12 July 2018
EMA/490585/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: cetirizine

Procedure no.: PSUSA/00000628/201711

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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
VIRLIX 10 MG, COMPRIME PELLICULE SECABLE	not available	330 235-1	SANOFI-AVENTIS FRANCE	FR
VIRLIX 10 MG/ML, SOLUTION BUVABLE EN GOUTTES	not available	341 772-3	SANOFI-AVENTIS FRANCE	FR
VIRLIX 10 MG, COMPRIME PELLICULE SECABLE	not available	330 234-5	SANOFI-AVENTIS FRANCE	FR
VIRLIX 10 MG/ML, SOLUTION BUVABLE EN GOUTTES	not available	341 774-6	SANOFI-AVENTIS FRANCE	FR
Alerlisin 10 mg comprimidos recubiertos con película	not available	58.499	LABORATORIOS MENARINI, S.A.	ES
Alerlisin 1 mg/ml solución oral	not available	62.787	LABORATORIOS MENARINI, S.A.	ES
Alerlisin 10 mg/ml gotas orales en solución	not available	62.788	LABORATORIOS MENARINI, S.A.	ES
FORMISTIN 10 mg/ml gocce orali, soluzione	not available	027329022	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
FORMISTIN 10 mg compresse rivestite con film	not available	027329034	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
FORMISTIN 10 mg compresse rivestite con film	not available	027329010	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
GENERIT 10 mg/ml gocce orali, soluzione	not available	038628020	GENETIC SPA	IT
RITECAM 10 mg/ml gocce orali, soluzione	not available	038629022	GENETIC SPA	IT
RAINGEN 10 mg/ml gocce orali, soluzione	not available	038630024	GENETIC SPA	IT
Piriteze Allergy 1mg/ml Syrup	not available	PL 44673/0095	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Piriteze Allergy Tablets	not available	PL 44673/0097	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	UK
Piriteze Allergy Tablets	not available	PL 44673/0098	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	UK
Piriteze Allergy Relief 1mg/ml Syrup	not available	PL 44673/0096	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	UK
Zyrlex 1 mg/ml oral lösning	IE/H/0209/003	12678	MACURE PHARMA APS	SE
Zyrlex 10 mg/ml orala droppar, lösning	IE/H/0209/002	11673	MACURE PHARMA APS	SE
Zyrlex 10 mg filmdragerade tabletter	IE/H/0209/001	11209	MACURE PHARMA APS	SE
Zyrtec 1 mg/ml drank	IE/H/0209/003	BE154673	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg filmomhulde tabletten	IE/H/0209/001	BE135877	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg/ml druppels voor oraal gebruik, oplossing	IE/H/0209/002	BE149746	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/001	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/002	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/003	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/004	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/005	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/006	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko	IE/H/0209/001	H/01/01728/007	UCB PHARMA S.A. (ANDERL	SI

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obložene tablete			BE)	
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/008	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/009	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/010	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/011	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/012	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/013	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/014	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/015	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/016	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmsko obložene tablete	IE/H/0209/001	H/01/01728/017	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 1 mg/ml peroralna raztopina	IE/H/0209/003	H/01/01728/018	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 1 mg/ml peroralna raztopina	IE/H/0209/003	H/01/01728/019	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 1 mg/ml peroralna raztopina	IE/H/0209/003	H/01/01728/020	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 1 mg/ml peroralna raztopina	IE/H/0209/003	H/01/01728/021	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 1 mg/ml peroralna raztopina	IE/H/0209/003	H/01/01728/022	UCB PHARMA S.A. (ANDERL BE)	SI
Zyrtec 10 mg filmtabletta	IE/H/0209/001	OGYI-T-1599/03	UCB MAGYARORSZÁG KFT (BUDAPEST HU)	HU
Zyrtec 10 mg	IE/H/0209/001	OGYI-T-1599/02	UCB MAGYARORSZÁG KFT	HU

