26 October 2017
EMA/663700/2017
Human Medicines Evaluation Division

# List of nationally authorised medicinal products

Active substance: cromoglicic acid

Procedure no.: PSUSA/00000883/201702

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
**Telephone** +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
**Send a question via our website** www.ema.europa.eu/contact

| 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 |
|---|---|---|---|---|
| Alleofta 20 mg/ml kapi za oko, otopina u jednodoznom spremniku | UK/H/5451/001 | HR-H-586030470 | SANOFI-AVENTIS CROATIA D.O.O. | HR |
| Alleofta 20 mg/ml kapi za oko, otopina u jednodoznom spremniku | UK/H/5451/001 | HR-H-586030470 | SANOFI-AVENTIS CROATIA D.O.O. | HR |
| Alleophta 20 mg/ml, očná roztoková instilácia v jednodávkovom obale | UK/H/5451/001 | 64/0414/16-S | SANOFI-AVENTIS SLOVAKIA SRO | SK |
| Alleophta 20 mg/ml, očná roztoková instilácia v jednodávkovom obale | UK/H/5451/001 | 64/0414/16-S | SANOFI-AVENTIS SLOVAKIA SRO | SK |
| Alleophta unidosis 20 mg/ml, collyre en solution en récipient unidose | UK/H/5451/01/DC | BE467022 | SANOFI BELGIUM | BE |
| Alleophta unidosis 20 mg/ml, collyre en solution en récipient unidose | UK/H/5451/01/DC | BE467022 | SANOFI BELGIUM | BE |
| Alleophta unidosis 20 mg/ml, collyre en solution en récipient unidose | UK/H/5451/01/DC | 0780335 | SANOFI BELGIUM | LU |
| Alleophta unidosis 20 mg/ml, collyre en solution en récipient unidose | UK/H/5451/01/DC | 0780321 | SANOFI BELGIUM | LU |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Alleophta unidosis 20 mg/ml, oogdruppels, oplossing in verpakking voor éénmalig gebruik | UK/H/5451/01/DC | BE467022 | SANOFI BELGIUM | BE |
| Alleophta unidosis 20 mg/ml, oogdruppels, oplossing in verpakking voor éénmalig gebruik | UK/H/5451/01/DC | BE467022 | SANOFI BELGIUM | BE |
| Alleopti 20 mg/ ml oldatos szemcsepp | not available | OGYI-T-22584/01 | SANOFI-AVENTIS ZRT | HU |
| ALLEOPTI 20 MG/ML OLDATOS SZEMCSEPP | not available | OGYI-T-22584/04 | SANOFI-AVENTIS ZRT | HU |
| Alleopti Komfort 20 mg/ ml oldatos szemcsepp egyadagos tartályban | not available | OGYI-T-22584/03 | SANOFI-AVENTIS ZRT | HU |
| Alleopti Komfort 20 mg/ ml oldatos szemcsepp egyadagos tartályban | not available | OGYI-T-22584/02 | SANOFI-AVENTIS ZRT | HU |
| Alleoptical, 20 mg/ml, krople do oczu, roztwór w pojemniku jednodawkowym | UK/H/5451/01/DC | 22656 | AVENTIS PHARMA LTD | PL |
| Alleoptical, 20 mg/ml, krople do oczu, roztwór w pojemniku jednodawkowym | UK/H/5451/01/DC | 22656 | AVENTIS PHARMA LTD | PL |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Allerg-Abak 20 mg/ml oogdruppels, oplossing | FR/H/0234/001 | RVG 29812 | LABORATOIRES THEA | NL |
| Allergo-COMOD eye drops | not available | MA063/00201 | URSAPHARM ARZNEIMITTEL GMBH | MT |
| Allergo-COMOD neusspray | not available | RVG 15592 | URSAPHARM BENELUX B.V. | NL |
| Allergo-COMOD ocní kapky, Oční kapky, roztok | not available | 64/020/01-C | URSAPHARM SPOL. S.R.O. | CZ |
| Allergo-COMOD oogdruppels | not available | RVG 15461 | URSAPHARM BENELUX B.V. | NL |
| Allergo-COMOD, 20 mg/ml, krople do oczu, roztwór | not available | 9762 | URSAPHARM POLAND SP. Z. O.O. | PL |
| ALLERGOCOMOD, collyre en solution | not available | 354 293-1 | LABORATOIRES URSAPHARM S.A.S. | FR |
