EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

03 October 2024
EMA/423872/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): dorzolamide / timolol

Procedure No. PSUSA/00001166/202402

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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
Amiptic 20 mg/5 mg/ml acu pilieni, šķīdums	PL/H/0616/001	13-0099	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
Amiptic 20 mg/ml + 5 mg/ml oční kapky, roztok	PL/H/0616/001	64/180/13-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Amiptinon 20 mg/5 mg/ml acu pilieni, šķīdums	PL/H/0446/001	17-0150	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
Arzotilol 20 mg/ml + 5 mg/ml augndropar, lausn	SE/H/1771/001	IS/1/11/035/01	TEVA B.V	IS
Betaclar 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0297/001	041858010	OMNIVISION ITALIA S.R.L.	IT
Betaclar 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0297/001	041858022	OMNIVISION ITALIA S.R.L.	IT
Betaclar 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0297/001	041858034	OMNIVISION ITALIA S.R.L.	IT
Codimaz, 20 mg/ml + 5 mg/ml, eye drops, solution	not available	PL 29556/0004	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Cosduo 20 mg/ml + 5 mg/ml colirio en solución	DK/H/2566/001	82086	SANTEN OY	ES
Cosduo 20 mg/ml + 5mg/ml Augentropfen, Lösung	DK/H/2566/001	95687.00.00	SANTEN OY	DE
COSIDIME 20 mg/ml + 5 mg/ml, collyre en solution	DK/H/2566/001	34009 300 850 2 8	SANTEN OY	FR
COSIDIME 20 mg/ml + 5 mg/ml, collyre en solution	DK/H/2566/001	134009 301 343 5 1	SANTEN OY	FR
COSIDIME 20 mg/ml + 5 mg/ml, collyre en solution	DK/H/2566/001	34009 300 850 3 5	SANTEN OY	FR
COSIDIME 20 mg/ml + 5 mg/ml, collyre en solution	DK/H/2566/001	34009 301 343 6 8	SANTEN OY	FR

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COSIDIME 20 mg/ml + 5 mg/ml, collyre en solution	DK/H/2566/001	34009 550 298 1 6	SANTEN OY	FR
COSOPT ® Unit Dose 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DK/H/134/02	BE 291663	SANTEN OY	BE
COSOPT 20 mg/5 mg v 1 ml kapljice za oko, raztopina	not available	H/99/00425/002	SANTEN OY	SI
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/003	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/005	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/004	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/007	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/008	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas vienadozėje talpyklėje)	DK/H/134/02	LT/1/98/0082/006	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas)	not available	LT/1/98/0082/001	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml akių lašai (tirpalas)	not available	LT/1/98/0082/002	SANTEN OY	LT
COSOPT 20 mg/5 mg/ml, acu pilieni, šķīdums	not available	99 - 0966	SANTEN OY	LV
Cosopt 20 mg/ml + 5 mg/ml colírio, solução em recipiente unidose	DK/H/134/02	5920988	SANTEN OY	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
(sem conservantes)				
Cosopt 20 mg/ml + 5 mg/ml colírio, solução em recipiente unidose (sem conservantes)	DK/H/134/02	5077722	SANTEN OY	PT
Cosopt 20 mg/ml + 5 mg/ml oldatos szemcsepp	not available	OGYI-T-7662/01	SANTEN OY	HU
Cosopt 20 mg/ml + 5 mg/ml øyedråper, oppløsning med konserveringsmiddel	not available	97-4128	SANTEN OY	NO
Cosopt 20 mg/ml + 5 mg/ml øyedråper, oppløsning uten konserveringsmiddel	not available	07-05545	SANTEN OY	NO
COSOPT 20 mg/ml + 5 mg/ml picaturi oftalmice, solutie	not available	6829/2014/01	SANTEN OY	RO
COSOPT 20 mg/ml + 5 mg/ml picaturi oftalmice, solutie	not available	6829/2014/02	SANTEN OY	RO
Cosopt 20 mg/ml + 5 mg/ml silmätipat, liuos, kerta-annospakkaus	DK/H/134/02	21944	SANTEN OY	FI
COSOPT 20 mg/ml + 5 mg/ml, collyre en solution en récipient unidose	DK/H/134/02	34009 377 057 2 1	SANTEN OY	FR
COSOPT 20 mg/ml + 5 mg/ml, collyre en solution en récipient unidose	DK/H/134/02	34009 377 058 9 9	SANTEN OY	FR
COSOPT 20 mg/ml + 5 mg/ml, kapi za oko, otopina	not available	HR-H-438170589-01	SANTEN OY	HR
COSOPT 20 mg/ml + 5	not available	HR-H-438170589-02	SANTEN OY	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml, kapi za oko, otopina				
COSOPT 20 mg/ml + 5 mg/ml, očná roztoková instilácia	not available	64/0218/00-S	SANTEN OY	SK
COSOPT 20 mg/ml + 5 mg/ml, oční kapky, roztok	not available	64/634/99-C	SANTEN OY	CZ
Cosopt 20 mg/ml + 5 mg/ml, ögondroppar, lösning i endosbehållare	DK/H/134/02	23456	SANTEN OY	SE
COSOPT 20 mg/ml +5 mg/ml οφθαλμικές σταγόνες, διάλυμα	not available	18201	VIANEX S.A.	CY
COSOPT 20 mg/ml +5 mg/ml οφθαλμικές σταγόνες, διάλυμα	DK/H/134/001	110211/09-11-2018	VIANEX S.A.	GR
Cosopt bez konzervačních přísad 20 mg/ml + 5 mg/ml oční kapky, roztok	DK/H/0134/003	64/118/17-C	SANTEN OY	CZ
COSOPT brez konzervansa 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DK/H/0134/003	H/06/01960/004	SANTEN OY	SI
COSOPT brez konzervansa 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DK/H/0134/003	H/06/01960/005	SANTEN OY	SI
COSOPT brez konzervansa 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DK/H/0134/003	H/06/01960/006	SANTEN OY	SI
COSOPT brez konzervansa 20 mg/ml + 5 mg/ml kapljice za oko, raztopina v	DK/H/134/02	H/06/01960/001	SANTEN OY	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
enoodmernem vsebniku				
COSOPT brez konzervansa 20 mg/ml + 5 mg/ml kapljice za oko, raztopina v enoodmernem vsebniku	DK/H/134/02	H/06/01960/002	SANTEN OY	SI
COSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing, flacon voor eenmalig gebruik 20 mg/ml + 5 mg/ml	DK/H/134/02	RVG 33660	SANTEN OY	NL
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie	DK/H/0134/003	11102/2018/01	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie	DK/H/0134/003	11102/2018/02	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie	DK/H/0134/003	11102/2018/03	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/01	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/02	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/03	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/04	SANTEN OY	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