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filmtabletta			(BUDAPEST HU)	
Zyrtec 10 mg/ml belsőleges oldatos cseppek	IE/H/0209/002	OGYI-T-4491/01	UCB MAGYARORSZÁG KFT (BUDAPEST HU)	HU
Zyrtec 10 mg Filmtabletten	IE/H/0209/001	1-19030	UCB PHARMA GMBH (VIENNA AT)	AT
Зиртек 10 mg филмирани таблетки	IE/H/0209/001	20020795	UCB PHARMA GMBH (MONHEIM DE)	BG
Zyrtec, 10 mg õhukese polümeerikattega tabletid	IE/H/0209/001	162097	UCB PHARMA OY FINLAND (ESPOO FI)	EE
Zyrtec 10 mg tabletti, kalvopäällysteinen	IE/H/0209/001	10385	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec 10 mg, comprimé pelliculé sécable	IE/H/0209/001	34009 329 944 2 7	UCB PHARMA S.A. (COLOMBES FR)	FR
Zyrtec 10 mg, comprimé pelliculé sécable	IE/H/0209/001	34009 329 945 9 5	UCB PHARMA S.A. (COLOMBES FR)	FR
Zyrtec 10 mg, comprimé pelliculé sécable	IE/H/0209/001	34009 336 135 9 4	UCB PHARMA S.A. (COLOMBES FR)	FR
Zyrtec 10 mg, comprimé pelliculé sécable	IE/H/0209/001	34009 339 615 1 0	UCB PHARMA S.A. (COLOMBES FR)	FR
Zirtek 10 mg film-coated tablets	IE/H/0209/001	PA0891/008/002	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtec 10 mg compresse rivestite con film	IE/H/0209/001	026894016	UCB PHARMA SPA (MILAN IT)	IT
Zirtec 10 mg compresse rivestite con film	IE/H/0209/001	026894042	UCB PHARMA SPA (MILAN IT)	IT
Zirtec 10 mg compresse rivestite con film	IE/H/0209/001	026894067	UCB PHARMA SPA (MILAN IT)	IT
Zyrtec 10 mg apvalkotās tabletes	IE/H/0209/001	98-0688	UCB PHARMA OY FINLAND (ESPOO FI)	LV
Zyrtec 10 mg plėvele dengtos tabletės	IE/H/0209/001	LT/1/97/2859/001	UCB PHARMA OY FINLAND (ESPOO FI)	LT
Zyrtec 10 mg plėvele	IE/H/0209/001	LT/1/97/2859/002	UCB PHARMA OY FINLAND	LT

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dengtos tabletės			(ESPOO FI)	
Zyrtec 10 mg plėvele dengtos tabletės	IE/H/0209/001	LT/1/97/2859/003	UCB PHARMA OY FINLAND (ESPOO FI)	LT
Zyrtec 10 mg potahované tablety	IE/H/0209/001	24/024/92-S/C	UCB S.R.O. (PRAGUE CZ)	CZ
Zyrtec® 10 mg Filmtabletten	IE/H/0209/001	14836.00.00	UCB PHARMA GMBH (MONHEIM DE)	DE
Ζιρτέκ 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0209/001	1075/08-01-2015	UCB A.E. (ATHENS GR)	GR
Zyrtec 10 mg filmomhulde tabletten	IE/H/0209/001	RVG 13010	UCB PHARMA B.V. (BREDA NL)	NL
Zyrtec 10 mg/ml perorální kapky, roztok	IE/H/0209/002	24/1030/92-S/C	UCB S.R.O. (PRAGUE CZ)	CZ
Zyrtec, 10 mg, tabletki powlekane	IE/H/0209/001	R/1846	VEDIM SP. Z O.O. (WARSAW PL)	PL
Zyrtec 1 mg/ml mikstur, oppløsning	IE/H/0209/003	7901	UCB NORDIC A/S (COPENHAGEN DK)	NO
Zyrtec 10 mg tabletter, filmdrasjerte	IE/H/0209/001	7594	UCB NORDIC A/S (COPENHAGEN DK)	NO
Zyrtec 1 mg/ml šķīdums iekšķīgai lietošanai	IE/H/0209/003	98-0791	UCB PHARMA OY FINLAND (ESPOO FI)	LV
Zyrtec 10 mg/ml pilieni iekšķīgai lietošanai, šķīdums	IE/H/0209/002	98-0730	UCB PHARMA OY FINLAND (ESPOO FI)	LV
Zyrtec 1 mg/ml solution buvable	IE/H/0209/003	2011101305	UCB PHARMA S.A. (ANDERL BE)	LU
Zyrtec 10 mg comprimés pelliculés	IE/H/0209/001	2011101303	UCB PHARMA S.A. (ANDERL BE)	LU
Zyrtec 10 mg/ml geriamieji lašai (tirpalas)	IE/H/0209/002	LT/1/97/2859/005	UCB PHARMA OY FINLAND (ESPOO FI)	LT
Zirtek 10 mg film-coated tablets	IE/H/0209/001	PL 00039/0542	UCB PHARMA LTD (SLOUGH GB)	UK
Zyrtec 2,5 mg/2,5 ml,	IE/H/0209/003	34009 333 019 8 9	UCB PHARMA S.A.	FR

