg/g gel	DE/H/5493/002	19-12810	TEVA B.V	NO
Diclofenac diethylamine Teva, 10 mg/g, żel	DE/H/5493/001	26295	TEVA B.V	PL
Diclofenac diethylamine Teva, 20 mg/g, żel	DE/H/5493/002	26296	TEVA B.V	PL
Diclofenac EG Forte 20 mg/g gel	DE/H/7329/001	BE661295	EUROGENERICS N.V./S.A.	BE
Diclofenac EG Forte 20 mg/g gel	DE/H/7329/001	BE661295	EUROGENERICS N.V./S.A.	BE
Diclofenac EG Forte 20 mg/g Gel	DE/H/7329/001	BE661295	EUROGENERICS N.V./S.A.	BE
Diclofenac EG Forte 20	DE/H/7329/001	2023060129	EUROGENERICS N.V./S.A.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/g gel				
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574019	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574045	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574021	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574033	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574058	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574060	EG S.P.A.	IT
Diclofenac EG STADA Italia 20 mg/g Gel	DE/H/7329/001	050574072	EG S.P.A.	IT
Diclofenac Eignapharma 20 mg/g gel	CZ/H/0935/001	29/056/19-C	EIGNAPHARMA SLU	CZ
Diclofenac epolamine Mylan 180 mg gyógyszeres tapasz	HU/H/0626/001	OGYI-T-23716/02	MYLAN IRELAND LIMITED	HU
Diclofenac Farmoz, 140 mg, emplastro medicamentoso	not available	5809223	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Diclofenac Farmoz, 140 mg, emplastro medicamentoso	not available	5809231	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Diclofenac Fidia 140 mg wirkstoffhaltiges Pflaster	DE/H/4645/001	96683.00.00	FIDIA FARMACEUTICI S.P.A	DE
Diclofenac HEUMANN Gel Gel mit 10 mg Diclofenac-Natrium/g	not available	19199.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Diclofenac IBSA Farmaceutici 140 mg, compresse imprégnée	not available	BE230404	IBSA FARMACEUTICI ITALIA	BE
Diclofenac IBSA Farmaceutici 140 mg, compresse imprégnée	not available	20100040763-0895199	IBSA FARMACEUTICI ITALIA	LU
Diclofenac IBSA	not available	BE230404	IBSA FARMACEUTICI ITALIA	BE

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Farmaceutici 140 mg, geïmpregneerd verbandgaas				
DICLOFENAC IBSA FARMACEUTICI 140 mg, imprägnierte Verbände	not available	BE230404	IBSA FARMACEUTICI ITALIA	BE
Diclofenac Liderlens 40 mg/ml cutaneous spray solution	SI/H/0234/001	PA23205/001/001	NUTRA ESSENTIAL OTC, S.L.	IE
DICLOFENAC LIDERLENS 40 mg/mL, solution pour pulvérisation cutanée	SI/H/0234/001	34009 302 617 3 6	NUTRA ESSENTIAL OTC, S.L.	FR
Diclofenac Linn 2,32% extra sterk, gel	SE/H/2259/001	RVG 129843	MAE HOLDING BV	NL
Diclofenac Mylan 180 mg vaistinis pleistras	HU/H/0626/001	LT/1/20/4605/001	MYLAN IRELAND LIMITED	LT
Diclofenac Mylan 180 mg vaistinis pleistras	HU/H/0626/001	LT/1/20/4605/002	MYLAN IRELAND LIMITED	LT
Diclofenac Mylan Pharma 180 mg cerotto medicato	IT/H/0559/001	045954017	MYLAN S.P.A.	IT
Diclofenac Mylan Pharma 180 mg cerotto medicato	IT/H/0559/001	045954029	MYLAN S.P.A.	IT
Diclofenac Mylan, 180 mg ravimplaaster	HU/H/0626/001	1008820	MYLAN IRELAND LIMITED	EE
Diclofenac Nocegap 10 mg/g Gel	PT/H/2250/001	5862735	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Diclofenac Nocegap 10 mg/g Gel	PT/H/2250/001	5862750	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Diclofenac Nocegap 10 mg/g Gel	PT/H/2250/001	5862727	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Diclofenac Nocegap 10 mg/g Gel	PT/H/2250/001	5862743	PHAGECON-SERVICOS E CONSULTORIA FARMACEUTICA	PT
Diclofenac Nutra Essential 40 mg/ml kožní	SI/H/0234/001	29/126/21-C	NUTRA ESSENTIAL OTC, S.L.	CZ

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sprej, roztok				
Diclofenac Nutra Essential 8 mg/Sprühstob Spray zur Anwendung auf der Haut, Lösung	SI/H/0234/001/DC	7005209.00.00	NUTRA ESSENTIAL OTC, S.L.	DE
Diclofenac Orifarm 11,6 mg/g gel	SE/H/1142/001	45481	ORIFARM GENERICS A/S	SE
Diclofenac Patch AB 140 mg emplâtre médicamenteux.	IT/H/0562/001	BE541715	AUROBINDO N.V.	BE
Diclofenac Patch AB 140 mg pleister.	IT/H/0562/001	BE541715	AUROBINDO N.V.	BE
Diclofenac Patch AB 140 mg wirkstoffhaltiges Pflaster	IT/H/0562/001	BE541715	AUROBINDO N.V.	BE
Diclofenac Patch EG 140 mg emplâtre médicamenteux	DE/H/4645/001	BE518631	EUROGENERICS N.V./S.A.	BE
Diclofenac Patch EG 140 mg emplâtre médicamenteux	DE/H/4645/001	2018060174	EUROGENERICS N.V./S.A.	LU
Diclofenac Patch EG 140 mg pleister	DE/H/4645/001	BE518631	EUROGENERICS N.V./S.A.	BE
Diclofenac Patch EG 140 mg wirkstoffhaltiges Pflaster	DE/H/4645/001	BE518631	EUROGENERICS N.V./S.A.	BE
Diclofenac Pharmakern 23,2 mg/g gel	not available	5752001	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Diclofenac Pharmakern 23,2 mg/g gel	not available	5751979	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Diclofenac Pharmakern 23,2 mg/g gel	not available	5751961	PHARMAKERN PORTUGAL – PRODUTOS	PT

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			FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	
DICLOFENAC PHARMAKI 1 %, gel	not available	NL53901	PHARMAKI GENERICS LTD	FR
Diclofenac ratiopharm 11,6 mg/g geeli	DE/H/5493/001	36997	TEVA B.V	FI
Diclofenac ratiopharm 11,6 mg/g gel	DE/H/5493/001	36997	TEVA B.V	FI
Diclofenac ratiopharm 23,2 mg/g geeli	DE/H/5493/002	36998	TEVA B.V	FI
Diclofenac ratiopharm 23,2 mg/g gel	DE/H/5493/002	36998	TEVA B.V	FI
Diclofenac Sandoz 20 mg/g gel	HU/H/0745/001	BE661232	SANDOZ N.V.	BE
Diclofenac Sandoz 50 mg/g gél	not available	OGYI-T-23245/01	SANDOZ HUNGÁRIA KFT	HU
Diclofenac Sandoz 50 mg/g gél	not available	OGYI-T-23245/02	SANDOZ HUNGÁRIA KFT	HU
Diclofenac Sodium 1% Gel	not available	PL 19255/0021	AMDEEPCHA LIMITED	XI
Diclofenac Sodium 1% Gel	not available	PL 19255/0022	AMDEEPCHA LIMITED	XI
Diclofenac sodium Nutra Essential, 40 mg/mL, aerozol na skóre, roztwór	SI/H/0234/001	27633	NUTRA ESSENTIAL OTC, S.L.	PL
Diclofenac Sodium Teva 1% w/w Gel	DE/H/5493/001	PA1986/093/001	TEVA B.V	IE
Diclofenac Sodium Teva 2% w/w Gel	DE/H/5493/002	PA1986/093/002	TEVA B.V	IE
Diclofenac STADA Schmerzgel forte 20 mg/g Gel	DE/H/7593/001	7010363.00.00	STADA GMBH	DE
Diclofenac STADA® 23,2 mg/g gel	not available	58771	STADA ARZNEIMITTEL AG	SE
Diclofenac Terapia 10 mg/g cremă	not available	11640/2019/01	TERAPIA S.A.	RO
Diclofenac Terapia 10	not available	11640/2019/02	TERAPIA S.A.	RO

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mg/g cremă				
Diclofenac Terapia 10 mg/g gel	not available	6513/2014/01	TERAPIA S.A.	RO
Diclofenac Terapia 50 mg/g gel	not available	11641/2019/01	TERAPIA S.A.	RO
Diclofenac Terapia 50 mg/g gel	not available	11641/2019/02	TERAPIA S.A.	RO
Diclofenac Teva 11,6 mg/g hlaup	DE/H/5493/001	IS/1/20/055/01	TEVA B.V	IS
DICLOFENAC TEVA 140 mg cerotti medicati	DE/H/1481/001	038721027	TEVA B.V	IT
DICLOFENAC TEVA 140 mg cerotti medicati	DE/H/1481/001	038721015	TEVA B.V	IT