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/05	SANTEN OY	RO
COSOPT fără conservant 20 mg/ml+5 mg/ml picături oftalmice, soluţie în recipient unidoză	DK/H/134/02	9197/2016/06	SANTEN OY	RO
COSOPT FREE 20 mg/ml + 5 mg/ml, očná roztoková instilácia v jednodávkovom obale	DK/H/134/02	64/0218/11-S	SANTEN OY	SK
Cosopt FREE bez konzervačních přísad 20 mg/ml + 5 mg/ml oční kapky, roztok v jednodávkovém obalu	DK/H/134/02	64/265/06-C	SANTEN OY	CZ
COSOPT iMulti 20 mg/ml + 5 mg/ml eye drops, solution	DK/H/0134/003	PL 16058/0025	SANTEN OY	XI
COSOPT iMulti 20 mg/ml + 5 mg/ml kapi za oko, otopina	DK/H/0134/003	HR-H-149562311-01	SANTEN OY	HR
COSOPT iMulti 20 mg/ml + 5 mg/ml kapi za oko, otopina	DK/H/0134/003	HR-H-149562311-02	SANTEN OY	HR
COSOPT iMulti 20 mg/ml + 5 mg/ml kapi za oko, otopina	DK/H/0134/003	HR-H-149562311-03	SANTEN OY	HR
COSOPT iMulti 20 mg/ml +5 mg/ml οφθαλμικές σταγόνες, διάλυμα	DK/H/0134/003	022891	VIANEX S.A.	CY
COSOPT iMulti 20 mg/ml +5 mg/ml οφθαλμικές σταγόνες, διάλυμα	DK/H/0134/003	91256/04-09-2023	VIANEX S.A.	GR
COSOPT iMulti Ukonserveret	DK/H/0134/003	59349	SANTEN OY	DK

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
COSOPT Multi 20 mg/ml + 5 mg/ml colírio, solução	DK/H/0134/003	5750062	SANTEN OY	PT
COSOPT Multi 20 mg/ml + 5 mg/ml tartósítószermentes oldatos szemcsepp	DK/H/0134/003	OGYI-T-7662/05	SANTEN OY	HU
COSOPT Multi 20 mg/ml + 5 mg/ml tartósítószermentes oldatos szemcsepp	DK/H/0134/003	OGYI-T-7662/06	SANTEN OY	HU
COSOPT Multi 20 mg/ml + 5 mg/ml tartósítószermentes oldatos szemcsepp	DK/H/0134/003	OGYI-T-7662/07	SANTEN OY	HU
COSOPT Multi Dose Free 20 mg/ml + 5 mg/ml očná roztoková instilácia	DK/H/0134/003	64/0355/18-S	SANTEN OY	SK
COSOPT Multidose conserveermiddelvrij 20 mg/ml + 5 mg/ml oogdruppels, oplossing	DK/H/0134/003	RVG 120935	SANTEN OY	NL
COSOPT PF 20 mg/5 mg/ml acu pilieni, šķīdums	DK/H/0134/003	18-0093	SANTEN OY	LV
COSOPT PF 20 mg/5 mg/ml, acu pilieni, šķīdums, vienreizējas devas traciņā	DK/H/134/02	11-0059	SANTEN OY	LV
COSOPT PF 20 mg/ml + 5 mg/ml colirio en solución	DK/H/0134/003	83389	SANTEN OY	ES
COSOPT PF 20 mg/ml + 5 mg/ml kapi za oko, otopina u jednodoznom spremniku	not available	HR-H-762834161	SANTEN OY	HR
Cosopt PF Multi, 20	DK/H/0134/003	25030	SANTEN OY	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
mg/ml + 5 mg/ml, krople do oczu, roztwór				
COSOPT PF, 20 mg/ml + 5 mg/ml, krople do oczu, roztwór w pojemniku jednodawkowym	DK/H/134/02	18307	SANTEN OY	PL
COSOPT Preservative-Free 20 mg/ml + 5 mg/ml, eye drops, solution in single-dose container	DK/H/134/02	PA0879/005/002	SANTEN OY	IE
COSOPT Preservative-Free 20 mg/ml + 5 mg/ml, eye drops, solution in single-dose container	DK/H/134/02	PL 16058/0015	SANTEN OY	XI
COSOPT senza conservante 20 mg/ml + 5 mg/ml, collirio, soluzione	DK/H/0134/003	034242077	SANTEN OY	IT
COSOPT senza conservante 20 mg/ml + 5 mg/ml, collirio, soluzione	DK/H/0134/003	034242089	SANTEN OY	IT
COSOPT senza conservante 20 mg/ml + 5 mg/ml, collirio, soluzione	DK/H/0134/003	034242091	SANTEN OY	IT
COSOPT sine 20 mg/5 mg/ml akių lašai (tirpalas)	DK/H/0134/003	LT/1/18/4276/001	SANTEN OY	LT
COSOPT sine 20 mg/5 mg/ml akių lašai (tirpalas)	DK/H/0134/003	LT/1/18/4276/002	SANTEN OY	LT
COSOPT sine 20 mg/5 mg/ml akių lašai (tirpalas)	DK/H/0134/003	LT/1/18/4276/003	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
COSOPT sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DK/H/0134/003	99981.00.00	SANTEN OY	DE
Cosopt sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DK/H/134/02	1-26570	SANTEN OY	AT
COSOPT sine 20 mg/ml + 5 mg/ml ögondroppar, lösning	DK/H/0134/003	35108	SANTEN OY	FI
Cosopt sine 20 mg/ml + 5 mg/ml ögondroppar, lösning	DK/H/0134/003	56217	SANTEN OY	SE
COSOPT sine 20 mg/ml + 5 mg/ml øyedråper, oppløsning	DK/H/0134/003	17-11651	SANTEN OY	NO
COSOPT sine 20 mg/ml + 5 mg/ml silmätipat, liuos	DK/H/0134/003	35108	SANTEN OY	FI
COSOPT sine 20 mg/ml + 5 mg/ml, augndropar, lausn	DK/H/0134/003	IS/1/18/058/01	SANTEN OY	IS
COSOPT Sine Conservans 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DK/H/0134/003	BE530337	SANTEN OY	BE
COSOPT Sine Conservans 20 mg/ml + 5 mg/ml collyre en solution	DK/H/0134/003	BE530337	SANTEN OY	BE
COSOPT Sine Conservans 20 mg/ml + 5 mg/ml collyre en solution	DK/H/0134/003	2018110318	SANTEN OY	LU
COSOPT Sine Conservans 20 mg/ml + 5 mg/ml collyre en	DK/H/0134/003	2018110318	SANTEN OY	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
solution				
COSOPT Sine Conservans 20 mg/ml + 5 mg/ml oogdruppels, oplossing	DK/H/0134/003	BE530337	SANTEN OY	BE
Cosopt Ukonserveret 20 mg/ml + 5 mg/ml øjendråber, opløsning i enkeltdosisbeholder	DK/H/134/02	36113	SANTEN OY	DK
COSOPT Unit Dose 20 mg/ml + 5 mg/ml collyre en solution en récipient unidose	DK/H/134/02	BE 291663	SANTEN OY	BE
COSOPT Unit Dose 20 mg/ml + 5 mg/ml collyre en solution en récipient unidose	DK/H/134/02	2007060029	SANTEN OY	LU
COSOPT UNIT DOSE 20 mg/ml + 5 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	DK/H/134/02	BE 291663	SANTEN OY	BE
COSOPT UNO 20 mg/ml + 5 mg/ml oldatos szemcsepp egyadagos tartályban	DK/H/134/02	OGYI-T-7662/02	SANTEN OY	HU
COSOPT UNO 20 mg/ml + 5 mg/ml oldatos szemcsepp egyadagos tartályban	DK/H/134/02	OGYI-T-7662/03	SANTEN OY	HU
COSOPT UNO 20 mg/ml + 5 mg/ml oldatos szemcsepp egyadagos tartályban	DK/H/134/02	OGYI-T-7662/04	SANTEN OY	HU
Cosopt, (20 mg+5 mg)/ml, krople do oczu, roztwór	not available	4424	SANTEN OY	PL

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COSOPT, 20 mg/5 mg/ml silmatilgad, lahus	not available	282199	SANTEN OY	EE
COSOPT-S® 20 mg/ml + 5 mg/ml Augentropfen im Einzeldosisbehältnis	DK/H/134/02	2007060029	SANTEN OY	LU
Costad, øjendråber, opløsning	DE/H/5435/001	45290	STADA ARZNEIMITTEL AG	DK
