5 May 2022
EMA/267394/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: tolterodine

Procedure no.: PSUSA 00002993 202109

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
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Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
Detrusitol 1 mg - Filmtabletten	SE/H/0139/001	1-22389	UPJOHN EESV	AT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 1 mg compresse rivestite con film	SE/H/0139/001	034168017	VIATRIS PHARMA S.R.L.	IT
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés	SE/H/0139/	BE191712	UPJOHN SRL	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
pelliculés	001			
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191721	UPJOHN SRL	BE
Detrusitol 1 mg comprimés pelliculés	SE/H/0139/001	BE191712	UPJOHN SRL	BE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PA 23055/014/001	UPJOHN EESV	IE
Detrusitol 1 mg film-coated tablets	not available	MA1396/00501	UPJOHN HELLAS L.T.D.	MT
Detrusitol 1 mg film-coated tablets	not available	MA1396/00501	UPJOHN HELLAS L.T.D.	MT
Detrusitol 1 mg film-coated tablets	not available	MA1396/00501	UPJOHN HELLAS L.T.D.	MT
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/0015	UPJOHN UK LIMITED	XI
Detrusitol 1 mg film-coated tablets	SE/H/0139/001	PL 50622/001	UPJOHN 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
		5		
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13101	UPJOHN EESV	FI
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
Detrusitol 1 mg filmdragerade tabletter	SE/H/0139/001	13475	UPJOHN EESV	SE
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
DETRUSITOL 1 mg Filmtabletten	SE/H/0139/001	2009020158	UPJOHN SRL	LU
Detrusitol 1 mg plėvele dengtos tabletės	not available	LT/1/99/1258/001	UPJOHN EESV	LT
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	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
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg,	SE/H/0139/	34009 346	PFIZER PFE FRANCE	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
comprimé pelliculé	001	161 2 9		
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	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
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 102 6 4	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 161 2 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 160 6 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 561 495 9 9	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 159 8 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 100 3 5	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 156 9 6	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 158 1 8	PFIZER PFE FRANCE	FR
DETRUSITOL 1 mg, comprimé pelliculé	SE/H/0139/001	34009 346 157 5 7	PFIZER PFE FRANCE	FR
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 1 mg, filmomhulde tabletten	SE/H/0139/001	RVG 22148	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
Detrusitol 2 mg - Filmtabletten	SE/H/0139/002	1-22390	UPJOHN EESV	AT
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 2 mg	SE/H/0139/	034168029	VIATRIS PHARMA 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
compresse rivestite con film	002			
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL 2 mg compresse rivestite con film	SE/H/0139/002	034168029	VIATRIS PHARMA S.R.L.	IT
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg comprimés pelliculés	SE/H/0139/002	2009020159	UPJOHN SRL	LU
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PA 23055/014/002	UPJOHN EESV	IE
Detrusitol 2 mg film-coated tablets	not available	MA1396/00502	UPJOHN HELLAS L.T.D.	MT
Detrusitol 2 mg film-coated tablets	not available	MA1396/00502	UPJOHN HELLAS L.T.D.	MT
Detrusitol 2 mg film-coated tablets	not available	MA1396/00502	UPJOHN HELLAS L.T.D.	MT
Detrusitol 2 mg film-coated	SE/H/0139/	PL	UPJOHN 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
tablets	002	50622/0016		
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg film-coated tablets	SE/H/0139/002	PL 50622/0016	UPJOHN UK LIMITED	XI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13102	UPJOHN EESV	FI
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmdragerade tabletter	SE/H/0139/002	13476	UPJOHN EESV	SE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg	SE/H/0139/	BE191694	UPJOHN SRL	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	002			
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191703	UPJOHN SRL	BE
Detrusitol 2 mg filmomhulde tabletten	SE/H/0139/002	BE191694	UPJOHN SRL	BE
Detrusitol 2 mg plėvele dengtos tabletės	not available	LT/1/99/1258/002	UPJOHN EESV	LT
Detrusitol 2 mg trde kapsule s podaljšanim sproščanjem	not available	H/99/00455/001	UPJOHN EESV	SI
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	CY
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	023122	UPJOHN HELLAS L.T.D.	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
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol 2 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0139/002	45485/03-07-2009	UPJOHN HELLAS L.T.D.	GR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	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
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg,	SE/H/0139/	34009 346	PFIZER PFE FRANCE	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
comprimé pelliculé	002	164 1 9		
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 165 8 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 109 0 5	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 561 496 5 0	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 162 9 7	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 164 1 9	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 163 5 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 166 4 8	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 108 4 4	PFIZER PFE FRANCE	FR
DETRUSITOL 2 mg, comprimé pelliculé	SE/H/0139/002	34009 346 167 0 9	PFIZER PFE FRANCE	FR
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol 2 mg, comprimidos recubiertos con película	SE/H/0139/002	62.022	PFIZER, S.L.	ES
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 2 mg, filmomhulde tabletten	SE/H/0139/002	RVG 22149	VIATRIS NETHERLANDS B.V.	NL
Detrusitol 4 mg trde kapsule s podaljšanim	not available	H/99/00455/002	UPJOHN EESV	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
sproščanjem				
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	FARMASIERRA LABORATORIOS, S.L.	ES
Detrusitol Neo 4 mg, cápsulas duras de liberación prolongada	SE/H/0139/004	64.245	PFIZER, S.L.	ES
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
Detrusitol retard 2 mg -	SE/H/0139/	1-24214	UPJOHN EESV	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
Kapseln	003			
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
Detrusitol retard 2 mg - Kapseln	SE/H/0139/003	1-24214	UPJOHN EESV	AT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio	SE/H/0139/003	034168106	VIATRIS PHARMA 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
prolungato				
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA 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
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg	SE/H/0139/	034168144	VIATRIS PHARMA S.R.L.	IT

