15 October 2025
EMADOC-1700519818-2654851
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): dorzolamide

EURD list No. PSUSA/00003168/202502

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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
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
DELAZO 20 mg/ml οφθαλμικές σταγόνες διάλυμα	DK/H/2604/001	50897/09-06-2017	PHARMATHEN S.A.	GR
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Delazo, øjendråber, opløsning	DK/H/2604/001	56875	PHARMATHEN S.A.	DK
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	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
Augentropfen, Lösung				
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml Augentropfen, Lösung	CZ/H/0972/001	2205242.00.00	TRB CHEMEDICA AG	DE
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE	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
			VIA FRATELLI	
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE	IT

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			VIA FRATELLI	
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153016	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153028	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml collirio, soluzione	CZ/H/1112/001	050153030	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NO
Dimaz 20 mg/ml øyedråper, oppløsning	CZ/H/1112/001	21-14101	ZAKLADY FARMACEUTYCZNE	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
			POLPHARMA S.A.	
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844477	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio solução	CZ/H/1112/001	5844501	BRILL PHARMA UNIPESSOAL LDA.	PT
Dimaz 20 mg/ml, colírio	CZ/H/1112/001	5844519	BRILL PHARMA UNIPESSOAL	PT

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solução			LDA.	
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz 20 mg/ml, ögondroppar, lösning	CZ/H/1112/001	62170	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	SE
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE	XI

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			POLPHARMA S.A.	
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz, 20 mg/ml, eye drops, solution	not available	PL 29556/0003	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	XI
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dimaz20 mg/ml colirio en solución	CZ/H/0972/001	86764	BRILL PHARMA S.L.	ES
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS	PT

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			FARMACÊUTICOS, S.A.	
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS	PT

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			FARMACÊUTICOS, S.A.	
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS	PT

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			FARMACÊUTICOS, S.A.	
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739057	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	5739065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcil 20 mg/ml Colírio, solução	not available	17/H/0034/001	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcilfree 20 mg/ml Colírio, solução	not available	5782065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcilfree 20 mg/ml Colírio, solução	not available	5782073	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcilfree 20 mg/ml Colírio, solução	not available	5782065	LABORATÓRIO EDOL - PRODUTOS	PT

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			FARMACÊUTICOS, S.A.	
Dorcilfree 20 mg/ml Colírio, solução	not available	5782073	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcilfree 20 mg/ml Colírio, solução	not available	5782065	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorcilfree 20 mg/ml Colírio, solução	not available	5782073	LABORATÓRIO EDOL - PRODUTOS FARMACÊUTICOS, S.A.	PT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	1-30561	DERMAPHARM GMBH	AT
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorlazept® 20 mg/ml Augentropfen, Lösung	AT/H/0546/001	83372.00.00	MIBE GMBH ARZNEIMITTEL	DE
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorvis 20 mg/ml kapi za oko, otopina	not available	HR-H-948759733	ZENTIVA, K.S.	HR
Dorzoclar 20 mg/ml collirio, soluzione	IT/H/0252/001/DC	039967017	OMNIVISION ITALIA S.R.L.	IT
Dorzoclar 20 mg/ml collirio, soluzione	IT/H/0252/001/DC	039967029	OMNIVISION ITALIA S.R.L.	IT
Dorzoclar 20 mg/ml collirio, soluzione	IT/H/0252/001/DC	039967031	OMNIVISION ITALIA S.R.L.	IT
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid - 1 A Pharma	DE/H/5864/001	2202750.00.00	1 A PHARMA 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
20 mg/ml Augentropfen, Lösung				
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid - 1 A Pharma 20 mg/ml Augentropfen, Lösung	DE/H/5864/001	2202750.00.00	1 A PHARMA GMBH	DE
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid ”STADA”, øjendråber, opløsning	DE/H/5445/001	45318	STADA ARZNEIMITTEL AG	DK
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid AL 20 mg/ml	DE/H/5445/001	79119.00.00	ALIUD PHARMA 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