| Allergo-COMOD® Augentropfen | not available | 1-24846 | URSAPHARM GES.M.B.H. | AT |
| Allergo-COMOD® Nasenspray | not available | 1-24841 | URSAPHARM GES.M.B.H. | AT |
| Allergocrom | not available | 64/0662/94-S | URSAPHARM ARZNEIMITTEL GMBH | SK |
| Allergocrom nosní sprej, Nosní sprej, roztok | not available | 69/945/97-C | URSAPHARM SPOL. S.R.O. | CZ |
| Allergocrom oční kapky, roztok | not available | 64/072/97-C | URSAPHARM SPOL. S.R.O. | CZ |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Allergocrom, 20 mg/ml, krople do oczu, roztwór | not available | 8980 | URSAPHARM POLAND SP. Z. O.O. | PL |
| Allergoval Kapseln | not available | 16619.00.03 | KOEHLER PHARMA GMBH | DE |
| Allgil 20 mg/ml ögondroppar, lösning | not available | 48236 | APOFRI AB | SE |
| Aspire Allergy Relief 2% w/v eye drops, Solution (eye drops) | not available | PL 35533/0033 | ASPIRE PHARMA LIMITED | UK |
| Aspire Hayfever Relief 2% w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| Boots Allergy Relief 2% w/v Eye Drops | not available | PL03468/0023 | BAUSCH & LOMB UK LTD. | UK |
| Boots Hayfever Relief 2% w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| Catacrom 2% w/v eye drops, solution | UK/H/1718/001 | PA 1464/1/1 | RAYNER PHARMACEUTICALS | IE |
| Catacrom 2% w/v eye drops, solution | UK/H/1718/001 | PL11412/0002 | MOORFIELDS EYE HOSPITAL NHS FOUNDATION TRUST | UK |
| COLIMUNE SACHETS 100 MG | not available | 355.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| COLIMUNE SACHETS 100 MG | not available | 355.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| COLIMUNE SACHETS 100 MG | not available | 355.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| COLIMUNE SACHETS 200 MG | not available | 355.01.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| COLIMUNE SACHETS 200 MG | not available | 355.01.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| COLIMUNE SACHETS 200 MG | not available | 355.01.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| CROMABAK 20 mg/ml colirio en solución | FR/H/0234/001 | 65867 | LABORATOIRES THEA | ES |
| CROMABAK 20 mg/ml collirio soluzione | FR/H/0234/001 | 036592020 | LABORATOIRES THEA | IT |
| Cromabak 20 mg/ml collyre en solution | not available | BE206333 | THEA PHARMA | BE |
| Cromabak 20 mg/ml collyre en solution | not available | BE206333 | THEA PHARMA | LU |
| Cromabak 20 mg/ml oogdruppels, oplossing | not available | BE206333 | THEA PHARMA | BE |
| Cromabak 20 mg/ml, colírio, solução | not available | 2612588 | LABORATOIRES THEA | PT |
| Cromabak 20 mg/ml, colírio, solução | not available | 2612687 | LABORATOIRES THEA | 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 |
|---|---|---|---|---|
| CROMABAK 20 mg/ml, collyre en solution | FR/H/0234/001 | 34009 340 530 6 1 | LABORATOIRES THEA | FR |
| Crominet 20 mg/ml ögondroppar, lösning | SE/H/1372/001 | 49988 | ORIFARM GENERICS A/S | SE |
| Crominet 20 mg/ml øyedråper, oppløsning | SE/H/1372/001 | 13-9749 | ORIFARM GENERICS A/S | NO |
| Crominet, øjendråber, opløsning | SE/H/1372/001 | 53098 | ORIFARM GENERICS A/S | DK |
| CROMOFREE 2 %, collyre en solution | not available | 34009 346 922 3 9 | LABORATOIRE CHAUVIN | FR |
| Cromoglicato de Sodio Brown 20 mglml colirio, solucao | UK/H/3500/001 | 5444906 | BROWN & BURK UK LIMITED | PT |
