16 April 2020
EMA/222089/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: finasteride

Procedure no.: PSUSA/00001392/201908

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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
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 248 4 5	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 379 090 7 5	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 379 091 3 6	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 688 4 9	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 249 0 6	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 690 9 9	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 689 0 0	MSD FRANCE	FR
CHIBRO-PROSCAR 5 mg, comprimé pelliculé	not available	34009 335 691 5 0	MSD FRANCE	FR
Finasteride 1 mg Film-coated Tablets	PT/H/2278/001	PL 00289/0826	TEVA UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Finasteride 5 mg film-coated tablets	not available	PL 21880/0052	MEDREICH PLC	UK
Finasteride 5 mg Film-coated tablets	not available	PL 36687/0290	TORRENT PHARMA (UK) LTD.	UK
Finasteride 5mg Tablets	AT/H/0268/001	PL 04416/0816	SANDOZ LTD	UK
Finasteride Accord 1 mg apvalkotās tabletes	NL/H/1149/001	09-0154	ACCORD HEALTHCARE B.V.	LV
Finasteride AHCL 1 mg compresse rivestite con film	NL/H/1149/001	039595020	ACCORD HEALTHCARE LIMITED	IT
Finasteride AHCL 1 mg compresse rivestite con film	NL/H/1149/001	039595184	ACCORD HEALTHCARE LIMITED	IT
Finasteride AHCL 1 mg compresse rivestite con film	NL/H/1149/001	039595196	ACCORD HEALTHCARE LIMITED	IT
Finasteride AHCL 1 mg compresse rivestite con film	NL/H/1149/001	039595018	ACCORD HEALTHCARE LIMITED	IT
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952057	TEVA ITALIA 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
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952020	TEVA ITALIA S.R.L.	IT
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952071	TEVA ITALIA S.R.L.	IT
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952032	TEVA ITALIA S.R.L.	IT
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952044	TEVA ITALIA S.R.L.	IT
Finasteride Teva Generics 1 mg compresse rivestite con film	PT/H/2278/001	040952069	TEVA ITALIA S.R.L.	IT
Finastid 5 mg compresse rivestite con film	not available	028309019	NEOPHARMED GENTILI SPA	IT
Finastid 5 mg compresse rivestite con film	not available	028309021	NEOPHARMED GENTILI SPA	IT
Genaprost 5 mg compresse rivestite con film	not available	028371021	NEOPHARMED GENTILI SPA	IT
Genaprost 5 mg compresse rivestite con film	not available	028371019	NEOPHARMED GENTILI SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pilus 1 mg compressa rivestita con film.	not available	034824021	NEOPHARMED GENTILI SPA	IT
Pilus 1 mg compressa rivestita con film.	not available	034824058	NEOPHARMED GENTILI SPA	IT
Pilus 1 mg compressa rivestita con film.	not available	034824045	NEOPHARMED GENTILI SPA	IT
Pilus 1 mg compressa rivestita con film.	not available	034824033	NEOPHARMED GENTILI SPA	IT
Pilus 1 mg compressa rivestita con film.	not available	034824019	NEOPHARMED GENTILI SPA	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237014	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237038	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237040	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237053	MSD ITALIA 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
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237026	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237014	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237038	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237040	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237053	MSD ITALIA S.R.L.	IT
Propecia 1 mg compresse rivestite con film	SE/H/0158/001	034237026	MSD ITALIA S.R.L.	IT
Propecia 1 mg comprimate filmate	not available	9711/2017/01	MERCK SHARP & DOHME ROMANIA SRL	RO
Propecia 1 mg comprimidos recubiertos con película	SE/H/0158/001	62.441	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832988	MERCK SHARP & DOHME, LDA.	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
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832988	MERCK SHARP & DOHME, LDA.	PT
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832780	MERCK SHARP & DOHME, LDA.	PT
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832889	MERCK SHARP & DOHME, LDA.	PT
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832582	MERCK SHARP & DOHME, LDA.	PT
Propecia 1 mg comprimidos revestidos por película	SE/H/0158/ 001	2832681	MERCK SHARP & DOHME, LDA.	PT
Propecia 1 mg filmdragerade tabletter	SE/H/0158/001	13713	MERCK SHARP & DOHME BV	FI
Propecia 1 mg filmsko obložene tablete	not available	H/99/01294/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Propecia 1 mg Filmtabletten	SE/H/0158/001	1-24532	MERCK SHARP & DOHME GES.M.B.H.	AT
Propecia 1 mg Filmtabletten	SE/H/0158/001	44270.00.00	MSD SHARP & DOHME GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Propecia 1 mg filmuhúðaðar töflur	SE/H/0158/001	IS/1/02/010/01	MERCK SHARP & DOHME BV	IS
Propecia 1 mg tabletti, kalvopäällysteinen	SE/H/0158/001	13713	MERCK SHARP & DOHME BV	FI
PROPECIA 1 mg, comprimé pelliculé	SE/H/0158/ 001	34009 349 065 4 1	MSD FRANCE	FR
PROPECIA 1 mg, comprimé pelliculé	SE/H/0158/ 001	34009 349 070 8 1	MSD FRANCE	FR
PROPECIA 1 mg, comprimé pelliculé	SE/H/0158/ 001	34009 349 066 0 2	MSD FRANCE	FR
PROPECIA 1 mg, comprimé pelliculé	SE/H/0158/ 001	34009 349 068 3 1	MSD FRANCE	FR
PROPECIA 1 mg, comprimé pelliculé	SE/H/0158/ 001	34009 349 067 7 0	MSD FRANCE	FR
PROPECIA 1 mg, film-coated tablets	not available	18318	MERCK SHARP & DOHME BV	CY
Propecia 1 mg, film-coated tablets	not available	MA224/01101	MERCK SHARP & DOHME BV	MT