Augentropfen				
Dorzolamid AL 20 mg/ml Augentropfen	DE/H/5445/001	79119.00.00	ALIUD PHARMA GMBH	DE
Dorzolamid Heumann 20 mg/ml Augentropfen	DE/H/1547/001	72412.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs	DE/H/5135/001	139308	MICRO LABS 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
20 mg/ml Augentropfen, Losung				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs	DE/H/5135/001	139308	MICRO LABS 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
20 mg/ml Augentropfen, Losung				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	139308	MICRO LABS GMBH	AT
Dorzolamid Micro Labs	DE/H/5135/001	139308	MICRO LABS 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
20 mg/ml Augentropfen, Losung				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs	DE/H/5135/001	99200.00.00	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
20 mg/ml Augentropfen, Losung				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs	DE/H/5135/001	99200.00.00	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
20 mg/ml Augentropfen, Losung				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Losung	DE/H/5135/001	99200.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs	DE/H/6618/001	2205614.00.00	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
20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs	DE/H/6618/001	2205614.00.00	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
20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis				
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Micro Labs 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	DE/H/6618/001	2205614.00.00	MICRO LABS GMBH	DE
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml očné roztokové	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	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
kvapky				
Dorzolamid Olikla 20 mg/ml očné roztokové kvapky	CZ/H/0761/001	64/0354/18-S	OLIKLA S.R.O.	SK
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid Olikla 20 mg/ml oční kapky, roztok	CZ/H/0761/001	64/217/17-C	OLIKLA S.R.O.	CZ
Dorzolamid STADA® 20 mg/ml Augentropfen	DE/H/5446/001	80856.00.00	STADAPHARM GMBH	DE
Dorzolamid STADA® 20 mg/ml Augentropfen	DE/H/5446/001	80856.00.00	STADAPHARM GMBH	DE
Dorzolamid STADA® 20 mg/ml Augentropfen	DE/H/5446/001	80856.00.00	STADAPHARM GMBH	DE
Dorzolamid STADA® 20 mg/ml Augentropfen	DE/H/5446/001	80856.00.00	STADAPHARM GMBH	DE
Dorzolamid Stulln 20 mg/ml Augentropfen, Lösung	DK/H/2604/001	96483.00.00	PHARMA STULLN 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
Dorzolamidă Arena 20 mg/ml picături oftalmice soluţie	not available	12262/2019/01	ARENA GROUP S.A	RO
Dorzolamida Aristo 20 mg/ml colirio en solución	PT/H/2352/001	72752	ARISTO PHARMA IBERIA, S.L.	ES
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas	PT/H/2105/001	5348842	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
20 mg/ml colírio, solução			S.A.	
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	5348842	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Aurovitas 20 mg/ml colírio, solução	PT/H/2105/001	PT/H/2105/001	GENERIS FARMACÊUTICA, S.A.	PT
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Farmalider 20 mg/ml colirio en solución	DE/H/6358/001	75.621	FARMALIDER, S.A.	ES
Dorzolamida Mylan 20 mg/ ml Colírio, Solução	NL/H/4581/001	5323928	MYLAN, LDA	PT
Dorzolamida Mylan 20 mg/ ml Colírio, Solução	NL/H/4581/001	5323944	MYLAN, LDA	PT
Dorzolamida Mylan 20 mg/ ml Colírio, Solução	NL/H/4581/001	5323936	MYLAN, LDA	PT
Dorzolamida Pharmathen	PT/H/2352/001	5324124	PHARMATHEN	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
20 mg/mL colírio, solução			INTERNATIONAL S.A.	
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen	PT/H/2352/001	5324124	PHARMATHEN	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
20 mg/mL colírio, solução			INTERNATIONAL S.A.	
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen	PT/H/2352/001	5324124	PHARMATHEN	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
20 mg/mL colírio, solução			INTERNATIONAL S.A.	
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324124	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324116	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamida Pharmathen 20 mg/mL colírio, solução	PT/H/2352/001	5324132	PHARMATHEN INTERNATIONAL S.A.	PT
Dorzolamide 20 mg/ml	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A	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
eye drops solution			SOMEX PHARMA	
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops solution	not available	PL 15764/0198	STRANDHAVEN LIMITED T/A SOMEX PHARMA	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	NL/H/3236/001	PL 20416/0559	CRESCENT PHARMA LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	NL/H/3236/001	PL 20416/0559	CRESCENT PHARMA LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	NL/H/3236/001	PL 20416/0559	CRESCENT PHARMA LIMITED	XI
Dorzolamide 20 mg/ml	PT/H/2352/001	PL 31225/0005	PHARMATHEN	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
eye drops, solution			INTERNATIONAL S.A.	