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capsule rigide a rilascio prolungato	003			
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio	SE/H/0139/003	034168068	VIATRIS PHARMA 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
prolungato				
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168272	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168120	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168144	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168106	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168082	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168118	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168094	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168068	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168031	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168070	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168043	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 2 mg capsule rigide a rilascio prolungato	SE/H/0139/003	034168056	VIATRIS PHARMA S.R.L.	IT
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
Detrusitol Retard 2 mg	SE/H/0139/	IS/1/01/02	UPJOHN EESV	IS

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
forðahylki, hörð.	003	8/01		
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
Detrusitol Retard 2 mg forðahylki, hörð.	SE/H/0139/003	IS/1/01/028/01	UPJOHN EESV	IS
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	BE228995	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération prolongée	SE/H/0139/003	2009020160	UPJOHN SRL	LU
DETRUSITOL Retard 2 mg gélules à libération	SE/H/0139/003	2009020160	UPJOHN SRL	LU

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prolongée				
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
DETRUSITOL Retard 2 mg harde capsules met verlengde afgifte.	SE/H/0139/003	BE229004	UPJOHN SRL	BE
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
Detrusitol retard 4 mg - Kapseln	SE/H/0139/004	1-24215	UPJOHN EESV	AT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg	SE/H/0139/	034168157	VIATRIS PHARMA S.R.L.	IT

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capsule rigide a rilascio prolungato	004			
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio	SE/H/0139/004	034168233	VIATRIS PHARMA 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
prolungato				
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA 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
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg	SE/H/0139/	034168221	VIATRIS PHARMA 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
capsule rigide a rilascio prolungato	004			
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168284	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168195	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio	SE/H/0139/004	034168260	VIATRIS PHARMA S.R.L.	IT

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prolungato				
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168207	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168219	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168157	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168171	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168169	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168183	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168221	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168233	VIATRIS PHARMA S.R.L.	IT
DETRUSITOL Retard 4 mg capsule rigide a rilascio prolungato	SE/H/0139/004	034168245	VIATRIS PHARMA S.R.L.	IT
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
Detrusitol Retard 4 mg forðahylki, hörð.	SE/H/0139/004	IS/1/01/028/02	UPJOHN EESV	IS
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg	SE/H/0139/	BE229022	UPJOHN SRL	BE

