EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

06 July 2023
EMA/316182/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): cetirizine

Procedure No. PSUSA/00000628/202211

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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
Zyrtec 1 mg/ml orale Lösung	IE/H/0209/003	1-20143	UCB PHARMA GMBH (VIENNA AT)	AT
Zyrtec 10 mg Filmtabletten	IE/H/0209/001	1-19030	UCB PHARMA GMBH (VIENNA AT)	AT
Zyrtec 10 mg/ml Tropfen	IE/H/0209/002	1-20142	UCB PHARMA GMBH (VIENNA AT)	AT
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 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 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 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	DE/H/3848/001	BE150491	JOHNSON & JOHNSON	BE

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comprimés pelliculés			CONSUMER N.V./S.A.	
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 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 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 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 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 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 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 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	DE/H/3848/001	BE150491	JOHNSON & JOHNSON	BE

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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 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 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 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 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 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 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 Filmtabletten	DE/H/3848/001	BE150491	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Zyrtec 1 mg/ml drank	IE/H/0209/003	BE154673	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 1 mg/ml solution buvable	IE/H/0209/003	BE154673	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg comprimés pelliculés	IE/H/0209/001	BE135877	UCB PHARMA S.A. (ANDERL BE)	BE
Zyrtec 10 mg	IE/H/0209/001	BE135877	UCB PHARMA S.A. (ANDERL	BE

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filmomhulde tabletten			BE)	
Zyrtec 10 mg Filmtabletten	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/ml solution buvable en gouttes	IE/H/0209/002	BE149746	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
Зиртек 10 mg филмирани таблетки	IE/H/0209/001	20020795	UCB PHARMA GMBH (MONHEIM DE)	BG
Зиртек 10 mg/ml перорални капки, разтвор	IE/H/0209/002	20020794	UCB PHARMA GMBH (MONHEIM DE)	BG
Zyrtec 1 mg/ml πόσιμο διάλυμα.	IE/H/0209/003	16365	UCB PHARMA S.A. (ANDERL BE)	CY
Zyrtec 10 mg επικαλυμμένα με λεπτό υμένιο δισκία.	IE/H/0209/001	13066	UCB PHARMA S.A. (ANDERL BE)	CY
Zyrtec 10 mg potahované tablety	IE/H/0209/001	24/024/92-S/C	UCB S.R.O. (PRAGUE CZ)	CZ
Zyrtec 10 mg/ml perorální kapky, roztok	IE/H/0209/002	24/1030/92-S/C	UCB S.R.O. (PRAGUE CZ)	CZ
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,	DE/H/3848/001	46103.00.00	JOHNSON & JOHNSON	DE

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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
Zyrtec® 10 mg Filmtabletten	IE/H/0209/001	14836.00.00	UCB PHARMA GMBH (MONHEIM DE)	DE
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
Benaday, filmovertrukne tabletter	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK
Benaday, filmovertrukne	DE/H/3848/001	30712	MCNEIL DENMARK APS	DK

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tabletter				
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
Zyrtec, filmovertrukne tabletter	IE/H/0209/001	13082	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec, oral opløsning	IE/H/0209/003	14784	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec, orale dråber, opløsning	IE/H/0209/002	13651	UCB NORDIC A/S (COPENHAGEN DK)	DK
Zyrtec, 10 mg õhukese polümeerikattega tabletid	IE/H/0209/001	162097	UCB PHARMA OY FINLAND (ESPOO FI)	EE
Zyrtec, 10 mg/ml suukaudsed tilgad, lahus	IE/H/0209/002	162197	UCB PHARMA OY FINLAND (ESPOO FI)	EE
Alerlisin 1 mg/ml solución oral.	not available	62.787	RETRAIN, S.A.U.	ES
Alerlisin 10 mg comprimidos recubiertos con película	not available	58.499	RETRAIN, S.A.U.	ES
Alerlisin 10 mg/ml gotas orales en solución.	not available	62.788	RETRAIN, S.A.U.	ES
Zyrtec 1 mg/ml solución oral.	IE/H/0209/003	60.279	UCB PHARMA S.A. (MADRID ES)	ES
Zyrtec 10 mg comprimidos recubiertos con película	IE/H/0209/001	58.481	UCB PHARMA S.A. (MADRID ES)	ES