COSTEC 20 mg/mL + 5 mg/mL, collyre en solution	DK/H/0134/003	34009 301 584 9 4	SANTEN OY	FR
COSTEC 20 mg/mL + 5 mg/mL, collyre en solution	DK/H/0134/003	34009 301 585 0 0	SANTEN OY	FR
COSTEC 20 mg/mL + 5 mg/mL, collyre en solution	DK/H/0134/003	34009 301 585 1 7	SANTEN OY	FR
Dorlatim 20 mg/ml + 5 mg/ml ögondroppar, lösning	DK/H/2977/001	58264	SANDOZ A/S	SE
Dorlatim, øjendråber, opløsning	DK/H/2977/001	61642	SANDOZ A/S	DK
Dortimvision, 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	AT/H/0486/001	48666	OMNIVISION GMBH	SE
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 301 917 3 6	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 301 917 4 3	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 301 917 5 0	PHARMA STULLN	FR
DORTIRUS 20 mg/mL +	AT/H/0736/001	34009 301 917 6 7	PHARMA STULLN	FR

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5 mg/mL, collyre en solution en récipient unidose				
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 301 917 7 4	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 550 694 3 0	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 550 694 4 7	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 302 208 0 1	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 302 208 1 8	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 302 208 2 5	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 302 208 4 9	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 302 208 5 6	PHARMA STULLN	FR
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en	AT/H/0736/001	34009 550 792 5 5	PHARMA STULLN	FR

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solution en récipient unidose				
DORTIRUS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	AT/H/0736/001	34009 550 792 6 2	PHARMA STULLN	FR
Dorvis plus 20 mg/ml + 5 mg/ml kapi za oko, otopina	not available	HR-H-091461330	ZENTIVA, K.S.	HR
DORZAMOX 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0727/001	040815019	DOC GENERICI S.R.L.	IT
DORZAMOX 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0727/001	040815021	DOC GENERICI S.R.L.	IT
DORZAMOX 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0727/001	040815033	DOC GENERICI S.R.L.	IT
Dorzastad 20 mg/ml + 5 mg/ml Augentropfen	DE/H/5435/001	1-30191	STADA ARZNEIMITTEL GMBH	AT
Dorzastad sine 20 mg/ml + 5 mg/ml Augentropfen, im Einzeldosisbehältnis	AT/H/1021/001	141186	STADA ARZNEIMITTEL GMBH	AT
Dorzo plus T STADA® 20 mg/ml + 5 mg/ml Augentropfen	DE/H/5435/001	79050.00.00	STADAPHARM GMBH	DE
Dorzocomp-Stulln 20 mg/ml + 5 mg/ml Augentropfen, Lösung	AT/H/0735/001	99326.00.00	PHARMA STULLN	DE
Dorzocomp-Stulln sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0736/001	99327.00.00	PHARMA STULLN	DE
DorzoComp-Vision 20 mg/ml + 5 mg/ml Augentropfen, Lösung	AT/H/1008/001	1-30190	OMNIVISION GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DorzoComp-Vision 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5998/001	79049.00.00	OMNIVISION GMBH	DE
DorzoComp-Vision sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0993/001/DC	139284	OMNIVISION GMBH	AT
DorzoComp-Vision sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0993/001/DC	2204211.00.00	OMNIVISION GMBH	DE
Dorzogen Combi 20 mg/ml + 5 mg/ml oční kapky, roztok	AT/H/1033/001	64/349/11-C	VIATRIS LIMITED	CZ
Dorzolamid + Timolol 1A Pharma 20 mg/ml + 5mg/ml - Augentropfen, Lösung	DK/H/2977/001	139158	1A PHARMA GMBH	AT
Dorzolamid + Timolol Arcana 20 mg/ml + 5 mg/ml Augentropfen	AT/H/1033/001	1-30171	ARCANA ARZNEIMITTEL GMBH	AT
Dorzolamid + Timolol Pharma Stulln 20 mg/ml + 5 mg/ml Augentropfen, Lösung	AT/H/0992/001	139285	PHARMA STULLN	AT
Dorzolamid + Timolol Pharma Stulln 20 mg/ml + 5 mg/ml Augentropfen, Lösung	AT/H/0992/001	2204210.00.00	PHARMA STULLN	DE
Dorzolamid + Timolol Stulln 20 mg/ml + 5 mg/ml Augentropfen, Lösung	AT/H/0735/001	139286	PHARMA STULLN	AT
Dorzolamid + Timolol Stulln sine 20 mg/ml + 5 mg/ml Augentropfen, Lösung im	AT/H/0736/001	139283	PHARMA STULLN	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Einzeldosisbehältnis				
Dorzolamid + Timolol TRB 20 mg/ml + 5 mg/ml Augentropfen	not available	89461.00.00	RAFARM SA.	DE
Dorzolamid AL comp. 20 mg/ml + 5 mg/ml Augentropfen	DE/H/5600/001	79051.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid comp - 1 A Pharma 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DK/H/2977/001	2202751.00.00	1 A PHARMA GMBH	DE
DORZOLAMID COMP. STULLN 20 mg/ml + 5 mg/ml Augentropfen, Lösung	not available	79269.00.00	PHARMA STULLN	DE
Dorzolamid sine AL comp. 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/1021/001	2204620.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid/ Timolol FDC Pharma 20 mg/ml / 5 mg/ml Augentropfen	DE/H/6357/001	80899	ASCEND GMBH	DE
Dorzolamid/Timolol "Medical Valley", øjendråber, opløsning	DK/H/3205/001	64861	MEDICAL VALLEY INVEST AB	DK
Dorzolamid/timolol "Pharmathen", øjendråber, opløsning	DK/H/2566/001	56282	PHARMATHEN S.A.	DK
Dorzolamid/Timolol Actavis 20 mg/ml + 5 mg/ml ögondroppar, lösning	SE/H/1771/001	42776	ACTAVIS GROUP PTC EHF.	SE
Dorzolamid/Timolol Heumann 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/6269/001	2203914.00.00	HEUMANN PHARMA GMBH & CO. GENERICA	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
Dorzolamid/Timolol Medical Valley 20 mg/ml + 5 mg/ml ögondroppar, lösning	DK/H/3205/001	61017	MEDICAL VALLEY INVEST AB	SE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5352/001	89000.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5352/001	89000.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5352/001	89000.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5352/001	89000.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/5352/001	89000.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen,	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lösung im Einzeldosisbehältnis.				