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	NL/H/4581/001	PL 04569/0911	GENERICS [UK] LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	NL/H/3236/001	PL 20416/0559	CRESCENT PHARMA LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 25258/0107	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml eye drops, solution	not available	PL 31103/0019	BLUMONT PHARMA LTD	XI
Dorzolamide 20 mg/ml eye drops, solution	PT/H/2352/001	PL 31225/0005	PHARMATHEN INTERNATIONAL S.A.	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	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 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20 mg/ml Eye Drops, Solution BP	not available	PL 15872/0025	FDC INTERNATIONAL LTD	XI
Dorzolamide 20mg/ml Eye Drops Solution	IE/H/0880/001	PL 0142/0986	ACCORD-UK LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide 20mg/ml Eye drops, solution	not available	PL35533/0146	ASPIRE PHARMA LIMITED	XI
Dorzolamide Actavis 20 mg/ml acu pilieni, šķīdums	LT/H/0139/001	11-0010	TEVA B.V	LV
Dorzolamide Actavis 20 mg/ml acu pilieni, šķīdums	LT/H/0139/001	11-0010	TEVA B.V	LV
Dorzolamide Actavis 20 mg/ml acu pilieni, šķīdums	LT/H/0139/001	11-0010	TEVA B.V	LV
Dorzolamide Actavis 20	LT/H/0139/001	LT/1/10/2335/002	TEVA B.V	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
mg/ml akiu lasai, tirpalas				
Dorzolamide Actavis 20 mg/ml akiu lasai, tirpalas	LT/H/0139/001	LT/1/10/2335/003	TEVA B.V	LT
Dorzolamide Actavis 20 mg/ml akiu lasai, tirpalas	LT/H/0139/001	LT/1/10/2335/001	TEVA B.V	LT
Dorzolamide Alvogen 20 mg/ml augndropar, lausn	IS/H/0375/001	IS/1/10/016/01	ALVOGEN EHF	IS
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo	PT/H/2105/001	040609036	AUROBINDO PHARMA	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
20 mg/ml collirio, soluzione			(ITALIA) S.R.L.	
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo 20 mg/ml collirio, soluzione	PT/H/2105/001	040609024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Dorzolamide Aurobindo	PT/H/2105/001	040609036	AUROBINDO PHARMA	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
20 mg/ml collirio, soluzione			(ITALIA) S.R.L.	
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
Dorzolamide Aurobindo 20 mg/ml oogdruppels, oplossing	PT/H/2105/001	RVG 105562	AUROBINDO PHARMA B.V.	NL
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400955043966	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20	not available	3400955043966	BIOGARAN	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
mg/ml, collyre en solution				
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400955043966	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400955043966	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400955043966	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20	not available	3400955043966	BIOGARAN	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
mg/ml, collyre en solution				
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	34009 301 190 1 3	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400955043966	BIOGARAN	FR
DORZOLAMIDE BGR 20 mg/ml, collyre en solution	not available	3400930119020	BIOGARAN	FR
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml	NL/H/5104/001	60358	BROWN & BURK IR 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
ögondroppar, lösning, endosbehållare				
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk 20 mg/ml ögondroppar, lösning, endosbehållare	NL/H/5104/001	60358	BROWN & BURK IR LIMITED	SE
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	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
conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik				
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	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
conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik				
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide Brown & Burk zonder conserveermiddel 20 mg/ml oogdruppels, oplossing in verpakking voor eenmalig gebruik	NL/H/5104/001	RVG 126313	BROWN & BURK IR LIMITED	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels, oplossing	DE/H/5445/001	RVG 105443	CENTRAFARM B.V.	NL
Dorzolamide CF 20 mg/ml, oogdruppels,	DE/H/5445/001	RVG 105443	CENTRAFARM 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
oplossing				
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 301 677 9 3	CRISTERS	FR
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 301 693 0 8	CRISTERS	FR
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 550 613 9 7	CRISTERS	FR
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 301 677 9 3	CRISTERS	FR
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 301 693 0 8	CRISTERS	FR
DORZOLAMIDE CRISTERS 20 mg/ml, collyre en solution	not available	34009 550 613 9 7	CRISTERS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	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 EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	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 EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 777 6 8	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	34009 498 778 2 9	EG LABO LABORATOIRES EUROGENERICS	FR
DORZOLAMIDE EG 20 mg/ml, collyre en solution	DE/H/5445/001	3400949877539	EG LABO LABORATOIRES EUROGENERICS	FR
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml acu pilieni, šķīdums	LV/H/0192/001	11-0011	SIA ELVIM	LV
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	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
(tirpalas)				
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	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
(tirpalas)				
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/001	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/002	SIA ELVIM	LT
Dorzolamide ELVIM 20 mg/ml akių lašai (tirpalas)	LV/H/0192/001	LT/1/10/2336/003	SIA ELVIM	LT
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE
Dorzolamide ELVIM, 20 mg/ml silmatilgad, lahus	LV/H/0192/001	724810	SIA ELVIM	EE

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 Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
Dorzolamide Farmaprojects 20 mg/ml oční kapky, roztok	CZ/H/0972/001	64/438/19-C	FARMAPROJECTS S.A.U.	CZ
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	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
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	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
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 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
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 510 1 3	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 2 1	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 3 8	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 4 5	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution	DE/H/5135/001	34009 301 892 5 2	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	34009 302 208 8 7	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL,	DE/H/6618/001	DE/H/6618/001	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
collyre en solution en récipient unidose				
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	34009 302 208 9 4	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	34009 302 208 8 7	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL, collyre en solution en récipient unidose	DE/H/6618/001	DE/H/6618/001	MICRO LABS GMBH	FR
DORZOLAMIDE MICROLABS 20 mg/mL,	DE/H/6618/001	34009 302 208 9 4	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
collyre en solution en récipient unidose				
Dorzolamide Mylan Generics 20 mg/ml collirio, soluzione	NL/H/4581/001	040083014	MYLAN S.P.A.	IT
Dorzolamide Mylan Generics 20 mg/ml collirio, soluzione.	NL/H/4581/001	040083026	MYLAN S.P.A.	IT
Dorzolamide Mylan Generics 20 mg/ml collirio, soluzione.	NL/H/4581/001	040083038	MYLAN S.P.A.	IT
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Polpharma 20 mg/ml oční kapky, roztok	CZ/H/1112/001	64/215/21-C	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	CZ
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose	not available	PLGB 25298/0363	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
container				
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	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 Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide	not available	PLGB 25298/0363	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
Preservative-Free 20mg/ml eye drops, solution in single-dose container				
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free	not available	PLGB 25298/0363	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
20mg/ml eye drops, solution in single-dose container				
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops,	not available	PLGB 25298/0363	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
solution in single-dose container				
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose	not available	PLGB 25298/0363	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
container				
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	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 Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
Dorzolamide Preservative-Free 20mg/ml eye drops, solution in single-dose container	not available	PLGB 25298/0363	BROWN & BURK UK LIMITED	XI
DORZOLAMIDE VIATRIS 2 %, collyre en solution	NL/H/4581/001	34009 496 063 6 8	VIATRIS SANTE	FR
DORZOLAMIDE VIATRIS 2 %, collyre en solution	NL/H/4581/001	34009 496 064 2 9	VIATRIS SANTE	FR
DORZOLAMIDE VIATRIS 2 %, collyre en solution	NL/H/4581/001	34009 496 061 3 9	VIATRIS SANTE	FR
Dorzolamide Viatris 20 mg/ml, oogdruppels, oplossing	NL/H/4581/001	RVG 101854	VIATRIS LIMITED	NL
Dorzolomide PharmaSwiss 20 mg/ml	DK/H/2476/001	PA23259/002/001	BAUSCH + LOMB IRELAND LIMITED	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
eye drops solution				
Dorzolomide PharmaSwiss 20 mg/ml eye drops solution	DK/H/2476/001	PA23259/002/001	BAUSCH + LOMB IRELAND LIMITED	IE
Dorzolomide PharmaSwiss 20 mg/ml eye drops solution	DK/H/2476/001	PA23259/002/001	BAUSCH + LOMB IRELAND LIMITED	IE
Dorzolomide PharmaSwiss 20 mg/ml eye drops solution	DK/H/2476/001	PA23259/002/001	BAUSCH + LOMB IRELAND LIMITED	IE
Dorzolomide PharmaSwiss 20 mg/ml eye drops solution	DK/H/2476/001	PA23259/002/001	BAUSCH + LOMB IRELAND LIMITED	IE
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189033	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189033	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189033	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml	PT/H/2352/001	040189033	DOC GENERICI 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
collirio, soluzione				
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189033	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189019	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189021	DOC GENERICI S.R.L.	IT
DORZONORM 20 mg/ml collirio, soluzione	PT/H/2352/001	040189033	DOC GENERICI S.R.L.	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio,	DK/H/1534/001	039620036	BRUSCHETTINI 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
soluzione				
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620012	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620024	BRUSCHETTINI S.R.L	IT
Dorzostill 2% collirio, soluzione	DK/H/1534/001	039620036	BRUSCHETTINI S.R.L	IT
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill 20 mg/ml, oogdruppels, oplossing	DK/H/1534/001	RVG 109493	BRUSCHETTINI S.R.L	NL
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, 20 mg/ml,	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	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				