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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 1 mg/ml oral lösning	IE/H/0209/003	11322	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec 10 mg filmdragerade tabletter	IE/H/0209/001	10385	UCB PHARMA OY FINLAND (ESPOO FI)	FI
Zyrtec 10 mg tabletti, kalvopäällysteinen	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 10 mg/ml tipat, liuos	IE/H/0209/002	11321	UCB PHARMA OY FINLAND (ESPOO FI)	FI
ALAIRGIX ALLERGIE CETIRIZINE 10 mg, comprimé à sucer sécable	not available	NL 30142-3400936707580	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
CETIRIZINE EG LABO CONSEIL 10 mg, comprimé à sucer	not available	NL30927	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
DRILL ALLERGIE CETIRIZINE 10 mg, comprimé à sucer	not available	34009 381 535 2 1	PIERRE FABRE MEDICAMENT	FR
REACTINE 10 mg, comprimé pelliculé sécable	DE/H/3848/001	34009 380 213 1 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
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	IE/H/0209/001	34009 339 615 1 0	UCB PHARMA S.A. (COLOMBES FR)	FR

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sécable				
ZYRTEC 10 mg/ml, solution buvable en gouttes	IE/H/0209/002	34009 341 770 0 2	UCB PHARMA S.A. (COLOMBES FR)	FR
ZYRTEC 2,5 mg/2,5 ml, solution buvable en flacon	IE/H/0209/003	34009 333 019 8 9	UCB PHARMA S.A. (COLOMBES FR)	FR
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
ZYRTECSET 10 mg, comprimé pelliculé sécable	not available	34009 364 616 8 0	UCB PHARMA S.A. (COLOMBES FR)	FR
ZYRTECSET 10 mg, comprimé pelliculé sécable	not available	34009 364 616 8 0	UCB PHARMA S.A. (COLOMBES FR)	FR
Ζιρτέκ 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0209/001	1075/08-01-2015	UCB A.E. (ATHENS GR)	GR
Ζιρτέκ 10 mg/ml πόσιμες σταγόνες, διάλυμα	IE/H/0209/002	1076/08-01-2015	UCB A.E. (ATHENS GR)	GR
Zyrtec 10 mg filmtabletta	IE/H/0209/001	OGYI-T-1599/03	UCB MAGYARORSZÁG KFT (BUDAPEST HU)	HU
Zyrtec 10 mg filmtabletta	IE/H/0209/001	OGYI-T-1599/02	UCB MAGYARORSZÁG KFT (BUDAPEST HU)	HU
Zyrtec 10 mg/ml belsőleges oldatos cseppek	IE/H/0209/002	OGYI-T-4491/01	UCB MAGYARORSZÁG KFT (BUDAPEST HU)	HU
Zirtek 1 mg/ml oral solution	IE/H/0209/003	PA0891/008/003	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek 1 mg/ml oral solution	IE/H/0209/003	PA0891/008/003	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek 10 mg film-coated tablets	IE/H/0209/001	PA0891/008/002	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek 10 mg film-coated tablets	IE/H/0209/001	PA0891/008/002	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek 10 mg/ml oral drops, solution	IE/H/0209/002	PA0891/008/004	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE

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Zirtek 10 mg/ml oral drops, solution	IE/H/0209/002	PA0891/008/004	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PA0891/008/005	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PA0891/008/005	UCB PHARMA IRELAND LTD (DUBLIN IE)	IE
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
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
Formistin 10 mg/ml gocce orali, soluzione.	not available	027329022	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
Formistin 10 mg/ml gocce orali, soluzione.	not available	027329022	ISTITUTO LUSO FARMACO D'ITALIA S.P.A.	IT
GENERIT 10 mg/ml gocce orali, soluzione	not available	038628020	GENETIC SPA	IT
GENERIT 10 mg/ml gocce orali, soluzione	not available	038628044	GENETIC SPA	IT
GENERIT 10 mg/ml gocce orali, soluzione	not available	038628057	GENETIC SPA	IT
RAINGEN 10 mg/ml gocce orali, soluzione	not available	038630024	GENETIC SPA	IT
RAINGEN 10 mg/ml gocce orali, soluzione	not available	038630048	GENETIC SPA	IT
RAINGEN 10 mg/ml gocce orali, soluzione	not available	038630051	GENETIC SPA	IT
RITECAM 10 mg/ml gocce orali, soluzione	not available	038629022	GENETIC SPA	IT
RITECAM 10 mg/ml gocce orali, soluzione	not available	038629046	GENETIC SPA	IT

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RITECAM 10 mg/ml gocce orali, soluzione	not available	038629059	GENETIC SPA	IT
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
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
Zirtec 10 mg/ml gocce orali, soluzione	IE/H/0209/002	026894028	UCB PHARMA SPA (MILAN IT)	IT
Zirtec 10 mg/ml gocce orali, soluzione	IE/H/0209/002	026894081	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 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 dengtos tabletės	IE/H/0209/001	LT/1/97/2859/002	UCB PHARMA OY FINLAND (ESPOO FI)	LT
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/ml geriamieji lašai (tirpalas)	IE/H/0209/002	LT/1/97/2859/005	UCB PHARMA OY FINLAND (ESPOO FI)	LT
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	2008099989	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Reactine 10 mg comprimés pelliculés	DE/H/3848/001	2008099989	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
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 solution buvable en gouttes	IE/H/0209/002	2011101304	UCB PHARMA S.A. (ANDERL BE)	LU
Zyrtec 1 mg/ml šķīdums	IE/H/0209/003	98-0791	UCB PHARMA OY FINLAND	LV