Diclofenac Teva 23,2 mg/g hlaup	DE/H/5493/002	IS/1/20/055/02	TEVA B.V	IS
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883018	TEVA B.V	IT
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883020	TEVA B.V	IT
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883032	TEVA B.V	IT
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883044	TEVA B.V	IT
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883057	TEVA B.V	IT
Diclofenac Teva BV 10 mg/g gel	DE/H/5493/001	047883069	TEVA B.V	IT
Diclofenac Teva BV 20 mg/g gel	DE/H/5493/002	047883071	TEVA B.V	IT
Diclofenac Teva BV 20 mg/g gel	DE/H/5493/002	047883083	TEVA B.V	IT
Diclofenac Teva BV 20 mg/g gel	DE/H/5493/002	047883095	TEVA B.V	IT
Diclofenac Teva BV 20 mg/g gel	DE/H/5493/002	047883107	TEVA B.V	IT
Diclofenac Teva BV 20 mg/g gel	DE/H/5493/002	047883119	TEVA B.V	IT
Diclofenac Teva BV 20	DE/H/5493/002	047883121	TEVA B.V	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/g gel				
DICLOFENAC TEVA CONSEIL 1 %, emplâtre médicamenteux	not available	NL37788	RATIOPHARM GMBH	FR
Diclofenac Teva Extra Sterk 2,32%, gel	not available	RVG 130435	TEVA B.V	NL
Diclofenac Teva Extra Sterk 2,32%, gel	not available	RVG 130431	TEVA NEDERLAND B.V.	NL
DICLOFENAC TEVA SANTE CONSEIL 1 %, gel	DE/H/6668/001	NL52461	TEVA B.V	FR
DICLOFENAC TEVA SANTE CONSEIL 2 %, gel	DE/H/6668/002	NL52462	TEVA B.V	FR
Diclofenac-Natrium Stragen 10 mg/g Gel	not available	2200473.00.00	FAIRMED HEALTHCARE GMBH	DE
Diclofenaco Abamed 1 mg/ml colirio en solución en envase unidosis.	not available	77503	QUALIX PHARMA S.L.	ES
Diclofenaco Kern Pharma 11,6 mg/g gel	not available	74.119	KERN PHARMA, S.L.	ES
Diclofenaco ratiopharm 30 mg/g gel	DE/H/5130/001	83055	TEVA PHARMA S.L.U.,	ES
Diclofenaco Teva 30 mg/g gel	DE/H/5138/001	83054	TEVA B.V	ES
Diclofenaco-lepori 1 mg/ml colirio en solución	not available	61.812	ANGELINI PHARMA ESPANA S.L	ES
Diclofenaco-lepori 1 mg/ml colirio en solución	not available	61.812	ANGELINI PHARMA ESPANA S.L	ES
Diclofenaco-lepori 1 mg/ml colirio en solución en envase unidosis	not available	63.643	ANGELINI PHARMA ESPANA S.L	ES
Diclofenaco-lepori 1 mg/ml colirio en solución en envase unidosis	not available	63.643	ANGELINI PHARMA ESPANA S.L	ES
Diclofenac-ratiopharm Schmerzpflaster 140 mg wirkstoffhaltiges Pflaster	DE/H/1479/001	62430.00.00	RATIOPHARM GMBH	DE
Diclofenac-ratiopharm® 30 mg/g Gel	DE/H/5130/001	99197.00.00	RATIOPHARM GMBH	DE

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Diclofenac-ratiopharm® 30 mg/g Gel	DE/H/5130/001	2018110319	RATIOPHARM GMBH	LU
DICLOFTIL "0,1% collirio, soluzione"	not available	029041011	FARMIGEA SPA	IT
Dicloftil "0,1% collirio, soluzione" - 30 contenitori monodose 0,5 ml	not available	029041023	FARMIGEA SPA	IT
DICLOFTIL 0,5 mg/0,5 ml colírio, solução em recipientes unidose.	PT/H/1749/001	5757067	FARMIGEA SPA	PT
Dicloftil 1 mg/ml collirio, soluzione	not available	029041035	FARMIGEA SPA	IT
Diclogel, 10 mg/g, żel	not available	26957	FORTIS PHARMACEUTICALS SP. Z O. O.	PL
Diclogel, 10 mg/g, żel	not available	26957	FORTIS PHARMACEUTICALS SP. Z O. O.	PL
Diclokern 11,6 mg/g gel	not available	76.209	KERN PHARMA, S.L.	ES
Diclokern Spray 39,2 mg/ml solución para pulverización cutánea	not available	86.279	KERN PHARMA, S.L.	ES
Diclokern® Forte 23,2 mg/g gel.	not available	83.900	KERN PHARMA, S.L.	ES
Dicloklaph 10 mg/g Gel	not available	2200472.00.00	AENOVA IP GMBH	DE
DicloMAX Mobilat, 20 mg/g, żel	DE/H/7329/001	28033	STADA ARZNEIMITTEL AG	PL
DICLOMED 0,3 mg/spruzzo, spray per mucosa orale	not available	032085033	FARMAKA S.R.L.	IT
DICLOMED 0,74 mg/ml Collutorio	not available	032085019	FARMAKA S.R.L.	IT
Diclomel Max Strength 2% w/w gel	DE/H/7329/001	PA0126/372/001	CLONMEL HEALTHCARE LTD.	IE
Diclomel Max Strength 2% w/w gel	DE/H/7329/001	MA1045/05501	CLONMEL HEALTHCARE LTD.	MT
Diclo-Mepha Schmerzpflaster 140 mg wirkstoffhaltiges Pflaster	DE/H/1481/001	70741.00.00	TEVA B.V	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
DICLOPLAST	not available	2309/14-2-2022	ANGELINI PHARMA HELLAS S.A.	GR
DICLOPLAST	not available	2309/14-2-2022	ANGELINI PHARMA HELLAS S.A.	GR
DICLOPLAST	not available	2309/14-2-2022	ANGELINI PHARMA HELLAS S.A.	GR
DICLOPLAST	not available	2309/14-2-2022	ANGELINI PHARMA HELLAS S.A.	GR
DICLORAL	not available	2451901	NESTORAS VLACHOS PC EPSILON HEALTH	GR
Dicloreum 30 mg/g spumă cutanată	not available	10603/2018/01	ALFASIGMA S.P.A.	RO
DICLOREUM ACTIGEL 1 % gel	not available	035450028	ALFASIGMA S.P.A.	IT
DICLOREUM ACTIGEL 1 % gel	not available	035450016	ALFASIGMA S.P.A.	IT
DICLOREUM ANTINFIAMMATORIO LOCALE 180 mg cerotti medicati	not available	042685026	ALFASIGMA S.P.A.	IT
DICLOREUM ANTINFIAMMATORIO LOCALE 180 mg cerotti medicati	not available	042685038	ALFASIGMA S.P.A.	IT
DICLOREUM ANTINFIAMMATORIO LOCALE 180 mg cerotti medicati	not available	042685014	ALFASIGMA S.P.A.	IT
DICLOREUM ANTINFIAMMATORIO LOCALE 3% Schiuma cutanea, contenitore sotto pressione	not available	042685040	ALFASIGMA S.P.A.	IT
Diclostim, 0,74 mg/mL (0,074%), roztwór do płukania gardła / jamy ustnej	not available	27044	SOLINEA SP. Z O.O. SP. K.	PL
Diclo-Vision sine 1 mg/ml Augentropfen, Lösung im	AT/H/0528/001	1-31525	OMNIVISION GMBH	AT

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Einzeldosisbehältnis				
Diclo-Vision sine 1 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0528/001	84680.00.00	OMNIVISION GMBH	DE
Diclox 10 mg/g Gel	DE/H/6245/001	2203844.00.00	RATIOPHARM GMBH	DE
Diclox forte 20 mg/g Gel	DE/H/6245/002	2203845.00.00	RATIOPHARM GMBH	DE
Diklofenak ABECE 11,6 mg/g gel	not available	55821	EVOLAN PHARMA AB	SE
Diklofenak Apofri 11,6 mg/g gel	not available	46780	EVOLAN PHARMA AB	SE
Diklofenak Apofri 11,6 mg/g hlaup.	SE/H/1873/001	IS/1/18/111/01	EVOLAN PHARMA AB	IS
Diklofenak Apofri 23,2 mg/g gel	not available	55648	EVOLAN PHARMA AB	SE
Diklofenak Mimer 23,2 mg/g gel	not available	58770	MIMER MEDICAL AB	SE
Diklofenak NET 23,2 mg/g gel	not available	55649	EVOLAN PHARMA AB	SE
Diklofenak Nutra Essential 40 mg/ml dermalno prsilo, raztopina	SI/H/0234/001	H/22/02923/001	NUTRA ESSENTIAL OTC, S.L.	SI
Diklofenak Teva 11,6 mg/g gel	DE/H/6668/001	60736	TEVA B.V	SE
Diklofenak Teva 23,2 mg/g gel	DE/H/6668/002	60737	TEVA B.V	SE
Diklofenak Viatris, 180 mg, plaster leczniczy (1,3%)	HU/H/0626/001	27498	VIATRIS LIMITED	PL
Diklofenakdietylamin Norfri 11,6 mg/g (1,16%) gel.	not available	17-11511	EVOLAN PHARMA AB	NO
Diklofenakdietylamin Norfri 23,2 mg/g (2,32 %) gel.	not available	17-11512	EVOLAN PHARMA AB	NO
Diklofenak-dietylamín Teva 11,6 mg/g gél	DE/H/5493/001	29/0160/20-S	TEVA B.V	SK
Diklofenak-dietylamín	DE/H/5493/002	29/0161/20-S	TEVA B.V	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Teva 23,2 mg/g gél				