| Cromoglicato de Sodio Brown 20 mglml colirio, solucao | UK/H/3500/001 | 5444914 | BROWN & BURK UK LIMITED | PT |
| CROMOHEXAL | not available | 64/142/96-C | LABORATOIRES THEA | CZ |
| CROMOHEXAL 20 mg/1 ml Nosní sprej, roztok | not available | 24/141/96-C | HEXAL AG | CZ |
| Cromolux 2% Eye Drops | not available | PL 03468/0022 | BAUSCH & LOMB UK LTD. | UK |
| Cromolux 2% Hay fever Eye Drops | not available | PL03468/0023 | BAUSCH & LOMB UK LTD. | UK |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Crom-Ophtal® 1 ml Augentropfen enthält 20 mg Natriumcromoglicat (Ph.Eur.) | not available | 7974.00.00 | DR. WINZER PHARMA GMBH | DE |
| Crom-Ophtal® Nasenspray | not available | 7975.00.00 | DR. WINZER PHARMA GMBH | DE |
| Crom-Ophtal® sine 1 ml Augentropfen enthält 20 mg Natriumcromoglicat (Ph.Eur.) | not available | 24158.00.00 | DR. WINZER PHARMA GMBH | DE |
| CROMOPTIC 2 %, collyre en solution | not available | 345 917-6 | LABORATOIRE CHAUVIN | FR |
| CROMOPTIC 2 %, collyre en solution en récipient unidose | not available | 34009 353 566 4 2 | LABORATOIRE CHAUVIN | FR |
| CUSICROM FUERTE OFTÁLMICO 40 mg/ml colirio en solución | not available | 57327 | NOVARTIS FARMACÉUTICA S.A. | ES |
| EM Pharma Allergy Relief 2%w/v Eye Drops, Solution (eye drops) | not available | PL 35533/0033 | ASPIRE PHARMA LIMITED | UK |
| EM Pharma Hayfever Relief 2%w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| Fenolip 20 mg/ml solução para inalação por nebulização | not available | 8707505 | ANGELINI FARMACÊUTICA, 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 |
|---|---|---|---|---|
| Fenolip, 10 mg / 0,5 ml, colírio, solução | not available | 2910883 | ANGELINI FARMACÊUTICA, LDA | PT |
| Fenolip, 20 mg/ml, colírio, solução | not available | 8707406 | ANGELINI FARMACÊUTICA, LDA | PT |
| INTAL CFC-FREE INHALER | not available | PL 04425/0179 | AVENTIS PHARMA LTD | UK |
| INTAL INHALATIONLÖSUNG 1% | not available | 7835.00.00 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL INHALATIONLÖSUNG 1% | not available | 7835.00.00 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL INHALATIONLÖSUNG 1% | not available | 7835.00.00 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL INHALATIONLÖSUNG 1% | not available | 7835.00.00 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |

| Product name (in authorisation country) | MRP/DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| INTAL N AEROSOL | not available | 354.00.01 | SANOFI-AVENTIS DEUTSCHLAND GMBH | DE |
| Lecrolyn | not available | 15022 | SANTEN OY | DK |
| LECROLYN 20 mg/ml – acu pilieni, šķīdums | not available | 99-0312 | SANTEN OY | LV |
| LECROLYN 20 mg/ml akių lašai (tirpalas) | not available | LT/1/96/2408/001 | SANTEN OY | LT |
| Lecrolyn 20 mg/ml ögondroppar, lösning, endosbehållare | not available | 11459 | SANTEN OY | SE |
| LECROLYN 20 mg/ml silmatilgad, lahus | not available | 103795 | SANTEN OY | EE |
| LECROLYN 20 mg/ml - silmätipat, liuos | not available | 11026 | SANTEN OY | FI |
| LECROLYN 40 mg/ml – acu pilieni, šķīdums | not available | 99-0314 | SANTEN OY | LV |
| LECROLYN 40 mg/ml akių lašai (tirpalas) | not available | LT/1/96/2408/002 | SANTEN OY | LT |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| LECROLYN 40 mg/ml akių lašai, tirpalas | not available | LT/1/96/2408/004 | SANTEN OY | LT |