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Micro Labs 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis.	DE/H/6622/001	7000180.00.00	MICRO LABS GMBH	DE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning	SE/H/2246/001	63282	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning	SE/H/2246/01/DC	63282	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5	SE/H/2246/01/DC	63282	MISOM LABS LIMITED	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
mg/ml ögondroppar, lösning				
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning	SE/H/2246/01/DC	63282	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning	SE/H/2246/01/DC	63282	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar,	SE/H/2252/001	63119	MISOM LABS LIMITED	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
lösning i endosbehållare				
Dorzolamid/Timolol Misom 20 mg/ml + 5 mg/ml ögondroppar, lösning i endosbehållare	SE/H/2252/001	63119	MISOM LABS LIMITED	SE
Dorzolamid/timolol Olikla 20 mg/ml + 5 mg/ml očné roztokové kvapky	CZ/H/0762/001	64/0371/18-S	OLIKLA S.R.O.	SK
Dorzolamid/timolol Olikla 20 mg/ml + 5 mg/ml oční kapky, roztok	CZ/H/0762/001	64/238/17-C	OLIKLA S.R.O.	CZ
Dorzolamid/Timolol OmniVision 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0486/001	89362.00.00	OMNIVISION GMBH	DE
Dorzolamid/Timolol OmniVision20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0486/001	135195	OMNIVISION GMBH	AT
Dorzolamida /Timolol Stulln 20 mg/ml + 5 mg/ml colirio en solución	AT/H/0735/001	85060	PHARMA STULLN	ES
Dorzolamida + Timolol Bruschettini, 20 mg/ml + 5 mg/ml, colírio, solução	not available	5578356	BRUSCHETTINI S.R.L	PT
Dorzolamida + Timolol Bruschettini, 20 mg/ml + 5 mg/ml, colírio, solução	not available	5578356	BRUSCHETTINI S.R.L	PT
Dorzolamida/Timolol Aurovitas 20 mg/ml + 5 mg/ml colirio en solución	PT/H/2116/001	74.906	AUROVITAS SPAIN,S.A.U.	ES
Dorzolamida/Timolol Farmalider 20 mg/ml + 5mg/ ml colirio en solución.	DE/H/6357/001	75037	FARMALIDER, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dorzolamida/Timolol MEIJI 20 mg/ml + 5 mg/ml colirio en solución.	not available	78.599	MEIJI PHARMA SPAIN, S.A.	ES
Dorzolamidă/Timolol Misom 20 mg/ml + 5 mg/ml picături oftalmice, soluție	SE/H/2246/001	15291/2024/01	MISOM LABS LIMITED	RO
Dorzolamidă/Timolol Misom 20 mg/ml + 5 mg/ml picături oftalmice, soluție	SE/H/2246/001	15291/2024/02	MISOM LABS LIMITED	RO
Dorzolamidă/Timolol Misom 20 mg/ml + 5 mg/ml picături oftalmice, soluție	SE/H/2246/001	15291/2024/03	MISOM LABS LIMITED	RO
Dorzolamidă/Timolol Misom 20 mg/ml + 5 mg/ml picături oftalmice, soluție	SE/H/2246/001	15291/2024/04	MISOM LABS LIMITED	RO
Dorzolamidă/Timolol Misom 20 mg/ml + 5 mg/ml picături oftalmice, soluție	SE/H/2246/001	15291/2024/05	MISOM LABS LIMITED	RO
Dorzolamida/Timolol STADA 20 mg/ml + 5 mg/ml colirio en solución	DE/H/5435/001	74.610	LABORATORIO STADA, S.L.	ES
Dorzolamida/Timolol Stulln PF 20 mg/ml + 5 mg/ml colirio en solución en envase unidosis	AT/H/0736/001	84817	PHARMA STULLN	ES
Dorzolamida/Timolol Viatris 20 mg/ml + 5 mg/ml colirio	AT/H/1033/001	74690	VIATRIS LIMITED	ES
Dorzolamida/Timolol VIR 20 mg/ml + 5 mg/ml colirio en solución	PT/H/0645/001	77386	INDUSTRIA QUÍMICA Y FARMACEÚTICA VIR, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dorzolamide / Timolol 20 mg/ml + 5 mg/ml Eye Drops, Solution	not available	PL 18956/0019	MEDICOM HEALTHCARE LTD	XI
Dorzolamide / Timolol 20 mg/ml + 5 mg/ml eye drops, solution	IT/H/0297/001	PL 01883/0358	MACARTHYS LABORATORIES LTD	XI
Dorzolamide + Timolol Stulln 20 mg/ml + 5 mg/ml oogdruppels, oplossing	AT/H/0735/001	RVG 120482	PHARMA STULLN	NL
Dorzolamide + Timolol Stulln zonder conserveermiddel 20 mg/ml + 5 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	AT/H/0736/001	RVG 120553	PHARMA STULLN	NL
Dorzolamide 20 mg/ml and Timolol 5 mg/ml Eye Drops, Solution BP	not available	PL 35638/0005	FDC PHARMA	XI
Dorzolamide and Timolol 20 mg/ml + 5 mg/ml, eye drops solution	DK/H/2477/001	PA23259/003/001	BAUSCH + LOMB IRELAND LIMITED	IE
Dorzolamide e Timololo Aurobindo 20 mg/ml + 5 mg/ml collirio, soluzione	PT/H/2116/001/DC	041137011	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide e Timololo Aurobindo 20 mg/ml + 5 mg/ml collirio, soluzione	PT/H/2116/001/DC	041137023	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide e Timololo Aurobindo 20 mg/ml + 5 mg/ml collirio, soluzione	PT/H/2116/001	041137035	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide e Timololo EG 20 mg/ml + 5 mg/ml collirio, soluzione	DE/H/5435/001	040797019	EG S.P.A.	IT
Dorzolamide e Timololo EG 20 mg/ml + 5 mg/ml	DE/H/5435/001	040797021	EG 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
collirio, soluzione				
Dorzolamide e Timololo EG 20 mg/ml + 5 mg/ml collirio, soluzione	DE/H/5435/001	040797033	EG S.P.A.	IT
Dorzolamide e Timololo Mylan Generics 20 mg/ml + 5 mg/ml, collirio	AT/H/1033/001	040792018	MYLAN S.P.A.	IT
Dorzolamide e Timololo Mylan Generics 20 mg/ml + 5 mg/ml, collirio	AT/H/1033/001	040792020	MYLAN S.P.A.	IT
Dorzolamide e Timololo Mylan Generics 20 mg/ml + 5 mg/ml, collirio	AT/H/1033/001	040792032	MYLAN S.P.A.	IT
DORZOLAMIDE E TIMOLOLO PENSA 20 mg/ml/ 5 mg/ml collirio, soluzione	not available	043861018	TOWA PHARMACEUTICAL.	IT
Dorzolamide e timololo Sandoz 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2977/001	047051053	SANDOZ S.P.A.	IT
Dorzolamide e timololo Sandoz 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2977/001	047051040	SANDOZ S.P.A.	IT
Dorzolamide e timololo Sandoz 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2977/001	047051014	SANDOZ S.P.A.	IT
Dorzolamide e timololo Sandoz 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2977/001	047051026	SANDOZ S.P.A.	IT
Dorzolamide e timololo Sandoz 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2977/001	047051038	SANDOZ S.P.A.	IT
Dorzolamide e Timololo Zentiva 20 mg/ml + 5	IT/H/0684/001	041580022	ZENTIVA 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
mg/ml collirio, soluzione				
Dorzolamide e Timololo Zentiva 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0684/001	041580010	ZENTIVA ITALIA S.R.L.	IT
Dorzolamide e Timololo Zentiva 20 mg/ml + 5 mg/ml collirio, soluzione	IT/H/0684/001	041580034	ZENTIVA ITALIA S.R.L.	IT
Dorzolamide/ Timolol 20 mg/ml + 5 mg/ml eye drops, solution in single-dose container	not available	PL35533/0086	ASPIRE PHARMA LIMITED	XI