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solution buvable en flacon			(COLOMBES FR)	
Zyrtec 10 mg/ml tipat, liuos	IE/H/0209/002	11321	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec, 10 mg/ml suukaudsed tilgad, lahus	IE/H/0209/002	162197	UCB PHARMA OY FINLAND (ESPOO FI)	EE
Зиртек 10 mg/ml перорални капки, разтвор	IE/H/0209/002	20020794	UCB PHARMA GMBH (MONHEIM DE)	BG
Zyrtec 10 mg comprimés pelliculés	IE/H/0209/001	BE135877	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg/ml Tropfen	IE/H/0209/002	1-20142	UCB PHARMA GMBH (VIENNA AT)	AT
Zyrtec, filmovertrukne tabletter	IE/H/0209/001	13082	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec, orale dråber, opløsning	IE/H/0209/002	13651	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec, oral opløsning	IE/H/0209/003	14784	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/02	UCB PHARMA GMBH (MONHEIM DE)	RO
Zyrtec, 10 mg/ml, krople doustne, roztwór	IE/H/0209/002	R/1847	VEDIM SP. Z O.O. (WARSAW PL)	PL
Zyrtec, 1 mg/ml, roztwór doustny	IE/H/0209/003	7815	VEDIM SP. Z O.O. (WARSAW PL)	PL
Zyrtec 10 mg/ml dråper, oppløsning	IE/H/0209/002	8173	UCB NORDIC A/S (COPENHAGEN DK)	NO
Zirtec 10 mg/ml gocce orali, soluzione	IE/H/0209/002	026894028	UCB PHARMA SPA (MILAN IT)	IT
Zirtek 10 mg/ml Oral Drops, Solution	IE/H/0209/002	PA0891/008/004	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zyrtec 1 mg/ml solución oral	IE/H/0209/003	60.279	UCB PHARMA S.A. (MADRID ES)	ES
Zyrtec, 1 mg/ml	IE/H/0209/003	162297	UCB PHARMA OY FINLAND	EE