| LECROLYN 40 mg/ml akių lašai, tirpalas | not available | LT/1/96/2408/005 | SANTEN OY | LT |
| Lecrolyn 40 mg/ml ögondroppar, lösning | not available | 12128 | SANTEN OY | SE |
| Lecrolyn 40 mg/ml ögondroppar, lösning, endosbehållare | not available | 12129 | SANTEN OY | SE |
| Lecrolyn 40 mg/ml øyedråper, oppløsning i dråpeflaske | not available | 8306 | SANTEN OY | NO |
| Lecrolyn 40 mg/ml øyedråper, oppløsning i endosebeholdere | not available | 03-2101 | SANTEN OY | NO |
| LECROLYN 40 mg/ml silmatilgad, lahus | not available | 103995 | SANTEN OY | EE |
| LECROLYN 40 mg/ml - silmätipat, liuos | not available | 11503 | SANTEN OY | FI |
| Lecrolyn sine 40 mg/ml akių lašai (tirpalas) | SE/H/1402/001 | LT/1/15/3711/001 | SANTEN OY | LT |
| Lecrolyn sine 40 mg/ml akių lašai (tirpalas) | SE/H/1402/001 | LT/1/15/3711/002 | SANTEN OY | LT |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Lecrolyn sine 40 mg/ml akių lašai (tirpalas) | SE/H/1402/001 | LT/1/15/3711/003 | SANTEN OY | LT |
| Lecrolyn sine 40 mg/ml augndropar, lausn | SE/H/1402/01/DC | IS/1/15/035/01 | SANTEN OY | IS |
| Lecrolyn sine 40 mg/ml ögondroppar, lösning | SE/H/1402/001 | 32110 | SANTEN OY | FI |
| Lecrolyn sine 40 mg/ml ögondroppar, lösning | SE/H/1402/001 | 50887 | SANTEN OY | SE |
| Lecrolyn sine 40 mg/ml øyedråper, oppløsning | SE/H/1402/01/DC | 14-9996 | SANTEN OY | NO |
| Lecrolyn sine 40 mg/ml silmatilgad, lahus | SE/H/1402/001 | 894915 | SANTEN OY | EE |
| Lecrolyn sine 40 mg/ml silmätipat, liuos | SE/H/1402/01/DC | 32110 | SANTEN OY | FI |
| Lecrolyn sine, øjendråber, opløsning | SE/H/1402/001 | 53827 | SANTEN OY | DK |
| Lecrolyn, øjendråber, opløsning | not available | 14960 | SANTEN OY | DK |
| LECROLYN® 20 mg/ml ögondroppar, lösning | not available | 11026 | SANTEN OY | FI |
| LECROLYN® 20 mg/ml ögondroppar, lösning i endosbehållare | not available | 11027 | SANTEN OY | FI |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| LECROLYN® 20 mg/ml - silmätipat, liuos kerta-annospakkauksessa | not available | 11027 | SANTEN OY | FI |
| LECROLYN® 40 mg/ml ögondroppar, lösning | not available | 11503 | SANTEN OY | FI |
| LECROLYN® 40 mg/ml ögondroppar, lösning i endosbehållare | not available | 11504 | SANTEN OY | FI |
| LECROLYN® 40 mg/ml - silmätipat, liuos kerta-annospakkauksessa | not available | 11504 | SANTEN OY | FI |
| LOMUDAL | not available | MT 10015 | SANOFI-AVENTIS DENMARK A/S | DK |
| LOMUDAL | not available | 022319038 | SANOFI SPA | IT |
| LOMUDAL | not available | 8593 | SANOFI AB | SE |
| LOMUDAL 20 MG | not available | RVG 05719 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| LOMUDAL 20 MG | not available | RVG 05719 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| Lomudal 20 mg inhalationspulver, hård kapsel | not available | 8593 | SANOFI AB | SE |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Lomudal 20 mg inhalationspulver, hård kapsel | not available | 8593 | SANOFI AB | SE |
| LOMUDAL 20 mg, solution pour inhalation par nébuliseur | not available | BE110503 | SANOFI BELGIUM | BE |
| Lomudal 20 mg, verneveloplossing | not available | BE110503 | SANOFI BELGIUM | BE |
| Lomudal 20 mg/ml ögondroppar, lösning | not available | 9326 | SANOFI AB | SE |
| Lomudal 20 mg/ml ögondroppar, lösning | not available | 9326 | SANOFI AB | SE |