Diltidol 10 mg/g gel	not available	73.473	ARISTO PHARMA IBERIA, S.L.	ES
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/002	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/004	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/005	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/003	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/001	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/007	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gelis	DE/H/7329/001	LT/1/23/5171/006	STADA ARZNEIMITTEL AG	LT
Ditel 23,2 mg/g gels	DE/H/7329/001	23-0085	STADA ARZNEIMITTEL AG	LV
Ditel, 23,2 mg/g geel	DE/H/7329/001	1109823	STADA ARZNEIMITTEL AG	EE
Dolifen 11,6 mg/g gel	not available	59.391	FAES FARMA, S.A.	ES
EVINOPON 16mg/ml δερματικό διάλυμα	not available	11054	BROS LTD	GR
FASTUFLEX 180 mg cerotto medicato.	IT/H/0561/001	045952013	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
FASTUFLEX 180 mg cerotto medicato.	IT/H/0561/001	045952025	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
FASTUM ANTIDOLORIFICO 10 mg/g GEL	not available	040657025	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
FASTUM ANTIDOLORIFICO 10 mg/g GEL	not available	040657013	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Flalgo 140 mg wirkstoffhaltiges Pflaster	BE/H/0244/001	BE498693	IBSA FARMACEUTICI ITALIA	BE
Flalgo 140 mg, léčivá náplast	BE/H/0244/001	29/402/16-C	IBSA SLOVAKIA S.R.O.	CZ
Flalgo emplâtre médicamenteux 140 mg	BE/H/0244/001	BE498693	IBSA FARMACEUTICI ITALIA	BE
Flalgo, 140 mg, pleister	BE/H/0244/001	BE498693	IBSA FARMACEUTICI ITALIA	BE
Flectoflex 140 mg wirkstoffhaltiges Pflaster	BE/H/0245/001	BE498666	IBSA FARMACEUTICI ITALIA	BE
Flectoflex, 140 mg, pleister	BE/H/0245/001	BE498666	IBSA FARMACEUTICI ITALIA	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
Flectoflex, 140mg emplâtre médicamenteux	BE/H/0245/001	BE498666	IBSA FARMACEUTICI ITALIA	BE
Flectopar	BE/H/0244/001	29/0392/16-S	IBSA SLOVAKIA S.R.O.	SK
Flector náplast	not available	29/0472/94-S	IBSA SLOVAKIA S.R.O.	SK
FLECTOR 1 POUR CENT, gel	not available	34009 333 846 1 6	IBSA PHARMA SAS	FR
FLECTOR 1 POUR CENT, gel	not available	34009 333 847 8 4	IBSA PHARMA SAS	FR
FLECTOR 1 POUR CENT, gel	not available	34009 333 848 4 5	IBSA PHARMA SAS	FR
FLECTOR 1 POUR CENT, gel	not available	34009 377 842 1 4	IBSA PHARMA SAS	FR
Flector 10 mg/g gel	not available	29/350/96-C	IBSA SLOVAKIA S.R.O.	CZ
Flector 10 mg/g gél	not available	OGYI-T-5033/01	IBSA PHARMA KFT	HU
Flector 10 mg/g gél	not available	OGYI-T-5033/02	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/05	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/06	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/07	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/08	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/09	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/10	IBSA PHARMA KFT	HU
Flector 140 mg gyógyszeres tapasz	not available	OGYI-T-5033/13	IBSA PHARMA KFT	HU
Flector 140 mg, medicinskt plåster	FR/H/0225/001	19221	IBSA FARMACEUTICI ITALIA	SE
FLECTOR 180 mg cerotto medicato	not available	027757032	IBSA FARMACEUTICI ITALIA	IT
FLECTOR 180 mg cerotto medicato	not available	027757069	IBSA FARMACEUTICI ITALIA	IT
FLECTOR 180 mg cerotto medicato	not available	027757044	IBSA FARMACEUTICI ITALIA	IT
FLECTOR 180 mg cerotto	not available	027757071	IBSA FARMACEUTICI ITALIA	IT

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medicato				
Flector 180 mg léčivá náplast	not available	29/361/96-C	IBSA SLOVAKIA S.R.O.	CZ
Flector EP - Pflaster	not available	1-23243	SANOVA PHARMA GESMBH	AT
Flector Extra 10 mg/g gél	not available	OGYI-T-5033/11	IBSA PHARMA KFT	HU
Flector Extra 10 mg/g gél	not available	OGYI-T-5033/12	IBSA PHARMA KFT	HU
Flector gél	not available	29/0471/94-S	IBSA SLOVAKIA S.R.O.	SK
Flector Schmerzpflaster 1%	FR/H/0267/001	62521.00.00	IBSA FARMACEUTICI ITALIA	DE
Flector Tissugel 1 % compresses imprégnées.	not available	BE208537	IBSA FARMACEUTICI ITALIA	BE
Flector Tissugel 1 % compresses imprégnées.	not available	2006038467-0302256	IBSA FARMACEUTICI ITALIA	LU
Flector Tissugel 1 % imprägnierte Verbände	not available	BE 208537	IBSA FARMACEUTICI ITALIA	BE
Flector Tissugel 1% geïmpregneerde verbandgazen.	not available	BE 208537	IBSA FARMACEUTICI ITALIA	BE
Flector Tissugel 1% geïmpregneerde verbandgazen.	not available	2006038467-0302256	IBSA FARMACEUTICI ITALIA	LU
FLECTOR TISSUGEL 140 mg , φαρμακευτικὸ ἔμπλαστρο	FR/H/0225/001	20149	IBSA FARMACEUTICI ITALIA	CY
Flector Tissugel 140 mg emplastro medicamentoso	not available	3231495	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Flector Tissugel 140 mg emplastro medicamentoso	not available	3231594	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Flector Tissugel 140 mg emplastro medicamentoso	not available	3231693	BGP PRODUCTS UNIPESSOAL, LDA.	PT
FLECTOR TISSUGEL 140 mg, emplâtre médicamenteux	FR/H/0225/001	34009 339 064 5 0	IBSA PHARMA SAS	FR
FLECTOR TISSUGEL 140 mg, emplâtre	FR/H/0225/001	34009 337 830 2 0	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
médicamenteux				
FLECTOR TISSUGEL 140 mg, emplâtre médicamenteux	FR/H/0225/001	34009 337 831 9 8	IBSA PHARMA SAS	FR
FLECTOR TISSUGEL 140 mg, emplâtre médicamenteux	FR/H/0225/001	34009 337 832 5 9	IBSA PHARMA SAS	FR
Flector Tissugel 140mg Medicated Plaster	FR/H/0225/001	PA1104/004/001	IBSA FARMACEUTICI ITALIA	IE
FLECTOR TISSUGEL 140mg medicated plaster	FR/H/0267/001	PL 21039/0004	IBSA FARMACEUTICI ITALIA	XI
Flector Tissugel, 1%, plaster leczniczy (Diclofenacum epolaminum)	FR/H/0225/001	11803	IBSA FARMACEUTICI ITALIA	PL
Flector, medicinsk plaster	FR/H/0225/001	34600	IBSA FARMACEUTICI ITALIA	DK
Flectorin 140 mg gyógyszeres tapasz	BE/H/0244/001	OGYI-T-23051/01	IBSA PHARMA KFT	HU
Flectorin 140 mg gyógyszeres tapasz	BE/H/0244/001	OGYI-T-23051/02	IBSA PHARMA KFT	HU
Flectorin 140 mg gyógyszeres tapasz	BE/H/0244/001	OGYI-T-23051/03	IBSA PHARMA KFT	HU
Flectorin 140 mg gyógyszeres tapasz	BE/H/0244/001	OGYI-T-23051/04	IBSA PHARMA KFT	HU
Flectormed 180 mg apósito adhesivo medicamentoso	BE/H/0245/001	82182	IBSA FARMACEUTICI ITALIA	ES
GLIMBAX 0,074 g/100 ml apă de gură	not available	12212/2019/01	FARMAKA S.R.L.	RO
GLIMBAX 0,074 g/100 ml orálny roztok	not available	95/0200/04-S	ANGELINI PHARMA ÖSTERREICH GMBH	SK
GLIMBAX 0,074 g/100 ml orálny roztok	not available	95/0200/04-S	ANGELINI PHARMA ÖSTERREICH GMBH	SK
Glimbax, 0,74 mg/ml (0,074%), roztwór do płukania jamy ustnej i gardła	not available	14588	ANGELINI PHARMA POLSKA SP. Z O.O.	PL
Glimbax, 0,74 mg/ml	not available	14588	ANGELINI PHARMA POLSKA	PL

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(0,074%), roztwór do płukania jamy ustnej i gardła			SP. Z O.O.	