Dorzolamide/Timolol "Stada"	AT/H/1021/001	62980	STADA ARZNEIMITTEL AG	DK
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution	AT/H/1033/001	PL 04569/1035	GENERICS [UK] LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution	not available	PL 35533/0147	ASPIRE PHARMA LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution	not available	PL31103/0012	BLUMONT PHARMA LTD	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution	not available	PL 25298/0257	BROWN & BURK UK LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution in single-dose container	SE/H/1966/001	PL31103/0026	BLUMONT PHARMA LTD	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml eye drops, solution.	IT/H/0684/001	PL 17780/0575	ZENTIVA PHARMA UK LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml, Preservative-Free, Eye drops, solution in single-dose container	not available	PL 25298/0252	BROWN & BURK 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
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml, Preservative-Free, Eye drops, solution in single-dose container	not available	PL 25298/0252	BROWN & BURK UK LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml, Preservative-Free, Eye drops, solution in single-dose container	not available	PL 25298/0252	BROWN & BURK UK LIMITED	XI
Dorzolamide/Timolol 20 mg/ml + 5 mg/ml, Preservative-Free, Eye drops, solution in single-dose container	not available	PL 25298/0252	BROWN & BURK UK LIMITED	XI
Dorzolamide/Timolol 20mg/ml + 5mg/ml Eye Drops, Solution	not available	PL 0142/0988	ACCORD-UK LIMITED	XI
DORZOLAMIDE/TIMOLOL ACTAVIS 20 mg/5 mg/ml acu pilieni, škidums	SE/H/1771/001	11-0103	TEVA B.V	LV
Dorzolamide/Timolol Actavis 20 mg/5 mg/ml akiu lasai tirpalas	SE/H/1771/001	LT/1/11/2454/002	TEVA B.V	LT
Dorzolamide/Timolol Actavis 20 mg/5 mg/ml akiu lasai tirpalas	SE/H/1771/001	LT/1/11/2454/003	TEVA B.V	LT
Dorzolamide/Timolol Actavis 20 mg/5 mg/ml akiu lasai tirpalas	SE/H/1771/001	LT/1/11/2454/001	TEVA B.V	LT
Dorzolamide/Timolol Alvogen 20 mg/ml + 5 mg/ml augndropar, lausn.	IS/H/0374/001	IS/1/10/142/01	ALVOGEN EHF	IS
Dorzolamide/Timolol Aurobindo 20/5 mg/ml, oogdruppels, oplossing	PT/H/2116/001	RVG 105541	AUROBINDO PHARMA 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
DORZOLAMIDE/TIMOLOL BGR 20 mg/ml + 5 mg/ml, collyre en solution	not available	34009 301 226 9 3	BIOGARAN	FR
DORZOLAMIDE/TIMOLOL BGR 20 mg/ml + 5 mg/ml, collyre en solution	not available	3400955045106	BIOGARAN	FR
DORZOLAMIDE/TIMOLOL BGR 20 mg/ml + 5 mg/ml, collyre en solution	not available	3400955045083	BIOGARAN	FR
DORZOLAMIDE/TIMOLOL CRISTERS 20 mg/ml + 5 mg/ml, collyre en solution	not available	34009 300 262 5 0	CRISTERS	FR
DORZOLAMIDE/TIMOLOL CRISTERS 20 mg/ml + 5 mg/ml, collyre en solution	not available	34009 550 091 4 6	CRISTERS	FR
DORZOLAMIDE/TIMOLOL CRISTERS 20 mg/ml + 5 mg/ml, collyre en solution	not available	34009 550 091 5 3	CRISTERS	FR
Dorzolamide/Timolol EG 20 mg/ml + 5 mg/ml Augentropfen	DE/H/5435/001	BE391447	EUROGENERICS N.V./S.A.	BE
Dorzolamide/Timolol EG 20 mg/ml + 5 mg/ml collyre en solution	DE/H/5435/001	BE391447	EUROGENERICS N.V./S.A.	BE
Dorzolamide/Timolol EG 20 mg/ml + 5 mg/ml collyre en solution	DE/H/5435/001	2012010020	EUROGENERICS N.V./S.A.	LU
Dorzolamide/Timolol EG 20 mg/ml + 5 mg/ml oogdruppels, oplossing	DE/H/5435/001	BE391447	EUROGENERICS N.V./S.A.	BE
DORZOLAMIDE/TIMOLOL	DE/H/5435/001	NL38129	EG LABO LABORATOIRES	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
EG 20 mg/ml + 5 mg/ml, collyre en solution			EUROGENERICS	
Dorzolamide/Timolol ELVIM 20 mg/5 mg/ml acu pilieni, šķidums	LV/H/0190/001/DC	11-0062	SIA ELVIM	LV
Dorzolamide/Timolol ELVIM 20 mg/5 mg/ml akių lašai (tirpalas)	LV/H/0190/001	LT/1/11/2420/002	SIA ELVIM	LT
Dorzolamide/Timolol ELVIM 20 mg/5 mg/ml akių lašai (tirpalas)	LV/H/0190/001	LT/1/11/2420/001	SIA ELVIM	LT
Dorzolamide/Timolol ELVIM 20 mg/5 mg/ml akių lašai (tirpalas)	LV/H/0190/001	LT/1/11/2420/003	SIA ELVIM	LT
Dorzolamide/Timolol ELVIM, 20 mg/5 mg/ml silmatilgad, lahus	LV/H/0190/001	738511	SIA ELVIM	EE
Dorzolamide/Timolol Farmaprojects 20 mg/ml + 5 mg/ml, oční kapky, roztok	CZ/H/0973/001	64/439/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide/Timolol Indoco 20 mg/ml + 5 mg/ml oční kapky, roztok	CZ/H/0564/001	64/267/16-C	INDOCO REMEDIES CZECH S.R.O.	CZ
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 349 9 0	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 0 3	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL	DE/H/6622/001	34009 302 350 1 0	MICRO LABS GMBH	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
+ 5 mg/mL, collyre en solution en récipient unidose				
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 2 7	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 3 4	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 5 8	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 6 5	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICRO LABS 20 mg/mL + 5 mg/mL, collyre en solution en récipient unidose	DE/H/6622/001	34009 302 350 7 2	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICROLABS 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/5352/001	34009 301 614 2 5	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICROLABS 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/5352/001	34009 301 343 8 2	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL	DE/H/5352/001	34009 301 343 7 5	MICRO LABS GMBH	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
MICROLABS 20 mg/ml + 5 mg/ml, collyre en solution				
DORZOLAMIDE/TIMOLOL MICROLABS 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/5352/001	34009 550 595 5 4	MICRO LABS GMBH	FR
DORZOLAMIDE/TIMOLOL MICROLABS 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/5352/001	34009 550 501 4 8	MICRO LABS GMBH	FR
Dorzolamide/Timolol Mylan 20/5 mg/ml, oogdruppels, oplossing	AT/H/1033/001	RVG 105655	VIATRIS LIMITED	NL
Dorzolamide/Timolol Polpharma 20 mg/ml + 5 mg/ml ocní kapky, roztok	CZ/H/1113/001	64/214/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide/Timolol STADA 20 mg/ml + 5 mg/ml silmätipat, liuos, kerta-annospakkaus	AT/H/1021/001	37160	STADA ARZNEIMITTEL AG	FI
Dorzolamide/Timolol STADA 20 mg/ml + 5 mg/ml, ögondroppar, lösning i endosbehållare	AT/H/1021/001	59403	STADA ARZNEIMITTEL AG	SE
Dorzolamide/Timolol Stada 20+5 mg/ml augndropar, lausn í stakskammtaíláti	AT/H/1021/001	IS/1/22/039/01	STADA ARZNEIMITTEL AG	IS