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
iekšķīgai lietošanai			(ESPOO FI)	
Zyrtec 10 mg apvalkotās tabletes	IE/H/0209/001	98-0688	UCB PHARMA OY FINLAND (ESPOO FI)	LV
Zyrtec 10 mg apvalkotās tabletes	IE/H/0209/001	98-0731	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 oral solution	IE/H/0209/003	MA030/00902	UCB PHARMA S.A. (ANDERL BE)	MT
Zyrtec 10 mg film-coated tablets	IE/H/0209/001	MA030/00901	UCB PHARMA S.A. (ANDERL BE)	MT
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 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

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 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
Zyrtec 1 mg/ml drank	IE/H/0209/003	RVG 14635	UCB PHARMA B.V. (BREDA NL)	NL
Zyrtec 10 mg filmomhulde tabletten	IE/H/0209/001	RVG 13010	UCB PHARMA B.V. (BREDA NL)	NL
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 10 mg/ml dråper, oppløsning	IE/H/0209/002	8173	UCB NORDIC A/S (COPENHAGEN DK)	NO
Zyrtec UCB, 10 mg, tabletki powlekane	not available	11212	VEDIM SP. Z O.O.	PL
Zyrtec UCB, 10 mg, tabletki powlekane	not available	11212	VEDIM SP. Z O.O.	PL
Zyrtec, 1 mg/ml, roztwór doustny	IE/H/0209/003	7815	VEDIM SP. Z O.O.	PL
Zyrtec, 1 mg/ml, roztwór doustny	IE/H/0209/003	7815	VEDIM SP. Z O.O.	PL
Zyrtec, 10 mg, tabletki powlekane	IE/H/0209/001	R/1846	VEDIM SP. Z O.O.	PL
Zyrtec, 10 mg, tabletki powlekane	IE/H/0209/001	R/1846	VEDIM SP. Z O.O.	PL
Zyrtec, 10 mg/ml, krople doustne, roztwór	IE/H/0209/002	R/1847	VEDIM SP. Z O.O.	PL
Zyrtec, 10 mg/ml, krople doustne, roztwór	IE/H/0209/002	R/1847	VEDIM SP. Z O.O.	PL
Zyrtec 1 mg/ml Solução oral	IE/H/0209/003	2051795	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA.	PT
Zyrtec 10 mg comprimidos revestidos por película	IE/H/0209/001	9717900	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA.	PT
Zyrtec 10 mg	IE/H/0209/001	5738000	UCB PHARMA (PRODUTOS	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
comprimidos revestidos por película			FARMACÊUTICOS), LDA.	
Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/02	UCB PHARMA GMBH (MONHEIM DE)	RO
Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/01	UCB PHARMA GMBH (MONHEIM DE)	RO
Zyrtec 10 mg/ml picături orale, soluţie	IE/H/0209/002	5076/2012/03	UCB PHARMA GMBH (MONHEIM DE)	RO
Zyrlex 1 mg/ml oral lösning	SE/H/2024/002	12678	MACURE PHARMA APS	SE
Zyrlex 10 mg filmdragerade tabletter	SE/H/2024/001	11209	MACURE PHARMA APS	SE
Zyrlex 10 mg/ml orala droppar, lösning	SE/H/2024/003	11673	MACURE PHARMA APS	SE
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 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 obložene tablete	IE/H/0209/001	H/01/01728/007	UCB PHARMA S.A. (ANDERL BE)	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 10 mg filmom obalené tablety cetirizíniumdichlorid	IE/H/0209/001	24/0024/92-S	UCB S.R.O. (PRAGUE CZ)	SK
Zyrtec 10 mg/ml perorálne roztokové kvapky cetirizíniumdichlorid	IE/H/0209/002	24/1030/92-S	UCB S.R.O. (PRAGUE CZ)	SK
Asda Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Asda Hayfever & Allergy Relief	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Asda Hayfever & Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
BecoAllergy 10mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Benadryl Allergy Children's 1mg/ml Oral Solution	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	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
Benadryl Allergy Children's 1mg/ml Oral Solution	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl Allergy Children's 1mg/ml Oral Solution	not available	PL 15513/0138	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl Allergy Children's 6+ 1mg/ml Oral Solution	not available	PL 15513/0124	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl Allergy One A Day 10mg Tablets	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Benadryl One A Day Relief	DE/H/3848/001	PL 15513/0118	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Boots Hayfever and allergy relief 10mg Tablets	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Boots Hayfever and allergy relief 10mg tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Careway Antihistamine Hayfever Relief 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Cetirizine dihydrochloride 10 mg film-coated tablets	not available	PL 20416/0566	CRESCENT PHARMA LIMITED	XI
Cetirizine Dihydrochloride 10mg Film-coated Tablets	not available	PL 36687/0242	TORRENT PHARMA (UK) LTD.	XI
Cetirizine Hydrochloride 10mg Film-coated	not available	PL 36687/0241	TORRENT PHARMA (UK) 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