| Lomudal 20 mg/ml øyedråper, oppløsning | not available | 6300 | SANOFI-AVENTIS NORGE AS | NO |
| Lomudal 20 mg/ml øyedråper, oppløsning | not available | 6300 | SANOFI-AVENTIS NORGE AS | NO |
| Lomudal 20 mg/ml øyedråper, oppløsning | not available | 6300 | SANOFI-AVENTIS NORGE AS | NO |
| LOMUDAL 20 MG/ML SILMATIPAT, LIUOS | not available | 7589 | SANOFI OY | FI |
| LOMUDAL 20 MG/ML SILMATIPAT, LIUOS | not available | 7589 | SANOFI OY | FI |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Lomudal 20 mg/ml, augndropar, lausn | not available | 792354 | SANOFI-AVENTIS NORGE AS | IS |
| Lomudal 40 mg/ml ögondroppar, lösning, endosbehållare | not available | 11683 | SANOFI AB | SE |
| Lomudal 40 mg/ml ögondroppar, lösning, endosbehållare | not available | 11683 | SANOFI AB | SE |
| Lomudal 40 mg/ml øyedråper, oppløsning i endosebeholder | not available | 7842 | SANOFI-AVENTIS NORGE AS | NO |
| Lomudal 40 mg/ml øyedråper, oppløsning i endosebeholder | not available | 7842 | SANOFI-AVENTIS NORGE AS | NO |
| LOMUDAL 40 MG/ML SILMATIPAT, LIUOS KERTA-ANNOSPIPETISSA | not available | 11080 | SANOFI OY | FI |
| LOMUDAL 40 MG/ML SILMATIPAT, LIUOS KERTA-ANNOSPIPETISSA | not available | 11080 | SANOFI OY | FI |
| LOMUDAL 40mg/ml collirio, soluzione | not available | 022319065 | SANOFI SPA | IT |
| LOMUDAL 40mg/ml spray nasale, soluzione | not available | 022319077 | SANOFI SPA | 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 |
|---|---|---|---|---|
| LOMUDAL FORTE | not available | RVG 11817 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| LOMUDAL FREONITON 5 MG/ANNOS INHALAATIOSUMUTE, SUSPENSIO | not available | 13264 | SANOFI OY | FI |
| LOMUDAL G.I. | not available | 10468 | SANOFI AB | SE |
| LOMUDAL G.I. | not available | 10468 | SANOFI AB | SE |
| LOMUDAL NASAL | not available | 13636 | SANOFI-AVENTIS DENMARK A/S | DK |
| LOMUDAL NASAL | not available | 10971 | SANOFI AB | SE |
| LOMUDAL NASAL | not available | 10971 | SANOFI AB | SE |
| LOMUDAL NASAL 5,2 MG/ANNOS NENASUMUTE, LIUOS | not available | 11373 | SANOFI OY | FI |
| LOMUDAL NASAL 5,2 MG/ANNOS NENASUMUTE, LIUOS | not available | 11373 | SANOFI OY | FI |
| Lomudal Nasal 5,2 mg/dose nesespray, oppløsning | not available | 00/7496 | SANOFI-AVENTIS NORGE AS | NO |
| Lomudal Nasal 5,2 mg/dose nesespray, oppløsning | not available | 00/7496 | SANOFI-AVENTIS NORGE AS | NO |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| LOMUDAL NEBULISER SOL. FOR INH. (GLASS VIALS) | not available | 0067254 | SANOFI BELGIUM | LU |
| Lomudal, øjendråber, opløsning | not available | 09241 | SANOFI-AVENTIS DENMARK A/S | DK |
| Lomudal, øjendråber, opløsning | not available | 09241 | SANOFI-AVENTIS DENMARK A/S | DK |
| Lomudal, øjendråber, opløsning, enkeltdosisbeholder | not available | 38011 | SANOFI-AVENTIS DENMARK A/S | DK |
| Lomudal, øjendråber, opløsning, enkeltdosisbeholder | not available | 38011 | SANOFI-AVENTIS DENMARK A/S | DK |
| LOMUDAL, SOLUTION POUR NEBULISATION | not available | 324 118-7 | SANOFI-AVENTIS FRANCE | FR |
| LOMUDAL, SOLUTION POUR NEBULISATION | not available | 324 119-3 | SANOFI-AVENTIS FRANCE | FR |
| Lomudal® Gastrointestinum 100 mg kapselit | not available | 8299 | ITALCHIMICI S.P.A. | FI |
| LOMUSOL 2% | not available | 0006/09100548 | SANOFI BELGIUM | LU |
| Lomusol 2% neusdruppels, oplossing | not available | BE158706 | SANOFI BELGIUM | BE |