Holactiv 10 mg/g gel	not available	5670716	HOLON S.A.	PT
Holactiv 10 mg/g gel	not available	5670708	HOLON S.A.	PT
Holactiv 10 mg/g gel	not available	5670674	HOLON S.A.	PT
Ilmodol dolori articolari e muscolari 180 mg cerotto medicato	IT/H/0560/001	045953015	FARMITALIA S.R.L.	IT
Ilmodol dolori articolari e muscolari 180 mg cerotto medicato	IT/H/0560/001	045953027	FARMITALIA S.R.L.	IT
Irfen 11,6 mg/g gel	DE/H/6668/001	86651	TEVA B.V	ES
Irfen Forte 23,2 mg/g gel	DE/H/6668/002	86650	TEVA B.V	ES
Itami 140 mg medicated plaster	DE/H/4645/001/DC	PA0814/002/001	FIDIA FARMACEUTICI S.P.A	IE
Itami 140 mg wirkstoffhaltiges pflaster	DE/H/2678/001	80421.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg wirkstoffhaltiges Pflaster	DE/H/2678/001	80421.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg wirkstoffhaltiges Pflaster	DE/H/2678/001	80421.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg wirkstoffhaltiges pflaster	DE/H/2677/001	80420.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg wirkstoffhaltiges Pflaster	DE/H/2677/001	80420.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg wirkstoffhaltiges Pflaster	DE/H/2677/001	80420.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itami 140 mg zdravilni obliž	DE/H/4645/001	H/18/02423/002	FIDIA FARMACEUTICI S.P.A	SI
Itami 140 mg zdravilni obliž	DE/H/4645/001	H/18/02423/003	FIDIA FARMACEUTICI S.P.A	SI
Itami 140 mg zdravilni obliž	DE/H/4645/001	H/18/02423/001	FIDIA FARMACEUTICI S.P.A	SI
Itami Diario 140 mg apósito adhesivo medicamentoso	DE/H/7389/001	88591	FIDIA FARMACEUTICI S.P.A	ES
Itami Diario 140 mg	DE/H/7389/001	88591	FIDIA FARMACEUTICI S.P.A	ES

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apósito adhesivo medicamentoso				
Itami Diario 140 mg apósito adhesivo medicamentoso	DE/H/7389/001	88591	FIDIA FARMACEUTICI S.P.A	ES
Itami Diario 140 mg apósito adhesivo medicamentoso	DE/H/7389/001	88591	FIDIA FARMACEUTICI S.P.A	ES
Itami Unidie 140 mg Cerotto Medicato	DE/H/7389/001/MR	050400011	FIDIA FARMACEUTICI S.P.A	IT
Itami Unidie 140 mg Cerotto Medicato	DE/H/7389/001	050400023	FIDIA FARMACEUTICI S.P.A	IT
Itami Unidie 140 mg Cerotto Medicato	DE/H/7389/001/MR	050400035	FIDIA FARMACEUTICI S.P.A	IT
Itami Unidie 140 mg Cerotto Medicato	DE/H/7389/001/MR	050400047	FIDIA FARMACEUTICI S.P.A	IT
Itami 140 mg liečivá náplasť	DE/H/7389/001	29/0068/23-S	FIDIA FARMACEUTICI S.P.A	SK
Itami 140 mg liečivá náplasť	DE/H/7389/001	29/0068/23-S	FIDIA FARMACEUTICI S.P.A	SK
Itami 140 mg liečivá náplasť	DE/H/7389/001	29/0068/23-S	FIDIA FARMACEUTICI S.P.A	SK
Itami 140 mg liečivá náplasť	DE/H/7389/001	29/0068/23-S	FIDIA FARMACEUTICI S.P.A	SK
Itami, 140 mg, plaster leczniczy	DE/H/2678/001	21301	FIDIA FARMACEUTICI S.P.A	PL
Itami, 140 mg, plaster leczniczy	DE/H/2678/001	21301	FIDIA FARMACEUTICI S.P.A	PL
Itami, 140 mg, plaster leczniczy	DE/H/2678/001	21301	FIDIA FARMACEUTICI S.P.A	PL
Itamiact 140 mg léčivá náplast	DE/H/7389/001	29/145/22-C	FIDIA FARMACEUTICI S.P.A	CZ
Itamiact 140 mg léčivá náplast	DE/H/7389/001	29/145/22-C	FIDIA FARMACEUTICI S.P.A	CZ
Itamiact 140 mg léčivá náplast	DE/H/7389/001	29/145/22-C	FIDIA FARMACEUTICI S.P.A	CZ
Itamiact 140 mg léčivá náplast	DE/H/7389/001	29/145/22-C	FIDIA FARMACEUTICI S.P.A	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Itamione 140 mg wirkstoffhaltiges Pflaster	DE/H/7389/001	7004505.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itamione 140 mg wirkstoffhaltiges Pflaster	DE/H/7389/001	7004505.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itamione 140 mg wirkstoffhaltiges Pflaster	DE/H/7389/001	7004505.00.00	FIDIA FARMACEUTICI S.P.A	DE
Itamione 140 mg wirkstoffhaltiges Pflaster	DE/H/7389/001/MR	7004505.00.00	FIDIA FARMACEUTICI S.P.A	DE
Kinespir 10 mg/g gel	DE/H/5493/001	BE567413	TEVA B.V	BE
Kinespir 10 mg/g Gel	DE/H/5493/001	BE567413	TEVA B.V	BE
Kinespir 10 mg/g gel	DE/H/5493/001	BE567413	TEVA B.V	BE
Kinespir Forte 20 mg/g gel	DE/H/5493/002	BE567422	TEVA B.V	BE
Kinespir Forte 20 mg/g gel	DE/H/5493/002	BE567422	TEVA B.V	BE
Kinespir Forte 20 mg/g Gel	DE/H/5493/002	BE567422	TEVA B.V	BE
Kinespir Patch 140 mg emplâtre médicamenteux	DE/H/1479/001	BE322016	TEVA PHARMA BELGIUM N.V./S.A	BE
Kinespir Patch 140 mg pleister	DE/H/1479/001	BE322016	TEVA PHARMA BELGIUM N.V./S.A	BE
KINESPIR PATCH 140 mg WIRKSTOFFHALTIGES PFLASTER	DE/H/1479/001	BE322016	TEVA PHARMA BELGIUM N.V./S.A	BE
Kruidvat Diclofenac 2,32% extra sterk, gel	not available	RVG 130455	MAE HOLDING BV	NL
Kruidvat Diclofenac 2,32% extra sterk, gel	not available	RVG 130455	MAE HOLDING BV	NL
Lokosol 10 mg/g gel	not available	HR-H-419662714	SALVUS D.O.O.	HR
MINIFLAM® 1,5 w/w Δερματικό διάλυμα	not available	64542/08.07.2021	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
MINIFLAMPATCH 180 mg φαρμακούχο έμπλαστρο	IT/H/0562/001	23659/22-03-2022	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
Motusol 1.16% w/w Gel	not available	PL 00289/2473	TEVA UK LIMITED	XI
Motusol MAX 12-Hour Pain Relief 2.32% w/w	not available	PL 00289/2538	TEVA UK LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gel				
Motusol Max 2.32% w/w Gel	not available	PL 00289/2474	TEVA UK LIMITED	XI
Naclof Unidose, oogdruppels 1 mg/ml.	not available	RVG 16483	LABORATOIRES THEA	NL
Naclof, 1 mg/ml, krople do oczu, roztwór	not available	R/1453	LABORATOIRES THEA	PL
Naclof, oogdruppels 1 mg/ml	not available	RVG 12800	LABORATOIRES THEA	NL
Olfen 10 mg/g gel	CZ/H/1066/001	HR-H-313859076	PLIVA HRVATSKA D.O.O.	HR
Olfen 140 mg emplastro medicamentoso	DE/H/1480/001	5128723	TEVA B.V	PT
Olfen 140 mg emplastro medicamentoso	DE/H/1480/001	5128715	TEVA B.V	PT
Olfen 140 mg léčivé náplasti léčivá náplast	DE/H/1479/001	29/509/08-C	RATIOPHARM GMBH	CZ
Olfen 140 mg liečivá náplasť	DE/H/1479/001	29/0426/08-S	TEVA B.V	SK
Olfen 20 mg/g gel	CZ/H/1066/002	HR-H-841350554	PLIVA HRVATSKA D.O.O.	HR
Olfen 23,2 mg/g gels	DE/H/5493/002	20-0086	MEPHA LDA	LV
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/001	MEPHA LDA	LT
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/002	MEPHA LDA	LT
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/003	MEPHA LDA	LT
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/004	MEPHA LDA	LT
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/005	MEPHA LDA	LT
Olfen 23,2 mg/g gelis	DE/H/5493/002	LT/1/20/4613/006	MEPHA LDA	LT
Olfen Artic 10 mg/g gel	DE/H/5493/001	5797337	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Olfen Artic 10 mg/g gel	DE/H/5493/001	5797352	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Olfen Artic 10 mg/g gel	DE/H/5493/001	5797345	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Olfen Forte 23,2 mg/g gél	CZ/H/1066/002	29/0159/20-S	TEVA B.V	SK
Olfen Max, 20 mg/g, żel	CZ/H/1066/002	26078	TEVA B.V	PL
Olfen Neo 10 mg/g gel	CZ/H/1066/001	29/459/19-C	TEVA B.V	CZ
Olfen Neo 11,6 mg/g gél	CZ/H/1066/001	29/0158/20-S	TEVA B.V	SK
Olfen Neo Forte 20 mg/g	CZ/H/1066/002	29/460/19-C	TEVA B.V	CZ

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gel				
Olfen Patch 140 mg plaster leczniczy	DE/H/1480/001	15463	TEVA B.V	PL
Olfen Patch 140 mg wirkstoffhaltiges Pflaster	DE/H/1480/001	70740.00.00	TEVA B.V	DE
Olfen żel, 10 mg/g, żel	CZ/H/1066/001	26077	TEVA B.V	PL
Olfen, 23,2 mg/g geel	DE/H/5493/002	1009420	MEPHA LDA	EE
Olfenex Artic 23,2 mg/g gel	DE/H/5493/002	5797360	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Olfenex Artic 23,2 mg/g gel	DE/H/5493/002	5797378	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Olfenex Artic 23,2 mg/g gel	DE/H/5493/002	5797402	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
PENNSAID 16 mg/ml soluzione cutanea	EL/H/0337/001	035719018	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
PENNSAID 16 mg/ml soluzione cutanea	EL/H/0337/001	035719032	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
PENNSAID 16 mg/ml soluzione cutanea	EL/H/0337/001	035719020	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
PENNSAID 16 mg/ml δερματικό διάλυμα	EL/H/0337/001/MR	39656/08-04-2019	VIANEX S.A.	GR
QPAIN 1,5% w/w δερματικό διάλυμα	not available	62581/19-06-2023	IASIS PHARMA	GR
ratioDolor Diclofenac Schmerzgel 1 % Gel	DE/H/5493/001	140246	TEVA B.V	AT
ratioDolor Diclofenac Schmerzgel 2 % Gel	DE/H/5493/002	140247	TEVA B.V	AT
SAILIB 1,5 % w/w δερματικό διάλυμα	not available	65259/24-06-2020	LIBYTEC PHARMACEUTICAL S.A.	GR
Solacutan 3% Gel	AT/H/0519/001	136996	DERMAPHARM GMBH	AT
Solacutan 3% gel	AT/H/0710/001/DC	045242029	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001/DC	045242043	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001/DC	045242056	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001/DC	045242068	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001/DC	045242070	MIBE PHARMA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solacutan 3% gel	AT/H/0710/001/DC	045242031	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001/DC	045242017	MIBE PHARMA ITALIA S.R.L.	IT