Tablets				
Cetirizine Hydrochloride 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Dr Reddy's Histease Allergy Relief 10mg Tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) LTD.	XI
EM Pharma Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
EM Pharma Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) LTD.	XI
essential Waitrose hayfever & allergy relief tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Galpharm Hayfever and Allergy Relief	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Galpharm Hayfever and Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Health Essential Allergy Relief 10 mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Lloyds Pharmacy Antihistamine Hayfever Relief 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Morrisons Hayfever & Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Morrisons Hayfever & Allergy Relief 10mg Tablets	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Numark Hayfever and Allergy Relief	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Numark Hayfever and Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Optipharma Hayfever and Allergy Relief 10mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Piriteze Allergy 1mg/ml Syrup	not available	PL 44673/0095	GLAXOSMITHKLINE CONSUMER HEALTHCARE	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
			(UK) TRADING LIMITED	
Piriteze Allergy Relief 1mg/ml Syrup	not available	PL 44673/0096	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	XI
Piriteze Allergy Tablets	not available	PL 44673/0098	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	XI
Piriteze Children's Hayfever & Allergy 1mg/ml Syrup	not available	PL 44673/0095	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	XI
Piriteze Children's Hayfever & Allergy 1mg/ml Syrup	not available	PL 44673/0096	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	XI
Piriteze Hayfever & Allergy 10mg Film Coated Tablets	not available	PL 44673/0097	GLAXOSMITHKLINE CONSUMER HEALTHCARE (UK) TRADING LIMITED	XI
Pollenase Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Pollenase Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) LTD.	XI
POLLENSHIELD HAYFEVER RELIEF Cetirizine hydrochloride 10mg Film-coated Tablets	not available	PL 0142/0606	ACCORD-UK LIMITED	XI
Sainsbury's hayfever and allergy relief	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Sainsbury's Healthcare hayfever and allergy relief 10mg tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Spar Hayfever & Allergy Relief 10mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Superdrug Hayfever & Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Tesco Allergy & Hayfever Relief 10mg Tablets	not available	PL 08553/0194	DR. REDDY'S LABORATORIES (UK) 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
Tesco Allergy and Hayfever Relief 10mg Tablets	not available	PL 08553/0193	DR. REDDY'S LABORATORIES (UK) LTD.	XI
Tesco Health Hayfever and Allergy Relief 10mg Tablets	not available	PL 16028/0066	GALPHARM HEALTHCARE LIMITED	XI
Tesco Health Hayfever and Allergy Relief 10mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
The Co-Operative Hayfever & Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
The Local Independent Trading Company Ltd Hayfever & Allergy Relief 10mg Tablets	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Value Health Hayfever & Allergy Relief	not available	PL 16028/0067	GALPHARM HEALTHCARE LIMITED	XI
Wockhardt Allergy and Hayfever Relief 10mg Film-coated Tablets	not available	PL 29831/0681	WOCKHARDT UK LTD	XI
Ziralton Allergy Relief for Children 5mg/5ml Oral Solution	not available	PL 04917/0140	PINEWOOD LABORATORIES LIMITED	XI
Zirtek Allergy 10 mg film-coated tablets	IE/H/0209/001	PL 00039/0542	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy 10 mg film-coated tablets	IE/H/0209/001	PL 00039/0542	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PL 00039/0561	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy Relief 10 mg film-coated tablets	not available	PL 00039/0561	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy Relief for Children 1 mg/mL oral solution	not available	PL 00039/0541	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy Relief for Children 1 mg/mL oral solution	not available	PL 00039/0541	UCB PHARMA LTD (SLOUGH GB)	XI
Zirtek Allergy Solution 1	IE/H/0209/003	PL 00039/0540	UCB PHARMA LTD (SLOUGH	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
mg/ml oral solution			GB)	
Zirtek Allergy Solution 1 mg/ml oral solution	IE/H/0209/003	PL 00039/0540	UCB PHARMA LTD (SLOUGH GB)	XI