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gélules à libération prolongée	004			
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	BE229022	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg gélules à libération prolongée	SE/H/0139/004	2009020161	UPJOHN SRL	LU
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
DETRUSITOL Retard 4 mg Hartkapseln, retardiert	SE/H/0139/004	BE229013	UPJOHN SRL	BE
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	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
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/003	32467	UPJOHN EESV	DK
Detrusitol Retard, hårde depotkapsler	SE/H/0139/004	32469	UPJOHN EESV	DK
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapsel, harde.	SE/H/0139/003	01-3746	UPJOHN EESV	NO
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg depotkapseli, kova	SE/H/0139/003	16472	UPJOHN EESV	FI
Detrusitol SR 2 mg	SE/H/0139/	16184	UPJOHN EESV	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
depotkapslar, hårda.	003			
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg depotkapslar, hårda.	SE/H/0139/003	16184	UPJOHN EESV	SE
Detrusitol SR 2 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/003	UPJOHN EESV	LT
Detrusitol SR 2 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/004	UPJOHN EESV	LT
Detrusitol SR 2 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/003	UPJOHN EESV	LT
Detrusitol SR 2 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/004	UPJOHN EESV	LT
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, capsules met verlengde afgifte, hard.	SE/H/0139/003	RVG 26669	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	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
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, prolonged-release capsules, hard	SE/H/0139/003	PA 23055/007/001	UPJOHN EESV	IE
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 2 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/003	45486/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapsel, harde.	SE/H/0139/004	01-3747	UPJOHN EESV	NO
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	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
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16473	UPJOHN EESV	FI
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg depotkapslar, hårda.	SE/H/0139/004	16185	UPJOHN EESV	SE
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/006	UPJOHN EESV	LT
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/005	UPJOHN EESV	LT
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/007	UPJOHN EESV	LT
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/006	UPJOHN EESV	LT
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/005	UPJOHN EESV	LT
Detrusitol SR 4 mg pailginto atpalaidavimo kietosios kapsulės	not available	LT/1/99/1258/007	UPJOHN EESV	LT
DETRUSITOL SR 4 mg tvrdé kapsuly s predĺženým uvoľňovaním	not available	73/0149/04-S	UPJOHN EESV	SK
DETRUSITOL SR 4 mg tvrdé tobolky s prodlouženým uvolňováním	not available	53/236/02-C	UPJOHN EESV	CZ
Detrusitol SR 4 mg, capsules met verlengde	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL

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afgifte, hard.				
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, capsules met verlengde afgifte, hard.	SE/H/0139/004	RVG 26670	VIATRIS NETHERLANDS B.V.	NL
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PA 23055/007/002	UPJOHN EESV	IE
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	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
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol SR 4 mg, καψάκιο παρατεταμένης αποδέσμευσης, σκληρό	SE/H/0139/004	45487/3-7-2009	UPJOHN HELLAS L.T.D.	GR
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol XL 4 mg, prolonged-release capsules, hard	SE/H/0139/004	PL 50622/0017	UPJOHN UK LIMITED	XI
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 1 mg Filmtabletten	SE/H/0139/001	42055.00.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE

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Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® 2 mg Filmtabletten	SE/H/0139/002	42055.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Detrusitol® retard 4 mg Hartkapsel	SE/H/0139/004	51841.01.00	PFIZER OFG GERMANY GMBH	DE
Protol SR 2 mg, prolonged-release capsules, hard	SE/H/0242/004	16186	PFIZER AB	SE
Protol SR 2 mg, prolonged-release capsules, hard	SE/H/0242/004	16186	PFIZER AB	SE
Protol SR 4 mg prolonged-release capsules, hard	SE/H/0242/004	16187	PFIZER AB	SE
Protol SR 4 mg prolonged-release capsules, hard	SE/H/0242/004	16187	PFIZER AB	SE
Urotrol 2 mg, comprimidos recubiertos con película	not available	62.534	ALMIRALL, S.A.	ES