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, 20 mg/ml, krople do oczu, roztwór	DK/H/1534/001	17041	BRUSCHETTINI S.R.L	PL
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzostill, øjendråber, opløsning	DK/H/1534/001	43562	BRUSCHETTINI S.R.L	DK
Dorzo-Vision 20 mg/ml Augentropfen	AT/H/1041/001	1-28927	OMNIVISION GMBH	AT
Dorzo-Vision 20 mg/ml Augentropfen	AT/H/1041/001	1-28927	OMNIVISION GMBH	AT
Dorzo-Vision 20 mg/ml Augentropfen, Lösung	not available	0608761	OMNIVISION GMBH	LU
Dorzo-Vision 20 mg/ml Augentropfen, Lösung	not available	0608761	OMNIVISION GMBH	LU
Dorzo-Vision sine 20 mg/ml Augentropfen, Lösung im	AT/H/0561/001	136729	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
Einzeldosisbehältnis				
Dorzo-Vision sine 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0561/001	136729	OMNIVISION GMBH	AT
Dorzo-Vision sine 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0561/001	92490.00.00	OMNIVISION GMBH	DE
Dorzo-Vision sine 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	AT/H/0561/001	92490.00.00	OMNIVISION GMBH	DE
Dorzo-Vision® 20 mg/ml Augentropfen, Lösung	AT/H/1041/001	74991.00.00	OMNIVISION GMBH	DE
Dorzo-Vision® 20 mg/ml Augentropfen, Lösung	AT/H/1041/001	74991.00.00	OMNIVISION GMBH	DE
Dorzo-Vision® 20 mg/ml Augentropfen, Lösung	not available	0608774	OMNIVISION GMBH	LU
Dorzo-Vision® 20 mg/ml Augentropfen, Lösung	not available	0608774	OMNIVISION GMBH	LU
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Dozotens 20 mg/ml očné roztokové kvapky	SK/H/0223/001	64/0928/10-S	BAUSCH + LOMB IRELAND LIMITED	SK
Eydelto 20 mg/ml eye	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA 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
drops solution				
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Eydelto 20 mg/ml eye drops solution	DK/H/2604/001	PL35533/0085	ASPIRE PHARMA LIMITED	XI
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml Augentropfen, Lösung	CZ/H/1112/001	141352	TRB CHEMEDICA (AUSTRIA) GMBH	AT
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml eye	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	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
drops, solution				
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml eye drops, solution	CZ/H/1112/001	PA1391/003/001	FARMAPROJECTS S.A.U.	IE
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml ögondroppar, lösning	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco 20 mg/ml silmätipat, liuos	CZ/H/1112/001	39509	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	FI
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	DK
Itoco, øjendråber, opløsning	CZ/H/1112/001	66190	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/001	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree 20 mg/ml akių lašai (tirpalas)	PL/H/0444/001	LT/1/17/4117/002	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LT
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Nodofree, 20 mg/ml, krople do oczu, roztwór	PL/H/0444/001	24239	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Odyral 20 mg/ml oogdruppels, oplossing in een verpakking voor eenmalig gebruik	NL/H/5895/001	RVG 131908	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	NL
Odyral 20 mg/ml oogdruppels, oplossing in een verpakking voor eenmalig gebruik	NL/H/5895/001	RVG 131908	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	NL
Odyral 20 mg/ml	NL/H/5895/001	RVG 131908	UNI-PHARMA KLEON TSETIS	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
oogdruppels, oplossing in een verpakking voor eenmalig gebruik			PHARMACEUTICAL LABORATORIES S.A.	
Odyral 20 mg/ml oogdruppels, oplossing in een verpakking voor eenmalig gebruik	NL/H/5895/001	RVG 131908	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	NL
Odyral 20 mg/ml silmatipat, liuos kerta-annospakkauksessa	NL/H/5895/001/DC	42757	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	FI
Odyral 20 mg/ml silmatipat, liuos kerta-annospakkauksessa	NL/H/5895/001/DC	42757	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	FI
Odyral 20 mg/ml silmatipat, liuos kerta-annospakkauksessa	NL/H/5895/001/DC	42757	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	FI
Odyral 20 mg/ml silmatipat, liuos kerta-annospakkauksessa	NL/H/5895/001/DC	42757	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	FI
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	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
endosbehållare				
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	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
endosbehållare				
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	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
endosbehållare				
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	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
endosbehållare				
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	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
endosbehållare				
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral 20 mg/ml, ögondroppar, lösning i endosbehållare	NL/H/5895/001	65069	ORIFARM HEALTHCARE A/S	SE
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001	024214	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001	024214	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY

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
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001	024214	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001	024214	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001/DC	(PRODUCT CODE) 3345701, (M.A NUMBER) 37433 I 01-04-2025	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001/DC	(PRODUCT CODE) 3345701, (M.A NUMBER) 37433 I 01-04-2025	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001/DC	(PRODUCT CODE) 3345701, (M.A NUMBER) 37433 I 01-04-2025	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