| Lomusol 2% solution pour instillation nasale | not available | BE158706 | SANOFI 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 |
|---|---|---|---|---|
| LOMUSOL 4 POUR CENT, solution pour pulvérisation nasale | not available | 337 537.3 | SANOFI-AVENTIS FRANCE | FR |
| LOMUSOL 4 POUR CENT, solution pour pulvérisation nasale | not available | 337 539.6 | SANOFI-AVENTIS FRANCE | FR |
| Lomusol 4% neusspray, oplossing | not available | BE172207 | SANOFI BELGIUM | BE |
| Lomusol 4% solution pour pulvérisation nasale | not available | BE172207 | SANOFI BELGIUM | BE |
| LOMUSOL 4% SPRAY | not available | 0216291 | SANOFI BELGIUM | LU |
| MULTICROM 2 %, collyre en solution | not available | 34009 340 532 9 0 | LABORATOIRES THEA | FR |
| MULTICROM UNIDOSES 2 POUR CENT, collyre en récipient unidose | not available | 348 847-9 | LABORATOIRES THEA | FR |
| MULTICROM UNIDOSES 2 POUR CENT, collyre en récipient unidose | not available | 349 775-1 | LABORATOIRES THEA | FR |
| MULTICROM UNIDOSES 2 POUR CENT, collyre en récipient unidose | not available | 357 483-6 | LABORATOIRES THEA | 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 |
|---|---|---|---|---|
| MULTICROM UNIDOSES 2 POUR CENT, collyre en récipient unidose | not available | 357 087-3 | LABORATOIRES THEA | FR |
| NALCROM | not available | RVG 10227 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| NALCROM | not available | RVG 10228 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| NALCROM | not available | 14/0305/89-S | ITALCHIMICI S.P.A. | SK |
| Nalcrom 100 mg capsule rigide | not available | 024861015 | ITALCHIMICI S.P.A. | IT |
| Nalcrom 100 mg, tvrdé tobolky | not available | 14/305/89-S/C | ITALCHIMICI S.P.A. | CZ |
| NALCROM 100MG CAPSULES | not available | PL 04425/0370 | AVENTIS PHARMA LTD | UK |
| Nalcrom 250 mg granulato per soluzione orale | not available | 024861039 | ITALCHIMICI S.P.A. | IT |
| NALCROM 250 mg granulato per soluzione orale | not available | 024861066 | ITALCHIMICI S.P.A. | IT |
| Nalcrom 500 mg granulato per soluzione orale | not available | 024861041 | ITALCHIMICI S.P.A. | IT |
| NALCROM 500 mg granulato per soluzione orale | not available | 024861078 | ITALCHIMICI 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 |
|---|---|---|---|---|
| NALCROM DRANK 100 MG | not available | RVG 12446 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| NALCROM DRANK 200 MG | not available | RVG 12447 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| Natriumcromoglicaat Apotex 20 mg/ml | not available | RVG 23703 | APOTEX EUROPE B.V. | NL |
| NATRIUMCROMOGLICAAT OOGDRUPPELS 20 G/ML | not available | RVG 57336 | BASIC PHARMA MANUFACTURING BV | NL |
| Natriumkromoglikat ABECE 20 mg/ml ögondroppar, lösning | not available | 55367 | EVOLAN PHARMA AB | SE |
| Numark Hayfever Relief 2% w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| OPHTACALM 2 %, collyre en solution en récipient unidose | not available | 34009 353 567 0 3 | LABORATOIRE CHAUVIN | FR |
| OPHTACALM 2 %, collyre en solution en récipient unidose | not available | 34009 353 568 7 1 | LABORATOIRE CHAUVIN | FR |
| OPHTACALM 2 %, collyre en solution en récipient unidose | not available | 34009 353 569 3 2 | LABORATOIRE CHAUVIN | FR |
| OPHTACALMFREE 2%, collyre en solution | not available | 34009 346 921 7 8 | LABORATOIRE CHAUVIN | FR |