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en solution	AT/H/0735/001	34009 301 915 9 0	PHARMA STULLN	FR
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en	AT/H/0735/001	34009 301 916 0 6	PHARMA STULLN	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
solution				
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en solution	AT/H/0735/001	34009 301 916 1 3	PHARMA STULLN	FR
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en solution	AT/H/0735/001	34009 301 916 2 0	PHARMA STULLN	FR
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en solution	AT/H/0735/001	34009 301 916 3 7	PHARMA STULLN	FR
DORZOLAMIDE/TIMOLOL STULLN 20 mg/mL + 5 mg/mL, collyre en solution	AT/H/0735/001	34009 301 916 5 1	PHARMA STULLN	FR
DORZOLAMIDE/TIMOLOL VIATRIS 20 mg/ml + 5 mg/ml, collyre en solution	AT/H/1033/001	NL 38247	VIATRIS SANTE	FR
DORZOLAMIDE/TIMOLOL ZENTIVA 20 mg/ml + 5 mg/ml, collyre en solution	IT/H/0684/001	34009 218 099 2 3	ZENTIVA FRANCE	FR
DORZOLAMIDE/TIMOLOL ZENTIVA 20 mg/ml + 5 mg/ml, collyre en solution	IT/H/0684/001	34009 218 096 3 3	ZENTIVA FRANCE	FR
DORZOLAMIDE/TIMOLOL ZENTIVA 20 mg/ml + 5 mg/ml, collyre en solution	IT/H/0684/001	34009 218 100 0 4	ZENTIVA FRANCE	FR
Dorzolamidum + Timololum Stulln, 20 mg/mL + 5 mg/mL, krople do oczu, roztwór	AT/H/0736/001	26039	PHARMA STULLN	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
w pojemniku jednodawkowym				
Dorzolamidum + Timololum Stulln, 20 mg/mL + 5 mg/mL, krople do oczu, roztwór	AT/H/0735/001	26038	PHARMA STULLN	PL
DORZYLEA 20 mg/ml + 5 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	AT/H/0736/001	99736/16/20-12-2019	PHARMA STULLN	GR
Dotizolil 20 mg/5 mg/ml acu pilieni, šķīdums	DK/H/2977/001	19-0119	SANDOZ PHARMACEUTICALS D.D.	LV
Dozopres Combi 20 mg/ml + 5 mg/ml oldatos szemcsepp	CZ/H/0937/001	OGYI-T-21780/01	BAUSCH + LOMB IRELAND LIMITED	HU
Dozopres Combi 20 mg/ml + 5 mg/ml oldatos szemcsepp	CZ/H/0937/001	OGYI-T-21780/02	BAUSCH + LOMB IRELAND LIMITED	HU
Dozopres Combi 20 mg/ml + 5 mg/ml oldatos szemcsepp	CZ/H/0937/001	OGYI-T-21780/03	BAUSCH + LOMB IRELAND LIMITED	HU
Dozopticum Duo 20 mg/ml + 5 mg/ml oldatos szemcsepp	DK/H/2477/001	OGYI-T-23066/01	BAUSCH + LOMB IRELAND LIMITED	HU
Dozopticum Duo 20 mg/ml + 5 mg/ml oldatos szemcsepp	DK/H/2477/001	OGYI-T-23066/02	BAUSCH + LOMB IRELAND LIMITED	HU
DOZOTIMA 20 mg/ml + 5 mg/ml oční kapky, roztok	CZ/H/0937/001	64/350/11-C	BAUSCH + LOMB IRELAND LIMITED	CZ
DUALKOPT 20 mg/ml + 5 mg/ml očná roztoková instilácia	DE/H/3682/001	64/0283/14-S	LABORATOIRES THEA	SK
DUALKOPT 20 mg/ml + 5 mg/ml ocní kapky, roztok	DE/H/3682/001	64/338/14-C	LABORATOIRES THEA	CZ
Dualkopt 20 mg/ml + 5	DE/H/3682/001	BE467404	LABORATOIRES THEA	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
mg/ml, Augentropfen, Lösung				
Dualkopt 20 mg/ml + 5 mg/ml, Augentropfen, Lösung	DE/H/3682/001	2015040064	LABORATOIRES THEA	LU
Dualkopt 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	BE467404	LABORATOIRES THEA	BE
DUALKOPT 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	34009 300 081 3 3	LABORATOIRES THEA	FR
DUALKOPT 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	34009 300 383 4 5	LABORATOIRES THEA	FR
DUALKOPT 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	34009 550 035 4 0	LABORATOIRES THEA	FR
DUALKOPT 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	34009 550 035 5 7	LABORATOIRES THEA	FR
Dualkopt 20 mg/ml + 5 mg/ml, collyre en solution	DE/H/3682/001	2015040064	LABORATOIRES THEA	LU
Dualkopt 20 mg/ml + 5 mg/ml, oogdruppels, oplossing	DE/H/3682/001	BE467404	LABORATOIRES THEA	BE
Dualkopt 20 mg/ml + 5 mg/ml, oogdruppels, oplossing	DE/H/3682/001	RVG 112681	LABORATOIRES THEA	NL
Duokopt 2|0 mg/ml + 5 mg/ml, ögondroppar, lösning	DE/H/3682/001	31096	LABORATOIRES THEA	FI
Duokopt 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/3682/001	135724	LABORATOIRES THEA	AT
Duokopt 20 mg/ml + 5 mg/ml Augentropfen,	DE/H/3682/001	88357.00.00	LABORATOIRES THEA	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
Lösung				
DUOKOPT 20 mg/ml + 5 mg/ml augndropar, lausn	DE/H/3682/001	IS/1/14/065/01	LABORATOIRES THEA	IS
DUOKOPT 20 mg/ml + 5 mg/ml kapi za oko, otopina	DE/H/3682/001	HR-H-620656079	LABORATOIRES THEA	HR
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/001	LABORATOIRES THEA	SI
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/002	LABORATOIRES THEA	SI
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/003	LABORATOIRES THEA	SI
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/004	LABORATOIRES THEA	SI
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/005	LABORATOIRES THEA	SI
DUOKOPT 20 mg/ml + 5 mg/ml kapljice za oko, raztopina	DE/H/3682/001	H/15/01987/006	LABORATOIRES THEA	SI
Duokopt 20 mg/ml + 5 mg/ml øyedråper, oppløsning	DE/H/3682/001	12-9254	LABORATOIRES THEA	NO
Duokopt 20 mg/ml + 5 mg/ml silmätipat, liuos	DE/H/3682/001	31096	LABORATOIRES THEA	FI
Duokopt 20 mg/ml + 5 mg/ml, ögondroppar, lösning	DE/H/3682/001	48572	LABORATOIRES THEA	SE
DUOKOPT 20 mg/ml + 5 mg/ml, οφθαλμικές σταγόνες, διάλυμα	DE/H/3682/001	22317	LABORATOIRES THEA	CY
DUOKOPT 20 mg/ml + 5 mg/ml, οφθαλμικές	DE/H/3682/001	39894/13-04-2020	LABORATOIRES THEA	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
σταγόνες, διάλυμα				
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/02	LABORATOIRES THEA	RO
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/05	LABORATOIRES THEA	RO
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/06	LABORATOIRES THEA	RO
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/04	LABORATOIRES THEA	RO
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/03	LABORATOIRES THEA	RO
DUOKOPT 20 mg/ml+5 mg/ml picături oftalmice soluţie	DE/H/3682/001	12977/2020/01	LABORATOIRES THEA	RO
Duokopt 20mg/ml + 5 mg/ml colirio en solución	DE/H/3682/001	79.101	LABORATOIRES THEA	ES
Duokopt 5 mg/ml + 20 mg/ml, colírio, solução	DE/H/3682/001	5628870	LABORATOIRES THEA	PT
DUOKOPT, 20 mg/ml + 5 mg/ml, krople do oczu, roztwór (Dorzolamidum + Timololum)	DE/H/3682/001	22333	LABORATOIRES THEA	PL
Duokopt, øjendråber, opløsning	DE/H/3682/001	51620	LABORATOIRES THEA	DK
Eylamdo 20 mg/ml + 5 mg/ml, eye drops solution	DK/H/2566/001	PL35533/0117	ASPIRE PHARMA LIMITED	XI
Eylamdo 20 mg/ml + 5 mg/ml, eye drops solution	DK/H/2566/001	PL35533/0117	ASPIRE PHARMA LIMITED	XI
EYROOBI 20 mg/ml + 5	DK/H/2566/001	044738045	FIDIA FARMACEUTICI S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml collirio, soluzione				