Odyral PF 20 mg/ml οφθαλμικές σταγόνες, διάλυμα σε περιέκτη μίας δόσης	NL/H/5895/001/DC	(PRODUCT CODE) 3345701, (M.A NUMBER) 37433 I 01-04-2025	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	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
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	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
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	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
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	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
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	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
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
Odyral, øjendråber, opløsning i enkeltdosisbeholder	NL/H/5895/001	69960	ORIFARM HEALTHCARE A/S	DK
ODYRAL® 20 mg/ml Οφθαλμικές σταγόνες, διάλυμα	NL/H/3236/001	3097301 (PRODUCT CODE)	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	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
ODYRAL® 20 mg/ml Οφθαλμικές σταγόνες, διάλυμα	NL/H/3236/001	3097301 (PRODUCT CODE)	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL,	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA 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
collyre en solution				
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 7 0	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 8 7	TRB CHEMEDICA SAS	FR
OFRIRI 20 mg/mL, collyre en solution	CZ/H/1112/001	34009 302 574 9 4	TRB CHEMEDICA SAS	FR
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor 20 mg/ml oční kapky, roztok	CZ/H/0582/001	64/1054/10-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Oftidor, 20 mg/ml, krople do oczu, roztwór	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND LIMITED	PL
Oftidor, 20 mg/ml,	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND	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			LIMITED	
Oftidor, 20 mg/ml, krople do oczu, roztwór	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND LIMITED	PL
Oftidor, 20 mg/ml, krople do oczu, roztwór	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND LIMITED	PL
Oftidor, 20 mg/ml, krople do oczu, roztwór	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND LIMITED	PL
Oftidor, 20 mg/ml, krople do oczu, roztwór	CZ/H/0582/001	17951	BAUSCH + LOMB IRELAND LIMITED	PL
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	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
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
Rezlod 20 mg/mL οφθαλμικές σταγόνες, διάλυμα	PT/H/2352/001	62965/11-07-2017	PHARMATHEN INTERNATIONAL S.A.	GR
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
SAKOR 20 mg/ml acu pilieni, šķīdums	PL/H/0444/001	17-0145	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	LV
Trusmono conserveermiddelvrij 20 mg/ml oogdruppels, oplossing	not available	RVG 123865	ROCKMED PHARMA B.V.	NL
Trusmono conserveermiddelvrij 20	not available	RVG 123865	ROCKMED 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
mg/ml oogdruppels, oplossing				
Trusmono conserveermiddelvrij 20 mg/ml oogdruppels, oplossing	not available	RVG 123865	ROCKMED PHARMA B.V.	NL
Trusmono conserveermiddelvrij 20 mg/ml oogdruppels, oplossing	not available	RVG 123865	ROCKMED PHARMA B.V.	NL
Trusmono conserveermiddelvrij 20 mg/ml oogdruppels, oplossing	not available	RVG 123865	ROCKMED PHARMA B.V.	NL
Trusmono conserveermiddelvrij 20 mg/ml oogdruppels, oplossing	not available	RVG 123865	ROCKMED PHARMA B.V.	NL
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
Trusopt	FR/H/0070/001	15910	SANTEN OY	DK
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im	FR/H/0070/002	BE285336	SANTEN OY	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
Einzeldosisbehältnis				
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 2% UNIT DOSE, 20 mg/ml Augentropfen im Einzeldosisbehältnis	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT ® 20 mg/ml, Augentropfen	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 2 % očné	not available	64/0905/96-S	SANTEN OY	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
roztokové kvapky				
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % očné roztokové kvapky	not available	64/0905/96-S	SANTEN OY	SK
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient	FR/H/0070/002	BE285336	SANTEN OY	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
unidose				
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2 % Unit Dose, 20 mg/ml collyre en solution en récipient unidose	FR/H/0070/002	2010090877	SANTEN OY	LU
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking	FR/H/0070/002	BE285336	SANTEN OY	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
voor éénmalig gebruik				
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 2% Unit Dose 20 mg/ml oogdruppels, oplossing in verpakking voor éénmalig gebruik	FR/H/0070/002	BE285336	SANTEN OY	BE
TRUSOPT 20 mg/ml acu pilieni, šķīdums	not available	98 - 0168	SANTEN OY	LV
TRUSOPT 20 mg/ml akių lašai (tirpalas)	not available	LT/1/96/1021/001	SANTEN OY	LT
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
Trusopt 20 mg/ml	FR/H/0070/001	1- 21324	SANTEN OY	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
Augentropfen Lösung				
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
Trusopt 20 mg/ml Augentropfen Lösung	FR/H/0070/001	1- 21324	SANTEN OY	AT
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colirio en solución	FR/H/0070/001	60.651	SANTEN OY	ES
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml colírio, solução	FR/H/0070/001	5770060	SANTEN OY	PT
TRUSOPT 20 mg/ml	FR/H/0070/001	AIC 031848056	SANTEN 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