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suukaudne lahus			(ESPOO FI)	
Zyrtec 10 mg/ml solution buvable en gouttes	IE/H/0209/002	BE149746	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 1 mg/ml orale Lösung	IE/H/0209/003	1-20143	UCB PHARMA GMBH (VIENNA AT)	AT
Zyrtec 1 mg/ml solution buvable	IE/H/0209/003	BE154673	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg επικαλυμμένα με λεπτό υμένιο δισκία.	IE/H/0209/001	13066	UCB PHARMA S.A. (ANDERL BE)	CY
Zyrtec 1 mg/ml πόσιμο διάλυμα.	IE/H/0209/003	16365	UCB PHARMA S.A. (ANDERL BE)	CY
Zyrtec 10 mg/ml gotas orales en solución	IE/H/0209/002	60.280	UCB PHARMA S.A. (MADRID ES)	ES
Zyrtec 1 mg/ml oraaliliuos	IE/H/0209/003	11322	UCB PHARMA OY FINLAND (ESPOO FI)	FI
ZYRTEC 2,5 mg/2,5 ml, solution buvable en flacon	IE/H/0209/003	34009 332 924 9 2	UCB PHARMA S.A. (COLOMBES FR)	FR
Zirtek Allergy Solution, 1 mg/ml oral solution	IE/H/0209/003	PL 00039/0540	UCB PHARMA LTD (SLOUGH GB)	UK
Zirtek 1 mg/ml oral solution	IE/H/0209/003	PA0891/008/003	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtec 1 mg/ml soluzione orale	IE/H/0209/003	026894030	UCB PHARMA SPA (MILAN IT)	IT
Zirtec 1 mg/ml soluzione orale	IE/H/0209/003	026894093	UCB PHARMA SPA (MILAN IT)	IT
Zyrtec 1 mg/ml geriamasis tirpalas	IE/H/0209/003	LT/1/97/2859/004	UCB PHARMA OY FINLAND (ESPOO FI)	LT
Zyrtec 1 mg/ml drank	IE/H/0209/003	RVG 14635	UCB PHARMA B.V. (BREDA NL)	NL
Zyrtec 1 mg/ml Solução oral	IE/H/0209/003	2051795	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT

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Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/01	UCB PHARMA GMBH (MONHEIM DE)	RO
Zyrtec 10 mg/ml perorálne roztokové kvapky	IE/H/0209/002	24/1030/92-S	UCB S.R.O. (PRAGUE CZ)	SK
ZYRTEC 10 mg/ml, solution buvable en gouttes	IE/H/0209/002	34009 341 770 0 2	UCB PHARMA S.A. (COLOMBES FR)	FR
Zirtec 10 mg/ml gocce orali, soluzione	IE/H/0209/002	026894081	UCB PHARMA SPA (MILAN IT)	IT
Zyrtec 10 mg comprimidos revestidos por película	IE/H/0209/001	9717900	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Zyrtec 10 mg comprimidos recubiertos con película	IE/H/0209/001	58.481	UCB PHARMA S.A. (MADRID ES)	ES
Zyrtec 10 mg/ml solution buvable en gouttes	IE/H/0209/002	2011101304	UCB PHARMA S.A. (ANDERL BE)	LU
Zyrtec 10 mg filmom obalené tablety	IE/H/0209/001	24/0024/92-S	UCB S.R.O. (PRAGUE CZ)	SK
Zyrtec 10 mg Filmtabletten	IE/H/0209/001	BE135877	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 1 mg/ml Lösung zum Einnehmen	IE/H/0209/003	BE154673	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg/ml Tropfen zum Einnehmen, Lösung	IE/H/0209/002	BE149746	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg filmdragerade tabletter	IE/H/0209/001	10385	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec 10 mg/ml orala droppar, lösning	IE/H/0209/002	11321	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec 1 mg/ml oral lösning	IE/H/0209/003	11322	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Ζιρτέκ 10 mg/ml πόσιμες σταγόνες, διάλυμα	IE/H/0209/002	1076/08-01-2015	UCB A.E. (ATHENS GR)	GR
Zyrtec 10 mg apvalkotās	IE/H/0209/001	98-0731	UCB PHARMA OY FINLAND	LV