EYROOBI 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2566/001	044738021	FIDIA FARMACEUTICI S.P.A	IT
EYROOBI 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2566/001	044738058	FIDIA FARMACEUTICI S.P.A	IT
EYROOBI 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2566/001	044738033	FIDIA FARMACEUTICI S.P.A	IT
EYROOBI 20 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2566/001	044738019	FIDIA FARMACEUTICI S.P.A	IT
Glaucopt 20 mg/ml + 5 mg/ml eye drops, solution	DE/H/6721/001	PL 52223/0005	VISUFARMA UK LIMITED	XI
Glaucopt 20 mg/ml + 5 mg/ml eye drops, solution in single-dose-container	DE/H/6721/002	PL 52223/0004	VISUFARMA UK LIMITED	XI
Glaukostad 20 mg/ml + 5 mg/ml silmätipat, liuos	DE/H/5435/001	27729	STADA ARZNEIMITTEL AG	FI
Kiranol 20 mg/ml + 5 mg/ml oldatos szemcsepp	SE/H/1771/001	OGYI-T-21778/03	ACTAVIS GROUP PTC EHF.	HU
Kiranol 20 mg/ml + 5 mg/ml oldatos szemcsepp	SE/H/1771/001	OGYI-T-21778/02	ACTAVIS GROUP PTC EHF.	HU
Kiranol 20 mg/ml + 5 mg/ml oldatos szemcsepp	SE/H/1771/001	OGYI-T-21778/01	ACTAVIS GROUP PTC EHF.	HU
Mardozia 20 mg/ml + 5 mg/ml οφθαλμικές σταγόνες, διάλυμα	EL/H/0270/001	129223/18-12-2018	PHARMATHEN S.A.	GR
MARDOZIA PF (χωρίς συντηρητικό)® 20 mg/ml + 5 mg/ ml οφθαλμικές σταγόνες διάλυμα	DK/H/2566/001	39912/09-05-2023	PHARMATHEN S.A.	GR
Nodofree Combi, 20 mg/ml + 5 mg/ml,	PL/H/0446/001	24215	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
krople do oczu, roztwór				
Nodom Combi, 20 mg/ml + 5 mg/ml, krople do oczu, roztwór	PL/H/0616/001	21226	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Oftidorix 20 mg/5 mg/ml akių lašai (tirpalas)	PL/H/0388/001	LT/1/11/2458/001	BAUSCH + LOMB IRELAND LIMITED	LT
Oftidorix 20 mg/5 mg/ml akių lašai (tirpalas)	PL/H/0388/001	LT/1/11/2458/002	BAUSCH + LOMB IRELAND LIMITED	LT
Oftidorix 20 mg/5 mg/ml akių lašai (tirpalas)	PL/H/0388/001	LT/1/11/2458/003	BAUSCH + LOMB IRELAND LIMITED	LT
Oftidorix 20 mg/ml + 5 mg/ml acu pilieni, šķīdums	PL/H/0388/001	11-0110	BAUSCH + LOMB IRELAND LIMITED	LV
OFTIDORIX 20 mg/ml + 5 mg/ml, kapi za oko, otopina	not available	HR-H-719166270	BAUSCH + LOMB IRELAND LIMITED	HR
OFTIDORIX PF 20 mg/ml + 5 mg/ml, kapi za oko, otopina	DK/H/2477/001	HR-H-458197035	BAUSCH + LOMB IRELAND LIMITED	HR
Oftidorix, 20 mg/ml + 5 mg/ml, krople do oczu, roztwór	PL/H/0388/001	18243	BAUSCH + LOMB IRELAND LIMITED	PL
Olatalin 20 mg/ml + 5 mg/ml collirio, soluzione	CZ/H/1113/001	050187018	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	IT
Olatalin 20 mg/ml + 5 mg/ml collirio, soluzione	CZ/H/1113/001	050187020	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	IT
Olatalin 20 mg/ml + 5 mg/ml collirio, soluzione	CZ/H/1113/001	050187032	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	IT
OLATALIN 20 mg/mL + 5 mg/mL, collyre en solution	CZ/H/1113/001	34009 302 575 0 0	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FR
OLATALIN 20 mg/mL + 5 mg/mL, collyre en solution	CZ/H/1113/001	34009 302 575 1 7	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FR
OLATALIN 20 mg/mL + 5 mg/mL, collyre en solution	CZ/H/1113/001	34009 302 575 2 4	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
Olatalin 5 mg/ml + 20 mg/ml colírio, solução	CZ/H/1113/001	5844527	BRILL PHARMA UNIPESSOAL LDA.	PT
Olatalin 5 mg/ml + 20 mg/ml colírio, solução	CZ/H/1113/001	5844535	BRILL PHARMA UNIPESSOAL LDA.	PT
Olatalin 5 mg/ml + 20 mg/ml colírio, solução	CZ/H/1113/001	5844543	BRILL PHARMA UNIPESSOAL LDA.	PT
Phardinol® 20mg/ml + 5mg/ml οφθαλμικές σταγόνες, διάλυμα	not available	26908/17/23-11-2018	PHARMEX S.A.	GR
Protizol 1.5 mg/0.3 ml + 6 mg/0.3 ml colírio, solução em recipiente unidose	not available	5775911	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 1.5 mg/0.3 ml + 6 mg/0.3 ml colírio, solução em recipiente unidose	not available	18/H/0099/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 1.5 mg/0.3 ml + 6 mg/0.3 ml colテュrio, soluテアテ」o em recipiente unidose	not available	5775929	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 1.5 mg/0.3 ml + 6 mg/0.3 ml colテュrio, soluテアテ」o em recipiente unidose	not available	18/H/0099/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 5 mg/ml + 20 mg/ml, colテュrio, soluテアテ」o	not available	5851415	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 5 mg/ml + 20 mg/ml, colテュrio, soluテアテ」o	not available	5851423	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Protizol 5 mg/ml + 20 mg/ml, colテュrio, soluテアテ」o	not available	22/H/0039/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
SifiOpt 20 mg/ 5 mg/mL picături oftalmice, soluţie	RO/H/0190/001	10640/2018/01	SIFI.	RO
SifiOpt 20 mg/ 5 mg/mL	RO/H/0190/001	10640/2018/02	SIFI.	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
picături oftalmice, soluţie				
SifiOpt 20 mg/ 5 mg/mL picături oftalmice, soluţie	RO/H/0190/001	10640/2018/03	SIFI.	RO
TensocViatris 20 mg/ml + 5 mg/ml oogdruppels	AT/H/1033/001	BE389916	VIATRIS GX	BE
Tidimaz 20 mg/ml + 5 mg/ml Augentropfen, Lösung	CZ/H/0973/01	2205243.00.00	TRB CHEMEDICA AG	DE
Tidimaz 20 mg/ml + 5 mg/ml colirio en solución	CZ/H/0973/001	86765	BRILL PHARMA S.L.	ES
Tidimaz 20 mg/ml + 5 mg/ml eye drops, solution	CZ/H/1113/001	PA1391/004/001	FARMAPROJECTS S.A.U.	IE
Tidimaz 20 mg/ml + 5 mg/ml ögondroppar, lösning	CZ/H/1113/001	39542	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Tidimaz 20 mg/ml + 5 mg/ml ögondroppar, lösning	CZ/H/1113/001	62171	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Tidimaz 20 mg/ml + 5 mg/ml øyedråper, oppløsning	CZ/H/1113/001	21-14102	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Tidimaz 20 mg/ml + 5 mg/ml silmätipat, liuos	CZ/H/1113/001	39542	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Tidimaz 20 mg/ml+5 mg/ml Augentropfen, Lösung	CZ/H/1113/001	141325	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	AT
Tidimaz, øjendråber, opløsning	CZ/H/1113/001	66193	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
TIDOCOMB 2,0% +0,5% collirio, soluzione	not available	041907015	OMIKRON	IT
TIDOCOMB 2,0% +0,5% collirio, soluzione	not available	041907027	OMIKRON	IT
Tidomat 20mg/ml + 5 mg/ml eye drops, solution	not available	PL 17277/0156	PHARMATHEN S.A.	XI
Timolol + Dorzolamida	PT/H/2116/001	5366430	GENERIS FARMACÊUTICA,	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
Aurovitas 5 mg/ml + 20 mg/ml colírio, solução			S.A.	