| OPTICROM | not available | RVG 07564 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |

| 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 |
|---|---|---|---|---|
| OPTICROM 2 % collyre en solution | not available | BE111343 | MELISANA NV | BE |
| OPTICROM 2 % collyre en solution | not available | 0082619 | MELISANA NV | LU |
| OPTICROM 2 % oogdruppels, oplossing Natrium cromoglicaat | not available | BE111343 | MELISANA NV | BE |
| Opticrom 2% Augentropfen | not available | BE111343 | MELISANA NV | BE |
| Opticrom 2% Augentropfen | not available | 0082619 | MELISANA NV | LU |
| Opticrom 20 mg/ ml oldatos szemcsepp | not available | OGYI-T-3691/02 | SANOFI-AVENTIS ZRT | HU |
| Opticrom 20 mg/ml colirio en solución en envase unidosis | UK/H/5451/01/DC | 80267 | SANOFI-AVENTIS, S.A. | ES |
| Opticrom 20 mg/ml colirio en solución en envase unidosis | UK/H/5451/01/DC | 80267 | SANOFI-AVENTIS, S.A. | ES |
| Opticrom 20 mg/ml Colírio, solução em frasco conta-gotas multidose | not available | 8606004 | SANOFI - PRODUTOS FARMACEUTICOS LDA | PT |
| Opticrom 20mg/ml Colírio, Solução em Unidose | not available | 5593314 | SANOFI - PRODUTOS FARMACEUTICOS 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 |
|---|---|---|---|---|
| Opticrom 20mg/ml Colírio, Solução em Unidose | not available | 5593322 | SANOFI - PRODUTOS FARMACEUTICOS LDA | PT |
| Opticrom Allergy 2%w/v Eye Drops, solution | not available | PA 540/116/2 | SANOFI-AVENTIS IRELAND LTD | IE |
| Opticrom Allergy 2%w/v Eye Drops, solution | not available | PA 540/116/2 | SANOFI-AVENTIS IRELAND LTD | IE |
| Opticrom Allergy 2%w/v Eye Drops, solution | not available | PA 540/116/2 | SANOFI-AVENTIS IRELAND LTD | IE |
| OPTICROM ALLERGY 2.0% W/W EYE DROPS, SOLUTION | not available | PL 04425/0323 | AVENTIS PHARMA LTD | UK |
| OPTICROM ALLERGY 2.0% W/W EYE DROPS, SOLUTION | not available | PL 04425/0323 | AVENTIS PHARMA LTD | UK |
| Opticrom Allergy Single Dose 2% w/v Eye Drops, Solution | UK/H/5451/01/DC | PA 540/116/3 | SANOFI-AVENTIS IRELAND LTD | IE |
| Opticrom Allergy Single Dose 2% w/v Eye Drops, Solution | UK/H/5451/01/DC | PA 540/116/3 | SANOFI-AVENTIS IRELAND LTD | IE |
| Opticrom Allergy Single Dose 2% w/v Eye Drops, Solution | UK/H/5451/01/DC | PL 04425/0688 | AVENTIS PHARMA LTD | UK |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Opticrom Allergy Single Dose 2% w/v Eye Drops, Solution | UK/H/5451/01/DC | PL 04425/0688 | AVENTIS PHARMA LTD | UK |
| OPTICROM AQUEOUS / VISICROM EYE DROPS | not available | PL 04425/0324 | AVENTIS PHARMA LTD | UK |
| OPTICROM AQUEOUS / VISICROM EYE DROPS | not available | PL 04425/0324 | AVENTIS PHARMA LTD | UK |
| OPTICROM EYE DROPS | not available | PA 540/116/1 | SANOFI-AVENTIS IRELAND LTD | IE |
| OPTICROM EYE DROPS | not available | PA 540/116/1 | SANOFI-AVENTIS IRELAND LTD | IE |
| OPTICROM EYE DROPS | not available | PA 540/116/1 | SANOFI-AVENTIS IRELAND LTD | IE |
| OPTICROM HAYFEVER 2% W/V EYE DROPS | not available | PL 04425/0628 | AVENTIS PHARMA LTD | UK |
| OPTICROM HAYFEVER 2% W/V EYE DROPS | not available | PL 04425/0628 | AVENTIS PHARMA LTD | UK |
| OPTICROM OLDATOS SZEMCSEPP | not available | OGYI-T-3691/01 | SANOFI-AVENTIS ZRT | HU |
| OPTICROM UNIT DOSE | not available | RVG 12039 | SANOFI-AVENTIS NETHERLANDS B.V. | NL |