Solacutan 3% gel	AT/H/0710/001	PL 49452/0001	MIBE PHARMA UK LTD	XI
Solacutan 30 mg/g Gel	AT/H/0519/001	90764.00.00	DERMAPHARM AG	DE
Solacutan 30 mg/g gel	AT/H/0710/001	81761	MIBE PHARMA ESPAÑA S.L.U.	ES
Solacutan, 30 mg/g, żel	AT/H/0519/001	23371	SUN-FARM SP. Z.O.O.	PL
Solaraze 3 % geeli	DE/H/5854/001	17888	ALMIRALL, S.A.	FI
Solaraze 3 % gel diklofenaknatrium	DE/H/5854/001	17888	ALMIRALL, S.A.	FI
SOLARAZE 3 %, gel	DE/H/5854/001	34009 349 080 3 3	ALMIRALL, S.A.	FR
SOLARAZE 3 %, gel	DE/H/5854/001	34009 349 082 6 2	ALMIRALL, S.A.	FR
SOLARAZE 3 %, gel	DE/H/5854/001	34009 387 108 9 2	ALMIRALL, S.A.	FR
SOLARAZE 3 %, gel	DE/H/5854/001	34009 387 110 3 5	ALMIRALL, S.A.	FR
SOLARAZE 3 %, gel	DE/H/5854/001	34009 387 112 6 4	ALMIRALL, S.A.	FR
Solaraze 3 %, gel	DE/H/5854/001	02-1566	ALMIRALL, S.A.	NO
Solaraze 3% Gel	DE/H/5854/001	1-24961	ALMIRALL, S.A.	AT
Solaraze 3% Gel	DE/H/5854/001	42752.00.00	ALMIRALL, S.A.	DE
Solaraze 3% gel	DE/H/5854/001	034129027	ALMIRALL, S.A.	IT
Solaraze 3% gel	DE/H/5854/001	034129054	ALMIRALL, S.A.	IT
Solaraze 3% gel	DE/H/5854/001	034129041	ALMIRALL, S.A.	IT
Solaraze 3% gel	DE/H/5854/001	034129015	ALMIRALL, S.A.	IT
Solaraze 3% gel	DE/H/5854/001	034129039	ALMIRALL, S.A.	IT
Solaraze 3% Gel	DE/H/5854/001	2003070004	ALMIRALL, S.A.	LU
Solaraze 3%, gel	DE/H/5854/001	14192	ALMIRALL, S.A.	SE
Solaraze 30 mg/g Gel	DE/H/5854/001	73.714	ALMIRALL, S.A.	ES
Solaraze gel a 3%	DE/H/5854/001	4392486	ALMIRALL, S.A.	PT
Solaraze gel a 3%	DE/H/5854/001	5184254	ALMIRALL, S.A.	PT
Solaraze gel a 3%	DE/H/5854/001	5184262	ALMIRALL, S.A.	PT
Solaraze gel a 3%	DE/H/5854/001	5184247	ALMIRALL, S.A.	PT
Solaraze gel a 3%	DE/H/5854/001	4392387	ALMIRALL, S.A.	PT
Solaraze, gel	DE/H/5854/001	34524	ALMIRALL, S.A.	DK
Solaraze3%, gel	DE/H/5854/001	PA 968/004/001	ALMIRALL, S.A.	IE
Solaraze3%, gel	DE/H/5854/001	PL 16973/0012	ALMIRALL, S.A.	XI
Solaraze™ 3%, hlaup	DE/H/5854/001	IS/1/03/009/01	ALMIRALL, S.A.	IS
TOFENAL 4% gel	not available	040957019	OP PHARMA S.R.L.	IT
TRAULEN 4% gel	not available	033420.050	OP PHARMA S.R.L.	IT

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Trekpleister Diclofenac 2,32% extra sterk, gel	not available	RVG 130456	MAE HOLDING BV	NL
VISUNAC 1 mg/ml collirio, soluzione	not available	040518019	VISUFARMA SPA	IT
VISUNAC 1 mg/ml collirio, soluzione	not available	040518021	VISUFARMA SPA	IT
Volini 50 mg/g gel	not available	10746/2018/01	TERAPIA S.A.	RO
Volini 50 mg/g gel	not available	10746/2018/02	TERAPIA S.A.	RO
Voltabak, øjendråber, opløsning	FR/H/0296/001	47537	LABORATOIRES THEA	DK
Voltadol 11,6 mg/g Gel	not available	71.620	HALEON SPAIN, S.A.	ES
Voltadol 140 mg apósito adhesivo medicamentoso	DE/H/6101/001	85300	HALEON SPAIN, S.A.	ES
VOLTADOL 140mg, Έμπλαστρο	DE/H/2679/001	300500102	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
VOLTADOL 140mg, Έμπλαστρο	DE/H/2679/001	300500101	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
VOLTADOL 140mg, Έμπλαστρο	DE/H/2679/001	300500103	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltadol Forte 23,2 mg/g gel.	not available	78407	HALEON SPAIN, S.A.	ES
Voltadol Forte Schmerzgel	not available	1-31531	GSK-GEBRO CONSUMER HEALTHCARE GMBH	AT
Voltadol Schmerzgel	not available	1-23820	GSK-GEBRO CONSUMER HEALTHCARE GMBH	AT
Voltadol Schmerzpflaster 140 mg wirkstoffhaltiges Pflaster	DE/H/4645/001	138228	GSK-GEBRO CONSUMER HEALTHCARE GMBH	AT
Voltadol Unidie 140 mg cerotto medicato	DE/H/6101/001	048717019	HALEON ITALY S.R.L.	IT
Voltadol Unidie 140 mg cerotto medicato	DE/H/6101/001	048717021	HALEON ITALY 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
Voltadol Unidie 140 mg cerotto medicato	DE/H/6101/001	048717033	HALEON ITALY S.R.L.	IT
Voltadol Unidie 140 mg cerotto medicato	DE/H/6101/001	048717045	HALEON ITALY S.R.L.	IT
Voltaren Emulgel 10 mg/g gel	not available	29/127/88-C	HALEON CZECH REPUBLIC S.R.O.	CZ
Voltaren ® Emulgel - Gel	not available	1-18355	NOVARTIS PHARMA GMBH	AT
Voltarén 1 mg/ml colirio en solución	not available	58.308	LABORATOIRES THEA	ES
Voltarén 1 mg/ml colirio en solución en envases unidosis	not available	62.326	LABORATOIRES THEA	ES
Voltaren 1 mg/ml Colírio, solução	not available	2223485	LABORATOIRES THEA	PT
Voltaren 11,6 mg/g gel.	not available	20227	HALEON DENMARK APS	SE
Voltaren 11,6 mg/g hlaup	not available	IS/1/04/156/01	HALEON DENMARK APS	IS
Voltaren 140 mg lääkelaastari	DE/H/2679/001	30251	HALEON DENMARK APS	FI
Voltaren 140 mg léčivá náplast	DE/H/2679/001	29/257/13-C	HALEON CZECH REPUBLIC S.R.O.	CZ
Voltaren 140 mg liečivá náplasť	DE/H/2679/001	29/0131/13-S	HALEON CZECH REPUBLIC S.R.O.	SK
Voltaren 140 mg medicinskt plåster	DE/H/2679/001	30251	HALEON DENMARK APS	FI
Voltaren 140 mg medicinskt plåster	DE/H/2679/001	47014	HALEON DENMARK APS	SE
Voltaren 1x denně 140 mg léčivá náplast	DE/H/6101/001	29/121/19-C	HALEON CZECH REPUBLIC S.R.O.	CZ
Voltaren 1x denne 140 mg liečivá náplasť	DE/H/6101/001	29/0122/20-S	HALEON CZECH REPUBLIC S.R.O.	SK
Voltaren 23,2 mg/g gel	not available	44669	HALEON DENMARK APS	SE
Voltaren 24 Stunden Schmerzpflaster 140 mg wirkstoffhaltiges Pflaster	DE/H/6101/001	2203408.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren ActiGo 140 mg gyógyszeres tapasz	DE/H/2679/001	OGYI-T-22466/02	GLAXOSMITHKLINE-CONSUMER KFT.	HU
VOLTAREN ACTIGO 140	DE/H/2679/001	OGYI-T-22466/03	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 GYÓGYSZERES TAPASZ			CONSUMER KFT.	
VOLTAREN ACTIGO 140 MG GYÓGYSZERES TAPASZ	DE/H/2679/001	OGYI-T-22466/01	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Aktigo 140 mg ārstnieciskais plāksteris	DE/H/2679/001	13-0111	GLAXOSMITHKLINE DUNGARVAN LIMITED	LV
Voltaren Aktigo 140 mg vaistinis pleistras	DE/H/2679/001	LT/1/13/3467/002	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Aktigo 140 mg vaistinis pleistras	DE/H/2679/001	LT/1/13/3467/001	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Aktigo 140 mg vaistinis pleistras	DE/H/2679/001	LT/1/13/3467/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Aktigo, 140 mg ravimplaaster	DE/H/2679/001	814113	GLAXOSMITHKLINE DUNGARVAN LIMITED	EE
Voltaren Colírio Unidoses 0,3 mg/0,3 ml Colírio, solução	not available	2911287	LABORATOIRES THEA	PT
Voltaren Colírio Unidoses 0,3 mg/0,3 ml Colírio, solução	not available	2911386	LABORATOIRES THEA	PT
Voltaren Colírio Unidoses 0,3 mg/0,3 ml Colírio, solução	not available	2911485	LABORATOIRES THEA	PT
Voltaren Emulgel	not available	MA1177/00401	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	MT
VOLTAREN EMULGEL 1% gel	not available	BE274574	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE135317	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE274574	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE294847	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE294847	HALEON BELGIUM	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTAREN EMULGEL 1% gel	not available	BE135317	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE553600	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% gel	not available	BE553600	HALEON BELGIUM	BE
VOLTAREN Emulgel 1% gel	not available	034548091	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548103	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548115	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548038	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548040	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548089	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548139	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548228	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548216	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548204	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548192	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548180	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 1% gel	not available	034548178	NOVARTIS FARMA S.P.A.	IT
VOLTAREN EMULGEL 1% gel	not available	2008049768	HALEON BELGIUM	LU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/69	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/64	GLAXOSMITHKLINE-CONSUMER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/70	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/63	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/66	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/65	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/71	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/67	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/68	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/19	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/20	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/18	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/13	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/10	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/17	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/16	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/14	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/12	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/15	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/11	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/32	GLAXOSMITHKLINE-CONSUMER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/45	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/44	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/46	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 1% gél	not available	OGYI-T-5572/47	GLAXOSMITHKLINE-CONSUMER KFT.	HU
