09 June 2017
EMA/36849/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): brimonidine / timolol

Procedure No.: PSUSA/00000431/201609

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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
Brimonidin/Timolol AL 2 mg/ml + 5 mg/ml Augentropfen, Lösung	DK/H/2372/001	95124.00.00	ALIUD PHARMA GMBH	DE
Brimonidin/Timolol Stada 2 mg/ml + 5 mg/ml Augentropfen	DK/H/2371/001	136430	STADA ARZNEIMITTEL AG	AT
BRIMONIDINA E TIMOLOLO EG 2 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2372/001	044245037	EG SPA	IT
BRIMONIDINA E TIMOLOLO EG 2 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2372/001	044245025	EG SPA	IT
BRIMONIDINA E TIMOLOLO EG 2 mg/ml + 5 mg/ml collirio, soluzione	DK/H/2372/001	044245013	EG SPA	IT
Brimonidina/Timolol ratiopharm 2 mg/ml + 5 mg/ml colirio en solución	DK-H-2129-001-DC	76415	RATIOPHARM ESPANA SA	ES
Brimonidina/Timolol STADA 2 mg/ml + 5 mg/ml colirio en solución	DK/H/2372/001	80.157	LABORATORIO STADA, S.L.	ES
Brimonidina/Timolol Teva 2 mg/ml + 5 mg/ml colirio en solución	DK/H/2114/001	76414	TEVA PHARMA S.L.U	ES
Brimonidine Tartrate /Timolol 2 mg/ml + 5 mg/ml eye drops, solution	DK/H/2526/001	PL04569/1631	GENERICS [UK] LIMITED	UK
Brimonidine Tartrate/Timolol 2 mg/ml + 5 mg/ml Eye Drops, Solution	DK/H/2114/001	PL 00289/2019	TEVA UK LIMITED	UK
Brimonidine/Timolol "Stada Arzneimittel AG", øjendråber, opløsning	DK/H/2371/001	53529	STADA ARZNEIMITTEL AG	DK
Brimonidine/Timolol "Stada", øjendråber, opløsning	DK/H/2372/001	53530	STADA ARZNEIMITTEL AG	DK

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Brimonidine/Timolol ratiopharm 2 mg/ml + 5 mg/ml, oogdruppels, oplossing	DK/H/2129/001	RVG 119617	RATIOPHARM NEDERLAND B.V	NL
Brimonidine/Timolol Teva 2 mg/ml + 5 mg/ml, oogdruppels, oplossing	DK/H/2114/001	RVG 119619	TEVA NEDERLAND B.V.	NL
Brimonidintartrat/Timolol "DOC Generici"	DK/H/2524/001	55948	DOC GENERICI S.R.L.	IT
Brimonidintartrat/Timolol Teva	DK/H/2114/001	49457	TEVA DENMARK A/S	DK
Brimoplus, 0,2% + 0,5%, krople do oczu, roztwór	DK/H/2114/001	20801	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Brimotimo, øjendråber, opløsning	DK/H/2129/001	49458	TEVA DENMARK A/S	DK
Combigan 2 mg/5 mg/ml acu pilieni, šķīdums	UK/H/0807/001	13-0276	ALLERGAN PHARMACEUTICALS IRELAND	LV
Combigan 2 mg/5 mg/ml akių lašai (tirpalas)	UK/H/0807/001	LT/1/13/3463/001	ALLERGAN PHARMACEUTICALS IRELAND	LT
Combigan 2 mg/5 mg/ml akių lašai (tirpalas)	UK/H/0807/001	LT/1/13/3463/002	ALLERGAN PHARMACEUTICALS IRELAND	LT
Combigan 2 mg/5 mg/ml silmatilgad, lahus	UK/H/0807/001	830513	ALLERGAN PHARMACEUTICALS IRELAND	EE
Combigan 2 mg/ml + 5 mg/ml Augentropfen	UK/H/0807/001	Z. NR. 1-26629	ALLERGAN PHARMACEUTICALS IRELAND	AT
Combigan 2 mg/ml + 5 mg/ml Augentropfen	UK/H/0807/001	63505.00.00	ALLERGAN PHARMACEUTICALS IRELAND	DE
COMBIGAN 2 mg/ml + 5 mg/ml Augentropfen, Lösung	UK/H/0807/001	BE280052	ALLERGAN PHARMACEUTICALS IRELAND	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
Combigan 2 mg/ml + 5 mg/ml augndropar, lausn	UK/H/0807/001	IS/1/05/053/01	ALLERGAN PHARMACEUTICALS IRELAND	IS
Combigan 2 mg/ml + 5 mg/ml colirio en solución.	UK/H/0807/001	67438	ALLERGAN PHARMACEUTICALS IRELAND	ES
Combigan 2 mg/ml + 5 mg/ml colírio, solução	UK/H/0807/001	5742788	ALLERGAN PHARMACEUTICALS IRELAND	PT
Combigan 2 mg/ml + 5 mg/ml collirio, soluzione	UK/H/0807/001	037083019	ALLERGAN PHARMACEUTICALS IRELAND	IT
Combigan 2 mg/ml + 5 mg/ml collirio, soluzione	UK/H/0807/001	037083021	ALLERGAN PHARMACEUTICALS IRELAND	IT
Combigan 2 mg/ml + 5 mg/ml eye drops, solution	UK/H/0807/001	PA 148/64/1	ALLERGAN PHARMACEUTICALS IRELAND	IE
Combigan 2 mg/ml + 5 mg/ml eye drops, solution	UK/H/0807/001	MA102/00701	ALLERGAN PHARMACEUTICALS IRELAND	MT
Combigan 2 mg/ml + 5 mg/ml eye drops, solution	UK/H/0807/001	PL 41443/0006	ALLERGAN PHARMACEUTICALS IRELAND	UK
COMBIGAN 2 mg/ml + 5 mg/ml kapi za oko, otopina	not available	UP/I-530-09/13-02/141	ALLERGAN PHARMACEUTICALS IRELAND	HR
Combigan 2 mg/ml + 5 mg/ml kapljice za oko, raztopina	UK/H/0807/001	5363-I-1310/10	ALLERGAN PHARMACEUTICALS IRELAND	SI
COMBIGAN 2 mg/ml + 5 mg/ml očná roztoková instilácia	UK/H/0807/001	64/0131/06-S	ALLERGAN PHARMACEUTICALS IRELAND	SK
Combigan 2 mg/ml + 5 mg/ml Ocní kapky, roztok	UK/H/0807/001	64/486/05-C	ALLERGAN PHARMACEUTICALS IRELAND	CZ