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tabletes			(ESPOO FI)	
Zyrtec 10 mg film-coated tablets	IE/H/0209/001	MA030/00901	UCB PHARMA S.A. (ANDERL BE)	MT
Zyrtec 1 mg/ml oral solution	IE/H/0209/003	MA030/00902	UCB PHARMA S.A. (ANDERL BE)	MT
Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/03	UCB PHARMA GMBH (MONHEIM DE)	RO
Cetirizine diHCl ADOH 10 mg, zuigtabletten	not available	RVG 30369	ADOH B.V.	NL
ZYRTECSET 10 mg, comprimé pelliculé sécable	not available	34009 364 616 8 0	UCB PHARMA S.A. (COLOMBES FR)	FR
Ziralton Allergy Relief for Children 5 mg/5 ml Oral Solution	not available	04917/0140	PINEWOOD LABORATORIES LIMITED	UK
CETIRIZINE EG 10 mg, comprimé à sucer	not available	NL30927	EG LABO LABORATOIRES EUROGENERICS	FR
DRILL ALLERGIE CETIRIZINE 10 mg, comprimé à sucer	not available	34009 381 535 2 1	PIERRE FABRE MEDICAMENT	FR
Wockhardt Allergy & Hayfever Relief 10mg Film-coated Tablets	not available	29831/0681	WOCKHARDT UK LTD	UK
Zyrtec UCB, 10 mg, tabletki powlekane	not available	11212	VEDIM SP. Z O.O. (WARSAW PL)	PL
Cetirizine 1mg/ml Oral Solution	not available	PL 04416/0410	SANDOZ LTD	UK
Cetirizine Dihydrochloride l0mg Film-coated Tablets	not available	PL 36687/0242	TORRENT PHARMA LTD.	UK
Benadryl Allergy Oral Syrup	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK
Benadryl for Children Allergy Solution	not available	PL 15513/0124	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK

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Children's 1mg/ml Oral Solution/ Benadryl Allergy Oral Syrup				
Benadryl Allergy Children's 1mg/ml Oral Solution/ Benadryl Allergy Oral Syrup	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy Oral Syrup	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy Oral Syrup	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy Children's 1mg/ml Oral Solution/ Benadryl Allergy Oral Syrup	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy Children's 6+ 1mg/ml Oral Solution	not available	PL 15513/0124	MCNEIL PRODUCTS LIMITED	UK
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PL 00039/0561	UCB PHARMA LTD (SLOUGH GB)	UK
Zirtek Allergy Relief for Children 1 mg/mL oral solution	not available	PL 00039/0541	UCB PHARMA LTD (SLOUGH GB)	UK
Cetirizine Dihydrochloride l0mg Film-coated Tablets	not available	PL 36687/0241	TORRENT PHARMA LTD.	UK
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PA0891/008/005	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
POLLENSHIELD HAYFEVER RELIEF Cetirizine hydrochloride 10mg Film-coated Tablets	not available	PL 0142/0606	ACTAVIS UK LTD.	UK
GENERIT 10 mg/ml gocce orali, soluzione	not available	038628020	GENETIC SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RITECAM 10 mg/ml gocce orali, soluzione	not available	038629022	GENETIC SPA	IT
RAINGEN 10 mg/ml gocce orali, soluzione	not available	038630024	GENETIC SPA	IT
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	2008099989	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED	UK
ACTIFED ALLERGIE CETIRIZINE 10 mg, comprimé pelliculé sécable	DE/H/3848/001	34009 380 213 1 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	2008099989	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER 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
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine®, 10 mg, Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
LivoReactine 10 mg	DE/H/3848/001	24/0028/04-S	MCNEIL PRODUCTS LIMITED	SK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
LivoReactine 10 mg	DE/H/3848/001	24/0028/04-S	MCNEIL PRODUCTS LIMITED	SK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	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
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
LivoReactine 10 mg	DE/H/3848/001	24/0028/04-S	MCNEIL PRODUCTS LIMITED	SK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine®	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON	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
10 mg Filmtabletten			GMBH	
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine® 10 mg Filmtabletten	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON GMBH	DE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON	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
filmomhulde tabletten			CONSUMER B.V.	
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	RVG 29446	JOHNSON & JOHNSON CONSUMER B.V.	NL
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg	DE/H/3848/001	BE150491	JOHNSON & JOHNSON	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
filmomhulde tabletten			CONSUMER N.V./S.A.	
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg	DE/H/3848/001	BE150491	JOHNSON & JOHNSON	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
Filmtabletten			CONSUMER N.V./S.A.	
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg filmomhulde tabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Reactine 10 mg Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
ALAIRGIX ALLERGIE CETIRIZINE 10 mg, comprimé à sucer sécable	not available	NL 30142-3400936707580	COOPERATION PHARMACEUTIQUE FRANCAISE	FR