VOLTAREN EMULGEL 1% Gel Diclofenac-Natrium	not available	BE294847	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% Gel Diclofenac-Natrium	not available	BE135317	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% Gel Diclofenac-Natrium	not available	BE274574	HALEON BELGIUM	BE
VOLTAREN EMULGEL 1% Gel Diclofenac-Natrium	not available	BE553600	HALEON BELGIUM	BE
Voltaren Emulgel 1,16 % Gel	not available	520.00.03	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
VOLTAREN EMULGEL 1.16%, gel	not available	RVG 31377	HALEON NETHERLANDS B.V.	NL
Voltaren Emulgel 10 mg/g gel	not available	HR-H-727399086	GLAXOSMITHKLINE DUNGARVAN LIMITED	HR
Voltaren Emulgel 10 mg/g gel	not available	5855598	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 10 mg/g gel	not available	5197868	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 10 mg/g gel	not available	9657502	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 10 mg/g gel	not available	9657510	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 10 mg/g gel	not available	5219316	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 10	not available	5035654	HALEON PORTUGAL, LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/g gel				
Voltaren Emulgel 10 mg/g gel	not available	5712740	HALEON PORTUGAL, LDA	PT
Voltaren Emulgel 11,6 mg/g (1,16%) geel	not available	061794	GLAXOSMITHKLINE DUNGARVAN LIMITED	EE
Voltaren Emulgel 11,6 mg/g geeli	not available	11748	HALEON DENMARK APS	FI
Voltaren Emulgel 11,6 mg/g gel	not available	58.402	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren Emulgel 11,6 mg/g gel	not available	11748	HALEON DENMARK APS	FI
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/07	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/05	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/06	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/09	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/08	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/10	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/11	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/12	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/13	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/14	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/15	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/16	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/17	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/18	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/19	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
VOLTAREN EMULGEL 11,6 mg/g gel	not available	13704/2021/20	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/002	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/005	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/004	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/001	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/009	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/010	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/011	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel	not available	H/92/01655/012	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gel, tlačni vsebnik	not available	H/92/01655/006	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/004	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/003	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/005	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/012	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/001	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/002	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/006	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gelis	not available	LT/1/94/0943/015	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 11,6 mg/g gels	not available	94-0179	GLAXOSMITHKLINE DUNGARVAN LIMITED	LV
VOLTAREN Emulgel 2% gel	not available	034548230	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548127	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548053	NOVARTIS FARMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTAREN Emulgel 2% gel	not available	034548065	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548077	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548141	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548154	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548166	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548230	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548127	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548053	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548065	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548077	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548141	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548154	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548166	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548230	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548127	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548053	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548065	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548077	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548141	NOVARTIS FARMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTAREN Emulgel 2% gel	not available	034548154	NOVARTIS FARMA S.P.A.	IT
VOLTAREN Emulgel 2% gel	not available	034548166	NOVARTIS FARMA S.P.A.	IT
Voltaren Emulgel 20 mg/g gel	not available	HR-H-828714474	GLAXOSMITHKLINE DUNGARVAN LIMITED	HR
Voltaren Emulgel 23,2 mg/g (2,32%) geel	not available	810513	GLAXOSMITHKLINE DUNGARVAN LIMITED	EE
Voltaren Emulgel 23,2 mg/g gel	not available	H/92/01655/008	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 23,2 mg/g gel	not available	H/92/01655/007	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 23,2 mg/g gel	not available	H/92/01655/013	GLAXOSMITHKLINE DUNGARVAN LIMITED	SI
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/010	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/011	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/013	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/008	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/007	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/009	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gelis	not available	LT/1/94/0943/014	GLAXOSMITHKLINE DUNGARVAN LIMITED	LT
Voltaren Emulgel 23,2 mg/g gels	not available	12-0188	GLAXOSMITHKLINE DUNGARVAN LIMITED	LV
Voltaren Emulgel Extra Sterk 2,32%, gel	not available	RVG 117734	HALEON NETHERLANDS B.V.	NL
Voltaren Emulgel Forte 2% Gel	not available	BE440422	HALEON BELGIUM	BE
Voltaren Emulgel Forte 2% gel	not available	BE440422	HALEON BELGIUM	BE
Voltaren Emulgel Forte 2% gel	not available	BE440422	HALEON BELGIUM	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Emulgel Forte 2% gel	not available	BE562231	HALEON BELGIUM	BE
Voltaren Emulgel Forte 2% gel	not available	BE562231	HALEON BELGIUM	BE
Voltaren Emulgel Forte 2% gel	not available	2013120670	HALEON BELGIUM	LU
Voltaren Emulgel Forte 2% gel Diclofenac-Natrium	not available	BE562231	HALEON BELGIUM	BE
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/55	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/60	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/51	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/59	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/53	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/73	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/52	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/72	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/49	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/58	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/56	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/48	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/62	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/54	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte	not available	OGYI-T-5572/57	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
20 mg/g gél			CONSUMER KFT.	