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Combigan 2 mg/ml + 5 mg/ml ögondroppar, lösning	UK/H/0807/001	22652	ALLERGAN PHARMACEUTICALS IRELAND	SE
Combigan 2 mg/ml + 5 mg/ml oogdruppels, oplossing	UK/H/0807/001	BE280052	ALLERGAN PHARMACEUTICALS IRELAND	LU
Combigan 2 mg/ml + 5 mg/ml øyedråper, oppløsning.	UK/H/0807/001	05-3535	ALLERGAN PHARMACEUTICALS IRELAND	NO
Combigan 2 mg/ml + 5 mg/ml picături oftalmice, soluție	UK/H/0807/001	6001/2013/01	ALLERGAN PHARMACEUTICALS IRELAND	RO
Combigan 2 mg/ml + 5 mg/ml picături oftalmice, soluție	UK/H/0807/001	6001/2013/02	ALLERGAN PHARMACEUTICALS IRELAND	RO
Combigan 2 mg/ml + 5 mg/ml silmätipat, liuos	UK/H/0807/001	21341	ALLERGAN PHARMACEUTICALS IRELAND	FI
Combigan 2 mg/ml + 5 mg/ml οφθαλμικές σταγόνες, διάλυμα	UK/H/0807/001	M.L.:022229	ALLERGAN PHARMACEUTICALS IRELAND	CY
Combigan 2 mg/ml + 5 mg/ml οφθαλμικές σταγόνες, διάλυμα	UK/H/0807/001	41726/20-05-2016	ALLERGAN PHARMACEUTICALS IRELAND	GR
Combigan 2 mg/ml + 5 mg/ml, collyre en solution	UK/H/0807/001	BE280052	ALLERGAN PHARMACEUTICALS IRELAND	BE
COMBIGAN 2 mg/ml + 5 mg/ml, collyre en solution	UK/H/0807/001	34009 371 722 4 0	ALLERGAN PHARMACEUTICALS IRELAND	FR
COMBIGAN 2 mg/ml + 5 mg/ml, collyre en solution	UK/H/0807/001	34009 375 410 7 7	ALLERGAN PHARMACEUTICALS IRELAND	FR
Combigan 2 mg/ml + 5 mg/ml, oogdruppels, oplossing	UK/H/0807/001	BE280052	ALLERGAN PHARMACEUTICALS IRELAND	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
Combigan 2 mg/ml + 5 mg/ml, oogdruppels, oplossing	UK/H/0807/001	RVG 32744	ALLERGAN PHARMACEUTICALS IRELAND	NL
Combigan szemcsepp 2 mg/ml + 5 mg/ml oldatos szemcsepp	UK/H/0807/001	OGYI-T-20114/02	ALLERGAN PHARMACEUTICALS IRELAND	HU
Combigan szemcsepp 2 mg/ml + 5 mg/ml oldatos szemcsepp	UK/H/0807/001	OGYI-T-20114/01	ALLERGAN PHARMACEUTICALS IRELAND	HU
Combigan, 2 mg/ml + 5 mg/ml øjendråber, opløsning	UK/H/0807/001	38127	ALLERGAN PHARMACEUTICALS IRELAND	DK
Combigan, 2 mg/ml + 5 mg/ml, krople do oczu, roztwór	UK/H/0807/001	12076	ALLERGAN PHARMACEUTICALS IRELAND	PL
COMBIGAN® 2 mg/ml + 5 mg/ml ögondroppar, lösning	UK/H/0807/001	21341	ALLERGAN PHARMACEUTICALS IRELAND	FI
Zolbrinex	DK/H/2525/001	55949	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	DK
Zolbrinex 2 mg/5 mg v 1 ml kapljice za oko, raztopina	DK/H/2525/001	H/16/02238/001	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Zolbrinex 2 mg/5 mg v 1 ml kapljice za oko, raztopina	DK/H/2525/001	H/16/02238/002	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Zolbrinex 2 mg/5 mg v 1 ml kapljice za oko, raztopina	DK/H/2525/001	H/16/02238/003	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SI
Комбиган 2 mg/ml + 5 mg/ml капки за очи, разтвор	UK/H/0807/001	20140073	ALLERGAN PHARMACEUTICALS IRELAND	BG