| OpticromTM Allergy 2.0% w/v Eye Drops, Solution | not available | MA 082/01901 | SANOFI MALTA LTD | MT |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| OpticromTM Allergy 2.0% w/v Eye Drops, Solution | not available | MA 082/01901 | SANOFI MALTA LTD | MT |
| OPTICRON 2 POUR CENT, collyre | not available | 320 298-0 | COOPERATION PHARMACEUTIQUE FRANCAISE | FR |
| OPTICRON UNIDOSE, collyre en récipient unidose | not available | 331 357.3 | COOPERATION PHARMACEUTIQUE FRANCAISE | FR |
| Optrex Allergy 2%w/v Eye Drops, Solution (eye drops) | not available | PL 35533/0033 | ASPIRE PHARMA LIMITED | UK |
| Optrex Allergy Eye Drops (sodium cromoglicate 2%) | not available | PL03468/0023 | BAUSCH & LOMB UK LTD. | UK |
| Optrex Hayfever Relief 2%w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| Pollenase Hayfever Relief 2%w/v Eye Drops | not available | PL 35533/0031 | ASPIRE PHARMA LIMITED | UK |
| SODIO CROMOGLICATO SANOFI | UK/H/5451/01/DC | 043692021 | SANOFI SPA | IT |
| SODIO CROMOGLICATO SANOFI | UK/H/5451/01/DC | 043692019 | SANOFI SPA | IT |
| Sodium Cromoglicate 2% w/v Eye Drops, Solution | not available | PL 35533/0032 | ASPIRE PHARMA LIMITED | UK |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Sodium Cromoglicate 2% w/v Eye Drops, Solution | UK/H/3500/001 | PL 25298/0033 | BROWN & BURK UK LIMITED | UK |
| Sodium Cromoglicate 2% w/v Eye Drops, Solution | UK/H/3500/001 | PL 25298/0033 | BROWN & BURK UK LIMITED | UK |
| Tesco Allergy Eye Drops sodium cromoglicate 2% | not available | PL03468/0023 | BAUSCH & LOMB UK LTD. | UK |
| Vivicrom Eye Drops | not available | PL 03468/0035 | BAUSCH & LOMB UK LTD. | UK |
| VIVIDRIN | not available | 75843/23.12.2015 | PHARMASWISS ČESKÁ REPUBLIKA S.R.O. | GR |
| VIVIDRIN | not available | 75843/23.12.2015 | PHARMASWISS ČESKÁ REPUBLIKA S.R.O. | GR |
| Vividrin 2% eye drops | not available | 11977 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | CY |
| Vividrin Antiallergic Eye Drops, Solution 2%w/v | not available | PA 555/5/2 | BAUSCH & LOMB UK LTD. | IE |
| Vividrin Eye Drops | not available | PL 03468/0035 | BAUSCH & LOMB UK LTD. | UK |
| Vividrin Hay Fever Eye Drops | not available | PL 03468/0035 | BAUSCH & LOMB UK LTD. | UK |
| Vividrin Preservative Free SDU 2% w/v Eye Drops solution | not available | PA 555/5/3 | BAUSCH & LOMB UK LTD. | IE |

| Product name (in authorisation country) | MRP / DCP Authorisation number | National Authorisation Number | MAH of product in the member state | Member State where product is authorised |
|---|---|---|---|---|
| Vividrin® antiallergische Augentropfen | not available | 4552.00.00 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | DE |
| Vividrin® antiallergische Augentropfen | not available | 1995012984 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | LU |
| Vividrin® iso EDO® antiallergische Augentropfen | not available | 30622.00.00 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | DE |
| Vividrin® MDO® | not available | 40229.00.00 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | DE |
| Vividrin® Nasenspray gegen Heuschnupfen | not available | 4553.00.00 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | DE |
| Vividrin® Nasenspray gegen Heuschnupfen | not available | 1989060399 | DR. GERHARD MANN CHEM.-PHARM. FABRIK GMBH | LU |