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/50	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/74	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/61	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/75	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/37	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/34	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/35	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/42	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/33	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/39	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/38	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/36	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/41	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/40	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel Forte 20 mg/g gél	not available	OGYI-T-5572/43	GLAXOSMITHKLINE-CONSUMER KFT.	HU
Voltaren Emulgel 11,6 mg/g gél	not available	29/0127/88-CS	HALEON CZECH REPUBLIC S.R.O.	SK
Voltaren Emulgelex 23,2 mg/g gel	not available	5864863	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5490602	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2	not available	5625025	HALEON PORTUGAL, LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/g gel				
Voltaren Emulgelex 23,2 mg/g gel	not available	5374848	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5380571	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5380563	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5781703	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5781711	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5781729	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5781737	HALEON PORTUGAL, LDA	PT
Voltaren Emulgelex 23,2 mg/g gel	not available	5781745	HALEON PORTUGAL, LDA	PT
Voltaren Forte 2.32% gel	not available	MA1177/00402	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	MT
Voltaren Forte 2,32 % gél	not available	29/0481/11-S	HALEON CZECH REPUBLIC S.R.O.	SK
Voltaren Forte 20 mg/g gel	not available	29/070/13-C	HALEON CZECH REPUBLIC S.R.O.	CZ
Voltaren Forte 23,2 mg/g geeli	not available	28882	HALEON DENMARK APS	FI
Voltaren Forte 23,2 mg/g gel	not available	28882	HALEON DENMARK APS	FI
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/28	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/25	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/27	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/30	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/26	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/29	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/02	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/04	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/01	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/03	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/08	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/07	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/09	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/05	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/06	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/10	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/11	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/12	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/13	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/14	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/15	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/16	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/17	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/18	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/19	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/20	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/21	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/22	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/23	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Forte 23,2 mg/g gel	not available	13380/2020/24	GLAXOSMITHKLINE CONSUMER HEALTHCARE S.R.L.	RO
Voltaren Forte 23,2 mg/g hlaup	not available	IS/1/20/025/01	HALEON DENMARK APS	IS
Voltaren Forte, gel	not available	60772	HALEON DENMARK APS	DK
Voltaren MAX, 23,2 mg/g, żel	not available	20030	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
VOLTAREN OFTA 0,1% collirio, soluzione	not available	027917018	LABORATOIRES THEA	IT
VOLTAREN OFTA 0,1% collirio, soluzione	not available	027917020	LABORATOIRES THEA	IT
VOLTAREN OFTA 0,1% collirio, soluzione	not available	027917032	LABORATOIRES THEA	IT
VOLTAREN OFTABAK 1 mg/ml collirio, soluzione	FR/H/0296/001	037696010	LABORATOIRES THEA	IT
Voltaren Ophtha – 1 mg/ml Augentropfen	not available	1–19877	LABORATOIRES THEA	AT
Voltaren Ophtha – 1 mg/ml Einmalaugentropfen	not available	1–23917	LABORATOIRES THEA	AT
Voltaren ophtha 1 mg/ml Augentropfen	not available	11202.00.00	LABORATOIRES THEA	DE
Voltaren Ophtha 1 mg/ml ögondroppar, lösning	not available	10888	LABORATOIRES THEA	FI
Voltaren Ophtha 1 mg/ml ögondroppar, lösning	not available	11451	LABORATOIRES THEA	SE
Voltaren Ophtha 1 mg/ml ögondroppar, lösning, endosbehållare	not available	11439	LABORATOIRES THEA	FI
Voltaren Ophtha 1 mg/ml ögondroppar, lösning, endosbehållare	not available	12085	LABORATOIRES THEA	SE
Voltaren Ophtha 1 mg/ml øyedråper, oppløsning i endosebeholdere.	not available	11-8492	LABORATOIRES THEA	NO
Voltaren Ophtha 1 mg/ml	not available	7379	LABORATOIRES THEA	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
øyedråper, oppløsning.				
Voltaren Ophtha 1 mg/ml silmätipat, liuos	not available	10888	LABORATOIRES THEA	FI
Voltaren Ophtha 1 mg/ml silmätipat, liuos kerta-annospakkauksessa	not available	11439	LABORATOIRES THEA	FI
Voltaren Ophtha Abak 1 mg/ml Augentropfen	FR/H/0296/001	1-30552	LABORATOIRES THEA	AT
Voltaren Ophtha Abak 1 mg/ml, øyedråper, oppløsning	FR/H/0296/001	10-7850	LABORATOIRES THEA	NO
Voltaren Ophtha CD 1 mg/ml oldatos szemcsepp	not available	OGYI-T-5572/07	LABORATOIRES THEA	HU
Voltaren Patch 140 mg emplâtre médicamenteux	DE/H/2679/001	BE439652	HALEON BELGIUM	BE
Voltaren Patch 140 mg pleister	DE/H/2679/001	BE439652	HALEON BELGIUM	BE
Voltaren Patch 140 mg wirkstoffhaltige Pflaster Für Jugendliche ab 16 Jahren und Erwachsene	DE/H/2679/001	BE439652	HALEON BELGIUM	BE
Voltaren Plast 140 mg emplastro medicamentoso	DE/H/2679/001	5569728	HALEON PORTUGAL, LDA	PT
Voltaren Plast 140 mg emplastro medicamentoso	DE/H/2679/001	5569744	HALEON PORTUGAL, LDA	PT
Voltaren Plast 140 mg emplastro medicamentoso	DE/H/2679/001	5569736	HALEON PORTUGAL, LDA	PT
Voltaren Schmerzgel 11,6 mg/g Gel	not available	9165.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Schmerzgel 11,6 mg/g Gel	not available	87009.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Schmerzgel	not available	82680.00.00	GLAXOSMITHKLINE	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
forte 23,2 mg/g Gel			CONSUMER HEALTHCARE GMBH & CO. KG	
Voltaren Schmerzgel forte 23,2 mg/g Gel	not available	82421.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren Schmerzpflaster 140 mg wirkstoffhaltiges Pflaster	DE/H/2679/001	80422.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Voltaren SPORT, 11,6 mg/g, żel	not available	R/1735	HALEON POLAND SP. Z O.O.	PL
Voltaren, gel	not available	35874	HALEON DENMARK APS	DK
Voltaren, gel i trykbeholder	not available	46325	HALEON DENMARK APS	DK
Voltaren® Emulgel®	not available	18989	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	CY
Voltaren® ophtha sine 1 mg/ml Augentropfen	not available	26204.00.00	LABORATOIRES THEA	DE
Voltaren® ophtha sine 1 mg/ml Augentropfen	not available	1536/11041074	LABORATOIRES THEA	LU
Voltaren® Ophtha Οφθαλμικές σταγόνες 0,1%	not available	13462	LABORATOIRES THEA	CY
VOLTARENACTIGO 1 %, gel en flacon pressurisé	not available	34009 359 314 7 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 %, gel en flacon pressurisé	not available	34009 280 079 1 9	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 340 765 3 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 372 302 9 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 372 303 5 3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-	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
			MALMAISON	