collirio, soluzione				
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848056	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848056	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848056	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848056	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848056	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml	FR/H/0070/001	AIC 031848056	SANTEN 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
collirio, soluzione				
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848068	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml collirio, soluzione	FR/H/0070/001	AIC 031848070	SANTEN ITALY S.R.L.	IT
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	P A0879/004/00 1	SANTEN OY	IE
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml eye drops, solution	FR/H/0070/001	PL 16058/0014	SANTEN OY	XI
TRUSOPT 20 mg/ml kapi za oko, otopina	not available	HR-H-471662399-02	SANTEN OY	HR
TRUSOPT 20 mg/ml kapi	not available	HR-H-471662399-02	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
za oko, otopina				
TRUSOPT 20 mg/ml kapi za oko, otopina	not available	HR-H-471662399-02	SANTEN OY	HR
TRUSOPT 20 mg/ml kapi za oko, otopina	not available	HR-H-471662399-02	SANTEN OY	HR
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml ögondroppar, lösning	FR/H/0070/001	12208	SANTEN OY	SE
Trusopt 20 mg/ml oldatos szemcsepp	not available	OGYI-T-7670/01	SANTEN OY	HU
TRUSOPT 20 mg/ml	FR/H/0070/001	BE208546	SANTEN OY	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
oogdruppels, oplossing				
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
TRUSOPT 20 mg/ml oogdruppels, oplossing	FR/H/0070/001	RVG 17618	SANTEN OY	NL
Trusopt 20 mg/ml øyedråper, oppløsning med konserveringsmiddel	not available	8172	SANTEN OY	NO
Trusopt 20 mg/ml øyedråper, oppløsning uten konserveringsmiddel	not available	06-3929	SANTEN OY	NO
TRUSOPT 20 mg/ml	not available	7216/2014/02	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
picături oftalmice, soluţie				
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/02	SANTEN OY	RO
TRUSOPT 20 mg/ml picături oftalmice, soluţie	not available	7216/2014/01	SANTEN OY	RO
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml	FR/H/0070/001	11787	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
silmätipat, liuos				
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
Trusopt 20 mg/ml silmätipat, liuos	FR/H/0070/001	11787	SANTEN OY	FI
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution	FR/H/0070/001	34009 301 660 9 3	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	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
récipient unidose				
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	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
récipient unidose				
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 139 2 8	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 366 140 0 0	SANTEN OY	FR
TRUSOPT 20 mg/ml, collyre en solution en récipient unidose	FR/H/0070/002	34009 370 616 6 7	SANTEN OY	FR
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
Trusopt 20 mg/ml, ögondroppar, lösning, endosbehållare	FR/H/0070/002	22553	SANTEN OY	SE
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	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
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
TRUSOPT CONSERVEERMIDDELVRIJ, oogdruppels, oplossing in flacon voor eenmalig gebruik 20 mg/ml	FR/H/0070/002	RVG 32633	SANTEN OY	NL
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
Trusopt free bez	FR/H/0070/002	64/417/05-C	SANTEN OY	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
konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu				
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
Trusopt free bez konzervačních přísad 20 mg/ml oční kapky, roztok v jednodávkovém obalu	FR/H/0070/002	64/417/05-C	SANTEN OY	CZ
TRUSOPT PF “ Χωρίς Συντηρητικό” 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT PF “ Χωρίς Συντηρητικό” 20mg/ml	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	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
Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης				
TRUSOPT PF " Χωρίς Συντηρητικό" 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT PF " Χωρίς Συντηρητικό" 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT PF " Χωρίς Συντηρητικό" 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT PF " Χωρίς Συντηρητικό" 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT PF " Χωρίς Συντηρητικό" 20mg/ml Οφθαλμικές σταγόνες διάλυμα, σταγονόμετρα μίας δόσης	FR/H/0070/002	99181/17/23-05-2018	VIANEX S.A.	GR
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN 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
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops solution, single dose container	FR/H/0070/002	PL 16058/0013	SANTEN UK LIMITED	XI
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	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
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
TRUSOPT Preservative-Free 20 mg/ml eye drops, solution, single-dose container	FR/H/0070/002	PA0879/004/002	SANTEN OY	IE
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
Trusopt Ukonserveret	FR/H/0070/002	38051	SANTEN OY	DK
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml silmatilgad, lahus	not available	155696	SANTEN OY	EE
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT, 20 mg/ml,	not available	R/6613	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
krople do oczu, roztwór				
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT, 20 mg/ml, krople do oczu, roztwór	not available	R/6613	SANTEN OY	PL
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml Augentropfen, Lösung	FR/H/0070/001	32120.00.00	SANTEN OY	DE