Timolol + Dorzolamida Aurovitas 5 mg/ml + 20 mg/ml colírio, solução	PT/H/2116/001	PT/H/2116/001	GENERIS FARMACÊUTICA, S.A.	PT
Timolol + Dorzolamida Aurovitas 5 mg/ml + 20 mg/ml colírio, solução	PT/H/2116/001	PT/H/2116/001	GENERIS FARMACÊUTICA, S.A.	PT
Timolol + Dorzolamida Germed 5 mg/ml + 20 mg/ml colírio, solução	PT/H/0645/001	5559117	TOWA PHARMACEUTICAL S.A.	PT
Timolol + Dorzolamida Mylan, 5 mg/ml + 20 mg/ml, colírio	AT/H/1033/001	5366448	MYLAN, LDA	PT
Timolol + Dorzolamida Mylan, 5 mg/ml + 20 mg/ml, colírio	AT/H/1033/001	5366455	MYLAN, LDA	PT
Timolol + Dorzolamida Mylan, 5 mg/ml + 20 mg/ml, colírio	AT/H/1033/001	5366463	MYLAN, LDA	PT
Tirzopt	AT/H/1033/001	45448	VIATRIS LIMITED	DK
Torzkom 20 mg/ml + 5 mg/ml, ögondroppar, lösning i endosbehållare	SE/H/1966/001	57594	BLUMONT OFTA TRADING LIMITED	SE
VisuCOPT 20 mg/ml + 5 mg/ml Augentropfen, Lösung	DE/H/6721/001	2206316.00.00	VISUFARMA SPA	DE
VisuCOPT 20 mg/ml + 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6721/002	7000950.00.00	VISUFARMA SPA	DE
Visucopt 20 mg/ml + 5 mg/ml, colirio en solución	DE/H/6721/001	86345	VISUFARMA SPA	ES
Visucopt 20 mg/ml + 5 mg/ml, colirio en solución en envase	DE/H/6721/002	86346	VISUFARMA SPA	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
unidosis				
Vizidor Duo	DK/H/2477/001	55284	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor Duo 20 mg/5 mg/ml silmatilgad, lahus	DK/H/2477/001	965318	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor Duo 20 mg/ml + 5 mg/ ml oční kapky, roztok	DK/H/2477/001	64/457/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
VIZIDOR DUO 20 mg/ml + 5 mg/ ml οφθαλμικές σταγόνες διάλυμα	DK/H/2477/001	73069/06-07-2022	BAUSCH + LOMB IRELAND LIMITED	GR
Vizidor Duo 20 mg/ml + 5 mg/ml Augentropfen	DK/H/2477/001	137278	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor Duo 20 mg/ml + 5 mg/ml Augentropfen Lösung	DK/H/2477/001	94298.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor Duo 20 mg/ml + 5 mg/ml očné roztokové kvapky	DK/H/2477/001	64/0160/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor Duo 20 mg/ml + 5 mg/ml, eye drops solution	not available	PL 03468/0088	BAUSCH & LOMB UK LTD.	XI
Vizidor Duo, (20 mg + 5 mg)/ml, krople do oczu, roztwór	DK/H/2477/001	23677	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidorix 20 mg/ 5 mg/ ml acu pilieni, šķīdums	DK/H/2477/001	18-0064	BAUSCH + LOMB IRELAND LIMITED	LV
ZITODOR 20mg/ml + 5mg/ml collirio, soluzione	not available	041908017	VISUFARMA SPA	IT
ZITODOR 20mg/ml + 5mg/ml collirio, soluzione	not available	041908029	VISUFARMA SPA	IT
Zycopt 5 mg/ml + 20 mg/ml colírio solução	not available	5705876	DAVI II - FARMACÊUTICA, S.A.	PT
Zycopt 5 mg/ml + 20 mg/ml colírio solução	not available	5705900	DAVI II - FARMACÊUTICA, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zycopt 5 mg/ml + 20 mg/ml, colírio solução	not available	5779079	DAVI II - FARMACÊUTICA, S.A.	PT
Zycopt 5 mg/ml + 20 mg/ml, colírio solução	not available	5779103	DAVI II - FARMACÊUTICA, S.A.	PT
Амиптифри 20 mg/ml + 5 mg/ml капки за очи, разтвор	PL/H/0446/001	20170272	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	BG
ДУОКОПТ 20 mg/ml + 5 mg/ml капки за очи, разтвор	DE/H/3682/001	20140347	LABORATOIRES THEA	BG
Косопт 2 %/0,5 % капки за очи, разтвор	not available	9900199	SANTEN OY	BG
Косопт PF 20 mg/ml + 5 mg/ml, капки за очи, разтвор в еднодозова опаковка	DK/H/134/02	20 I I 063 7	SANTEN OY	BG
КОСОПТ айМулти 20 mg/ml + 5 mg/ml капки за очи, разтвор	DK/H/0134/003	MA NUMBER 20180223	SANTEN OY	BG
Офтидорикс 20 mg/ml + 5 mg/ml капки за очи, разтвор	PL/H/0388/001	20110404	BAUSCH + LOMB IRELAND LIMITED	BG
Офтидорикс PF 20 mg/ml + 5 mg/ml капки за очи, разтвор	DK/H/2477/001	20160309	BAUSCH + LOMB IRELAND LIMITED	BG