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 375 463 3 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 363 540 8 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 375 465 6 0	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 280 077 9 7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 280 078 5 8	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 301 289 7 8	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 1 POUR CENT, gel	not available	34009 301 480 5 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENACTIGO 2 % INTENSE, gel	not available	34009 274 075 8 1	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel	not available	34009 328 869 7 5	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel	not available	34009 372 304 1 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel	not available	34009 372 305 8 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel	not available	34009 328 870 5 7	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel en flacon	not available	34009 358 947 6 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-	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
pressurisé			MALMAISON	
VOLTARENE EMULGEL 1 %, gel en flacon pressurisé	not available	34009 358 945 3 3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENE EMULGEL 1 %, gel en flacon pressurisé	not available	34009 358 951 3 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENOPHTA 0,1 POUR CENT (0,3 mg/0,3 ml), collyre en solution en récipient unidose	not available	338 685-6	LABORATOIRES THEA	FR
VOLTARENOPHTA 0,1 POUR CENT (0,3 mg/0,3 ml), collyre en solution en récipient unidose	not available	338 686-2	LABORATOIRES THEA	FR
VOLTARENOPHTA 0,1 POUR CENT (0,3 mg/0,3 ml), collyre en solution en récipient unidose	not available	338 687-9	LABORATOIRES THEA	FR
VOLTARENOPHTABAK 1 mg/ml, collyre en solution	FR/H/0296/001	34009 366 822 4 5	LABORATOIRES THEA	FR
VOLTARENPLAST 1%, emplâtre médicamenteux	DE/H/1480/001	34009 394 222 8 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENPLAST 1%, emplâtre médicamenteux	DE/H/1480/001	394 225-7 OU 34009 394 225 7 2	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENPLAST 1%, emplâtre médicamenteux	DE/H/1480/001	394 223-4 OU 34009 394 223 4 3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENPLAST 1%, emplâtre médicamenteux	DE/H/1480/001	394 224-0 OR 34009 394 224 0 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENSPE 1 %, gel	not available	34009 372 300 6 3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENSPE 1 %, gel	not available	34009 340 761 8 3	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-	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
			MALMAISON	
VOLTARENSPE 1 %, gel	not available	34009 340 762 4 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
VOLTARENSPE 1 %, gel	not available	34009 372 301 2 4	GLAXOSMITHKLINE SANTÉ GRAND PUBLIC, RUEIL-MALMAISON	FR
Voltarol 1% w/w Gel	not available	PA0678/140/001	HALEON IRELAND LIMITED	IE
Voltarol 1.16% Emulgel, gel	not available	PL 44673/0157	HALEON UK TRADING LIMITED	XI
Voltarol 11,6 mg/g (1,16%) gel.	not available	03-2300	HALEON DENMARK APS	NO
Voltarol 12 Hour Emulgel P 2.32% Gel	not available	PL 44673/0154	HALEON UK TRADING LIMITED	XI
Voltarol 12 Hour Joint Pain Relief 2.32% Gel	not available	PL 44673/0154	HALEON UK TRADING LIMITED	XI
Voltarol 140 mg Medicated Plaster	not available	PL 15545/0012	GLAXOSMITHKLINE DUNGARVAN LIMITED	XI
Voltarol 140 mg medisinert plaster	DE/H/2679/001	11-8696	HALEON DENMARK APS	NO
Voltarol Back & Muscle Pain Relief 1.16% Gel	not available	PL 44673/0156	HALEON UK TRADING LIMITED	XI
Voltarol Emulgel	not available	321470104	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470101	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470102	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470105	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	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
Voltarol Emulgel	not available	321470106	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470103	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470107	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470108	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel	not available	321470109	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Emulgel 1% w/w Gel	not available	PA0678/140/002	HALEON IRELAND LIMITED	IE
Voltarol Emulgel Extra Strength 2% w/w Gel	not available	PA 678/140/003	HALEON IRELAND LIMITED	IE
Voltarol Emulgel P	not available	PL 44673/0155	HALEON UK TRADING LIMITED	XI
Voltarol Extra Strength Emulgel 2.32% Gel	not available	PL 44673/0160	HALEON UK TRADING LIMITED	XI
Voltarol Forte 2% w/w γέλη	not available	321470206	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470208	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470207	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER	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
			SOCIETE ANONYME	
Voltarol Forte 2% w/w γέλη	not available	321470201	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470202	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470203	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470205	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 2% w/w γέλη	not available	321470204	GLAXOSMITHKLINE CONSUMER HEALTHCARE HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Forte 23,2 mg/g (2,32 %) gel.	not available	10-7734	HALEON DENMARK APS	NO
Voltarol Joint & Back Pain Relief 2.32% Gel	not available	PL 44673/0160	HALEON UK TRADING LIMITED	XI
Voltarol Joint Pain Relief 2.32% Gel	not available	PL 44673/0160	HALEON UK TRADING LIMITED	XI
Voltarol Max Strength Pain Relief 2.32% Gel	not available	PL 44673/0154	HALEON UK TRADING LIMITED	XI
Voltarol Ophtha	not available	PL 20162/0018	LABORATOIRES THEA	XI
Voltarol Ophtha Multidose 0.1% Eye Drops	not available	PL 20162/0017	LABORATOIRES THEA	XI
Voltarol Ophtha Multidose 1 mg/ml Eye Drops Solution.	not available	PA 1107/009/002	LABORATOIRES THEA	IE
Voltarol Osteoarthritis Joint Pain Relief 1.16% Gel	not available	PL 44673/0155	HALEON UK TRADING LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltarol Pain-eze Emulgel	not available	PL 44673/0156	HALEON UK TRADING LIMITED	XI
Voltarol, medicinsk plaster	DE/H/2679/001	49708	HALEON DENMARK APS	DK
Vurdon 16 mg/ml δερματικό διάλυμα	not available	64540/09-07-2021	HELP ABEE	GR
Vurdon 16 mg/ml δερματικό διάλυμα	not available	64540/09-07-2021	HELP ABEE	GR
Zaflex , 10 mg/g, gel	not available	5651542	CUIDAFARMA, LDA.	PT
Zaflex , 10 mg/g, gel	not available	5651526	CUIDAFARMA, LDA.	PT
Zaflex , 10 mg/g, gel	not available	5651534	CUIDAFARMA, LDA.	PT
ZARANNY 180 mg cerotto medicato	IT/H/0562/001	045951011	MIAT	IT
ZARANNY 180 mg cerotto medicato	IT/H/0562/001	045951023	MIAT	IT
Zaranny 140 mg emplastro medicamentoso.	IT/H/0562/001	5770623	MIAT	PT
Zaranny 140 mg emplastro medicamentoso.	IT/H/0562/001	5770631	MIAT	PT
ZEROFLOG 0,011 g/15 ml collutorio, 12 bustine	not available	034373 023	VALEAS S.P.A.	IT
ZEROFLOG 0,074 g/100 ml collutorio, 1 flacone	not available	034373 011	VALEAS S.P.A.	IT
ZEROFLOG 0,022 g /15 ml soluzione, spray per mucosa orale	not available	034373 035	VALEAS S.P.A.	IT
ZILIPROL 1,5% w/w de?µat??? d????µa	not available	25107/18/11-01-2019	MEDICAIR BIOSCIENCE LABORATORIES ΑΝΩΝΥΜΗ ΦΑΡΜΑΚΕΥΤΙΚΗ ΕΤΑΙΡΕΙΑ	GR
ВОЛТАРЕН ЕМУЛГЕЛ 1,16% Гел	not available	20011208	GLAXOSMITHKLINE DUNGARVAN LIMITED	BG
ВОЛТАРЕН ФОРТЕ 2,32% ГЕЛ	not available	20110722	GLAXOSMITHKLINE DUNGARVAN LIMITED	BG
ГЛИМБАКС 0,074 % промивка за уста	not available	20050526	ANGELINI PHARMA BULGARIA EOOD	BG
ГЛИМБАКС 0,074 %	not available	20050526	ANGELINI PHARMA	BG

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
промивка за уста			BULGARIA EOOD	
ГЛИМБАКС 0,074 % промивка за уста	not available	20050526	ANGELINI PHARMA BULGARIA EOOD	BG
ГЛИМБАКС 0,148 % спрей за устна лигавица, разтвор	not available	20130096	ANGELINI PHARMA BULGARIA EOOD	BG
ГЛИМБАКС 0,148 % спрей за устна лигавица, разтвор	not available	20130096	ANGELINI PHARMA BULGARIA EOOD	BG
Диклак 12 часа 23,2 mg/g гел	HU/H/0745/001	20230057	SANDOZ PHARMACEUTICALS D.D.	BG
Диклоабак 1 mg/ml капки за очи, разтвор	FR/H/0296/001	20120184	LABORATOIRES THEA	BG
Мобилат Емулгел 2,32% гел	DE/H/7329/001	20230109	STADA ARZNEIMITTEL AG	BG