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml	FR/H/0070/001	BE208546	SANTEN OY	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
collyre en solution				
TRUSOPT® 20 mg/ml collyre en solution	FR/H/0070/001	BE208546	SANTEN OY	BE
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	FR/H/0070/001	53809/03-05-2019	VIANEX S.A.	GR
TRUSOPT® 20 mg/ml, oční kapky, roztok	not available	64/567/97-C	SANTEN OY	CZ
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
TRUSOPT-S® 20 mg/ml	FR/H/0070/002	63083.00.00	SANTEN OY	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
Augentropfen, Lösung im Einzeldosisbehältnis				
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
TRUSOPT-S® 20 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	FR/H/0070/002	63083.00.00	SANTEN OY	DE
UNIDROPS 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	NL/H/3236/001	022708	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY
UNIDROPS 20 mg/ml οφθαλμικές σταγόνες, διάλυμα	NL/H/3236/001	022708	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	CY
Vizidor	DK/H/2476/001	55282	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor	DK/H/2476/001	55282	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor	DK/H/2476/001	55282	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor	DK/H/2476/001	55282	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor	DK/H/2476/001	55282	BAUSCH + LOMB IRELAND LIMITED	DK
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND	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
Augentropfen, Lösung			LIMITED	
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	94297.00.00	BAUSCH + LOMB IRELAND LIMITED	DE
Vizidor 20 mg/ml eye drops solution	DK/H/2476/001	PL 03468/0087	BAUSCH & LOMB UK LTD.	XI
Vizidor 20 mg/ml eye drops solution	DK/H/2476/001	PL 03468/0087	BAUSCH & LOMB UK LTD.	XI
Vizidor 20 mg/ml eye drops solution	DK/H/2476/001	PL 03468/0087	BAUSCH & LOMB UK LTD.	XI
Vizidor 20 mg/ml eye drops solution	DK/H/2476/001	PL 03468/0087	BAUSCH & LOMB UK LTD.	XI
Vizidor 20 mg/ml eye drops solution	DK/H/2476/001	PL 03468/0087	BAUSCH & LOMB UK LTD.	XI
Vizidor 20 mg/ml kapi za oko, otopina	DK/H/2476/001	HR-H-162487466	BAUSCH + LOMB IRELAND LIMITED	HR
Vizidor 20 mg/ml kapi za oko, otopina	DK/H/2476/001	HR-H-162487466	BAUSCH + LOMB IRELAND LIMITED	HR
Vizidor 20 mg/ml kapi za	DK/H/2476/001	HR-H-162487466	BAUSCH + LOMB IRELAND	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
oko, otopina			LIMITED	
Vizidor 20 mg/ml kapi za oko, otopina	DK/H/2476/001	HR-H-162487466	BAUSCH + LOMB IRELAND LIMITED	HR
Vizidor 20 mg/ml kapi za oko, otopina	DK/H/2476/001	HR-H-162487466	BAUSCH + LOMB IRELAND LIMITED	HR
Vizidor 20 mg/ml očné roztokové kvapky	DK/H/2476/001	64/0161/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor 20 mg/ml očné roztokové kvapky	DK/H/2476/001	64/0161/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor 20 mg/ml očné roztokové kvapky	DK/H/2476/001	64/0161/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor 20 mg/ml očné roztokové kvapky	DK/H/2476/001	64/0161/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor 20 mg/ml očné roztokové kvapky	DK/H/2476/001	64/0161/18-S	BAUSCH + LOMB IRELAND LIMITED	SK
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND	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
kapky, roztok			LIMITED	
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml oční kapky, roztok	DK/H/2476/001	64/456/17-C	BAUSCH + LOMB IRELAND LIMITED	CZ
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND	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
ögondroppar, lösning			LIMITED	
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml ögondroppar, lösning	DK/H/2476/001	65350	BAUSCH + LOMB IRELAND LIMITED	SE
Vizidor 20 mg/ml oldatos szemcsepp	DK/H/2476/001	OGYI-T-24425/01	BAUSCH + LOMB IRELAND LIMITED	HU
Vizidor 20 mg/ml oldatos	DK/H/2476/001	OGYI-T-24425/01	BAUSCH + LOMB IRELAND	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
szemcsepp			LIMITED	
Vizidor 20 mg/ml oldatos szemcsepp	DK/H/2476/001	OGYI-T-24425/01	BAUSCH + LOMB IRELAND LIMITED	HU
Vizidor 20 mg/ml oldatos szemcsepp	DK/H/2476/001	OGYI-T-24425/01	BAUSCH + LOMB IRELAND LIMITED	HU
Vizidor 20 mg/ml oldatos szemcsepp	DK/H/2476/001	OGYI-T-24425/01	BAUSCH + LOMB IRELAND LIMITED	HU
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml øyedråper, oppløsning	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND LIMITED	NO
Vizidor 20 mg/ml	DK/H/2476/001	23-15789	BAUSCH + LOMB IRELAND	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			LIMITED	
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmatilgad, lahus	DK/H/2476/001	965218	BAUSCH + LOMB IRELAND LIMITED	EE
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND	FI

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silmätipat, liuos			LIMITED	
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor 20 mg/ml silmätipat, liuos	DK/H/2476/001	42987	BAUSCH + LOMB IRELAND LIMITED	FI
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml,	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND	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			LIMITED	
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor, 20 mg/ml, krople do oczu, roztwór	DK/H/2476/001	23676	BAUSCH + LOMB IRELAND LIMITED	PL
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND	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
Augentropfen, Lösung			LIMITED	
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Vizidor® 20 mg/ml Augentropfen, Lösung	DK/H/2476/001	137277	BAUSCH + LOMB IRELAND LIMITED	AT
Zoliop 20 mg/ml collirio, soluzione	IT/H/0658/001	042570010	FARMIGEA SPA	IT
Zoliop 20 mg/ml collirio, soluzione	IT/H/0658/001	042570022	FARMIGEA SPA	IT
Zoliop 20 mg/ml collirio, soluzione	IT/H/0658/001	042570034	FARMIGEA SPA	IT