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
Propecia 1 mg, filmomhulde tabletten	SE/H/0158/001	RVG 27397	MERCK SHARP & DOHME BV	NL
PROPECIA 1 mg, tabletki powlekane	not available	4602	MSD POLSKA SP. Z O.O.	PL
PROPECIA 1mg, επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/158/01/R02	52369/30-06-2016	VIANEX S.A.	GR
Propecia, filmovertrukne tabletter	SE/H/0158/ 001	30252	MERCK SHARP & DOHME BV	DK
Propecia® 1 mg Film-Coated Tablets	not available	PL 0025/0351	MERCK SHARP & DOHME LTD.	UK
PROPECIA® 1 mg, comprimé pelliculé	SE/H/0158/001	2009030231	MSD BELGIUM BVBA/SPRL	LU
PROSCAR 5 mg compresse rivestite con film	not available	028308017	MSD ITALIA S.R.L.	IT
PROSCAR 5 mg compresse rivestite con film	not available	028308029	MSD ITALIA S.R.L.	IT
PROSCAR 5 mg compresse rivestite con film	not available	028308017	MSD ITALIA 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
PROSCAR 5 mg compresse rivestite con film	not available	028308029	MSD ITALIA S.R.L.	IT
PROSCAR 5 mg comprimate filmate	not available	5885/2013/01	MERCK SHARP & DOHME ROMANIA SRL	RO
PROSCAR 5 mg comprimate filmate	not available	5885/2013/02	MERCK SHARP & DOHME ROMANIA SRL	RO
PROSCAR 5 mg comprimidos recubiertos con película	not available	59.830	LABORATORIOS CHIBRET, S.A.	ES
Proscar 5 mg comprimidos revestidos por película	not available	2133189	MERCK SHARP & DOHME, LDA.	PT
Proscar 5 mg comprimidos revestidos por película	not available	2133387	MERCK SHARP & DOHME, LDA.	PT
PROSCAR 5 mg film-coated tablets	not available	PA 1286/017/001	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Proscar 5 mg filmdragerad tablett	not available	11644	MERCK SHARP & DOHME BV	SE
Proscar 5 mg filmdragerade tabletter	not available	10751	MERCK SHARP & DOHME BV	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
PROSCAR 5 mg filmom obložene tablete	not available	HR-H-039526411	MERCK SHARP & DOHME D.O.O.	HR
Proscar 5 mg Filmtabletten	not available	1-19689	MERCK SHARP & DOHME GES.M.B.H.	AT
Proscar 5 mg tabletter	not available	7841	MERCK SHARP & DOHME BV	NO
PROSCAR 5 mg tabletti, kalvopäällysteinen	not available	10751	MERCK SHARP & DOHME BV	FI
PROSCAR 5 mg, filmomhulde tabletten	not available	RVG 15482	MERCK SHARP & DOHME BV	NL
PROSCAR επικαλυμμένο με λεπτό υμένιο δισκίο 5mg/TAB	not available	119786/18/12-06-2019	VIANEX S.A.	GR
PROSCAR, 5 mg, tabletki powlekane	not available	R/3621	MSD POLSKA SP. Z O.O.	PL
PROSCAR® 5 mg, comprimés pelliculés	not available	BE160124	MSD BELGIUM BVBA/SPRL	BE
PROSCAR® 5 mg, comprimés pelliculés	not available	2003107721	MSD BELGIUM BVBA/SPRL	LU

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
PROSCAR® 5 mg, filmomhulde tabletten	not available	BE160124	MSD BELGIUM BVBA/SPRL	BE
PROSCAR® 5 mg, Filmtabletten	not available	BE160124	MSD BELGIUM BVBA/SPRL	BE
PROSCAR® 5mg film-coated Tablets (finasteride)	not available	PL 00025/0279	MERCK SHARP & DOHME LTD.	UK
PROSCAR°5 mg Filmtabletten Wirkstoff: Finasterid Zur Anwendung bei männlichen Patienten	not available	42859.00.00	MSD SHARP & DOHME GMBH	DE
PROSTIDE 5 mg compresse rivestite con film	not available	028356018	ALFASIGMA S.P.A.	IT
PROSTIDE 5 mg compresse rivestite con film	not available	028356020	ALFASIGMA S.P.A.	IT
Prostide 5 mg filmsko obložene tablete	not available	H/97/01301/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI