12 July 2018
EMA/475651/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): methylprednisolone

Procedure No.: PSUSA/00002026/201711

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Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
A d v a n t a n ® δερματικό διάλυμα 0,1% w/v	not available	3888/1-2-2002	BAYER HELLAS SA	GR
A d v a n t a n ®, δερματικό γαλάκτωμα (0,1% w/w)	not available	3887/01-02-2002	BAYER HELLAS SA	GR
A d v a n t a n αλοιφή εξωτερικής χρήσης 0,1%	not available	4938 / 28-01-2013	BAYER HELLAS SA	GR
A d v a n t a n κρέμα εξωτερικής χρήσης 0,1%	not available	69049/12/28-01-2013	BAYER HELLAS SA	GR
Advantan - dermatologische Lösung	not available	1-21795	BAYER AUSTRIA GMBH	AT
Advantan ® 0,1 % emulsiovoide	not available	11082	BAYER OY	FI
Advantan ® 0,1 % voide	not available	11083	BAYER OY	FI
Advantan ® 0,1% Creme	not available	22296.00.00	JENAPHARM GMBH & CO KG	DE
Advantan ® 0,1% Lösung, Lösung zur Anwendung auf der Haut	not available	22299.00.03	JENAPHARM GMBH & CO KG	DE
Advantan ® Lotion 0,1 % emulsio iholle	AT/H/0102/001	13292	BAYER OY	FI
Advantan 0,1 % Creme	not available	22299.00.00	JENAPHARM GMBH & CO KG	DE
Advantan 0,1 % emulsione cutanea	AT/H/0102/001	028159061	BAYER SPA	IT
Advantan 0,1 % emulsione cutanea	AT/H/0102/001	028159059	BAYER SPA	IT
Advantan 0,1 % Fettsalbe, Salbe	not available	22299.00.02	JENAPHARM GMBH & CO KG	DE
Advantan 0,1 % Salbe	not available	22299.00.01	JENAPHARM GMBH & CO KG	DE
Advantan 0,1% - Creme	not available	1-19575	BAYER AUSTRIA GMBH	AT
Advantan 0,1% crema	not available	028159010	BAYER SPA	IT
Advantan 0,1% crema	not available	028159073	BAYER SPA	IT
Advantan 0,1% crema idrofoba	not available	028159022	BAYER SPA	IT
Advantan 0,1% soluzione cutanea	not available	028159046	BAYER SPA	IT
Advantan 0,1% unguento	not available	028159034	BAYER SPA	IT
Advantan 0.1% cream	not available	MA896/00201	BAYER SPA	MT
Advantan 0.1% ointment	not available	MA896/00202	BAYER SPA	MT
Advantan 0.1% w/w Fatty Ointment	not available	PA 1410/70/3	BAYER LTD	IE

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Advantan 0.1% w/w Ointment	not available	PA 1410/70/2	BAYER LTD	IE
Advantan 1 mg/g cremă	not available	7187/2006/01	BAYER AG	RO
Advantan 1 mg/g cremă	not available	7187/2006/05	BAYER AG	RO
Advantan 1 mg/g cremă	not available	7187/2006/04	BAYER AG	RO
Advantan 1 mg/g cremă	not available	7187/2006/02	BAYER AG	RO
Advantan 1 mg/g cremă	not available	7187/2006/03	BAYER AG	RO
Advantan 1 mg/g creme	not available	2150886	BAYER PORTUGAL LDA	PT
Advantan 1 mg/g dermalna emulzija	not available	H/95/00126/005	BAYER D.O.O	SI
Advantan 1 mg/g kenőcs	not available	OGYI-T-20448/05	BAYER AG	HU
Advantan 1 mg/g kenőcs	not available	OGYI-T-20448/06	BAYER AG	HU
Advantan 1 mg/g kreem	not available	357701	BAYER AG	EE
Advantan 1 mg/g krém	not available	OGYI-T-20448/03	BAYER AG	HU
Advantan 1 mg/g krém	not available	OGYI-T-20448/04	BAYER AG	HU
Advantan 1 mg/g krema	not available	HR-H-429835039-01	BAYER DOO	HR
Advantan 1 mg/g krema	not available	H/95/00126/004	BAYER D.O.O	SI
Advantan 1 mg/g krema	not available	H/95/00126/003	BAYER D.O.O	SI
Advantan 1 mg/g kremas	not available	LT/1/97/2738/001	BAYER AG	LT
Advantan 1 mg/g krēms	not available	00-0245	BAYER AG	LV
Advantan 1 mg/g külsőleges emulzió	not available	OGYI-T-20448/10	BAYER AG	HU
Advantan 1 mg/g külsőleges emulzió	not available	OGYI-T-20448/09	BAYER AG	HU
Advantan 1 mg/g mast	not available	HR-H-434685681-01	BAYER DOO	HR
Advantan 1 mg/g mazilo	not available	H/95/00126/001	BAYER D.O.O	SI
Advantan 1 mg/g mazilo	not available	H/95/00126/002	BAYER D.O.O	SI
Advantan 1 mg/g pomada	not available	2151082	BAYER PORTUGAL LDA	PT
Advantan 1 mg/g salv	not available	171797	BAYER AG	EE
Advantan 1 mg/g tepalas	not available	LT/1/97/2738/002	BAYER AG	LT
Advantan 1 mg/g unguent	not available	7188/2006/02	BAYER AG	RO
Advantan 1 mg/g unguent	not available	7188/2006/04	BAYER AG	RO
Advantan 1 mg/g unguent	not available	7188/2006/03	BAYER AG	RO
Advantan 1 mg/g unguent	not available	7188/2006/05	BAYER AG	RO
Advantan 1 mg/g ziede	not available	00-0244	BAYER AG	LV
Advantan 1 mg/ml külsőleges oldat	not available	OGYI-T-20448/01	BAYER AG	HU

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
Advantan 1 mg/ml solução cutânea	not available	5297239	BAYER PORTUGAL LDA	PT
Advantan 1mg/ml külsőleges oldat	not available	OGYI-T-20448/02	BAYER AG	HU
Advantan cream 0.1% w/w	not available	PA1410/70/1	BAYER LTD	IE
Advantan krém	not available	46/0185/03-S	BAYER SPOL SRO	SK
Advantan krém 1 mg/g krém	not available	46/512/96-C	BAYER AG	CZ
Advantan masť	not available	46/0187/03-S	BAYER SPOL SRO	SK
Advantan mastný krém	not available	46/0186/03-S	BAYER SPOL SRO	SK
Advantan mastný krém 1 mg/g krém	not available	46/511/96-C	BAYER AG	CZ
Advantan Milch 0,1% Emulsion zur Anwendung auf der Haut	AT/H/0102/001	1-22211	BAYER AUSTRIA GMBH	AT
Advantan Milk 1 mg/g nahaemulsioon	not available	429203	BAYER AG	EE
Advantan Milk 1 mg/g odos emulsija	not available	LT/1/97/0148/001	BAYER AG	LT
Advantan Milk 1 mg/g odos emulsija	not available	LT/1/97/0148/002	BAYER AG	LT
Advantan Milk 1 mg/g uz ādas lietojama emulsija	not available	04-0250	BAYER AG	LV
Advantan Milk 1mg/g emulsie cutanată	not available	5507/2013/01	BAYER AG	RO
Advantan Milk 1mg/g emulsie cutanată	not available	5507/2013/02	BAYER AG	RO
Advantan mléko 1 mg/g Kožní emulze	not available	46/031/03-C	BAYER AG	CZ
Advantan zsíros 1 mg/g kenőcs	not available	OGYI-T-20448/08	BAYER AG	HU
Advantan zsíros 1 mg/g kenőcs	not available	OGYI-T-20448/07	BAYER AG	HU
Advantan, 1 mg/g unguent	not available	7188/2006/01	BAYER AG	RO
Advantan, 1 mg/g, Creme	not available	1993010082	BAYER SA NV	LU
Advantan, 1 mg/g, crème	not available	BE159327	BAYER SA NV	BE
Advantan, 1 mg/g, crème	not available	BE159327	BAYER SA NV	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
Advantan, 1 mg/g, crème	not available	1993010082	BAYER SA NV	LU
Advantan, 1 mg/g, emulsão cutânea	AT/H/0102/001	2710283	BAYER PORTUGAL LDA	PT
Advantan, 1 mg/g, emulsão cutânea	AT/H/0102/001	2710184	BAYER PORTUGAL LDA	PT
Advantan, 1 mg/g, emulsja na skórę	not available	R/7180	BAYER SP.Z.O.O	PL
Advantan, 1 mg/g, pommade	not available	BE159311	BAYER SA NV	BE
Advantan, 1 mg/g, pommade	not available	1993010081	BAYER SA NV	LU
Advantan, 1 mg/g, Salbe	not available	1993010081	BAYER SA NV	LU
Advantan, 1 mg/g, zalf	not available	BE159311	BAYER SA NV	BE
Advantan, 1 mg/g, Creme	not available	BE159327	BAYER SA NV	BE
Advantan, 1 mg/g, Salbe	not available	BE159311	BAYER SA NV	BE
Advantan, 1mg/g, krem	not available	R/7179	BAYER SP.Z.O.O	PL
Advantan,1 mg/g, maść	not available	R/7181	BAYER SP.Z.O.O	PL
Advantan® 0,1 % voide	not available	11083	BAYER OY	FI
Advantan® 0,1% - Fettsalbe	not available	1-19574	BAYER AUSTRIA GMBH	AT
Advantan® 0,1% - Salbe	not available	1-19576	BAYER AUSTRIA GMBH	AT
Advantan® 0,1% Fettsalbe, Salbe	not available	22296.00.02	JENAPHARM GMBH & CO KG	DE
Advantan® 0,1% Salbe	not available	22296.00.01	JENAPHARM GMBH & CO KG	DE
Advantan® Lotion 0,1 % kutan emulsion	AT/H/0102/001	13292	BAYER OY	FI
Advantan® Milch 0,1 % Emulsion zur Anwendung auf der Haut	AT/H/0102/001	22299.00.04	JENAPHARM GMBH & CO KG	DE
Advantan®0,1 % emulsiovoide	not available	11082	BAYER OY	FI
Adventan 1 mg/g crema	not available	60.137	BAYER HISPANIA SL	ES
Adventan 1 mg/g emulsión cutánea	AT/H/0102/001	62.662	BAYER HISPANIA SL	ES
Adventan 1 mg/g pomada	not available	60.138	BAYER HISPANIA SL	ES
Adventan 1 mg/g ungüento	not available	60.139	BAYER HISPANIA SL	ES
Adventan 1 mg/ml solución cutánea	not available	63.521	BAYER HISPANIA SL	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
DEPO MEDROL 40mg/mL sospensione iniettabile	not available	017932017	PFIZER ITALIA S.R.L.	IT
DEPO MEDROL 40mg/mL sospensione iniettabile	not available	017932029	PFIZER ITALIA S.R.L.	IT
Depo-Medrol 200 mg/5 ml Injektionssuspension	not available	BE124512	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 200 mg/5 ml suspensie voor injectie	not available	BE124512	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 200 mg/5 ml Suspension Injectable	not available	BE124512	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Injektionssuspension	not available	BE124537	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Injektionssuspension	not available	BE061844	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Injektionssuspension	not available	2008019606	PFIZER S.A. (BELGIUM)	LU
Depo-Medrol 40 mg/1 ml suspensie voor injectie	not available	BE124537	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml suspensie voor injectie	not available	BE061844	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Suspension Injectable	not available	BE124537	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Suspension Injectable	not available	BE061844	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 40 mg/1 ml Suspension Injectable	not available	2008019606	PFIZER S.A. (BELGIUM)	LU
DEPO-MEDROL 40 mg/1 ml, suspension injectable en flacon	not available	34009 553 153 5 3	PFIZER HOLDING FRANCE	FR
DEPO-MEDROL 40 mg/1 ml, suspension injectable en flacon	not available	34009 336 978 6 0	PFIZER HOLDING FRANCE	FR
DEPO-MEDROL 40 mg/1 ml, suspension injectable en flacon	not available	34009 302 935 2 2	PFIZER HOLDING FRANCE	FR
DEPO-MEDROL 40 mg/ml injekčná suspenzia	not available	56/0191/71-C/S	PFIZER EUROPE MA EEIG	SK

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
DEPO-MEDROL 40 mg/ml Injekční suspenze	not available	56/191/71-C	PFIZER, SPOL. S R.O.	CZ
Depo-Medrol 40 mg/ml injeksjonsvæske, suspensjon	not available	5074	PFIZER AS	NO
DEPO-MEDROL 40 mg/ml injektioneste, suspensio	not available	4491	PFIZER OY	FI
Depo-Medrol 40 mg/ml injektionsvätska, suspension	not available	4491	PFIZER OY	FI
Depo-Medrol 40 mg/ml injektionsvätska, suspension	not available	6941	PFIZER AB	SE
Depo-Medrol 40 mg/ml injektionsvätska, suspension, förfylld spruta	not available	9127	PFIZER AB	SE
Depo-Medrol 40 mg/ml stungulyf, dreifa	not available	691220	PFIZER APS	IS
Depo-Medrol 40 mg/ml suspensão injectável	not available	8114314	LABORATÓRIOS PFIZER, LDA.	PT
Depo-Medrol 40 mg/ml suspensão injectável	not available	5109038	LABORATÓRIOS PFIZER, LDA.	PT
Depo-Medrol 40 mg/ml suspenzija za injekcije	not available	UP/I-530-09/13-02/20	PFIZER CROATIA D.O.O.	HR
Depo-Medrol 40 mg/ml suspenzija za injiciranje	not available	H/93/00451/001	PFIZER LUXEMBOURG SARL	SI
DEPO-MEDROL 40 mg/ml szuszpenziós injekció	not available	OGYI-T-6384/01	PFIZER KFT.	HU
Depo-Medrol 40 mg/ml, suspensie voor injectie	not available	RVG 00605	PFIZER B.V.	NL
Depo-Medrol 80 mg/2 ml Injektionssuspension	not available	BE061835	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 80 mg/2 ml Injektionssuspension	not available	BE124521	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 80 mg/2 ml suspensie voor injectie	not available	BE124521	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 80 mg/2 ml suspensie voor injectie	not available	BE061835	PFIZER S.A. (BELGIUM)	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
Depo-Medrol 80 mg/2 ml Suspension Injectable	not available	BE061835	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 80 mg/2 ml Suspension Injectable	not available	BE124521	PFIZER S.A. (BELGIUM)	BE
Depo-Medrol 80 mg/2ml suspensão injectável	not available	8114306	LABORATÓRIOS PFIZER, LDA.	PT
DEPO-MEDROL, 40 mg/ml süstesuspensioon	not available	056994	PFIZER EUROPE MA EEIG	EE
DEPO-MEDROL, 40 mg/ml, zawiesina do wstrzykiwań	not available	R/1548	PFIZER EUROPE MA EEIG	PL
Depo-Medrol, injektionsvæske, suspension	not available	03239	PFIZER APS	DK
Depo-Medrone 40 mg/ml	not available	MA505/02001	PFIZER HELLAS, A.E.	MT
Depo-Medrone 40 mg/ml	not available	PL 00057/0963	PFIZER LIMITED	UK
Depo-Medrone® 40 mg/ml Suspension for Injection 1 ml vial	not available	PA 822/122/1	PFIZER HEALTHCARE IRELAND	IE
Depo-Medrone® 40 mg/ml Suspension for Injection 2 ml vial	not available	PA 822/122/2	PFIZER HEALTHCARE IRELAND	IE
Depo-Medrone® 40 mg/ml Suspension for Injection 3 ml vial	not available	PA 822/122/3	PFIZER HEALTHCARE IRELAND	IE
Fodier 0.1% w/w κρέμα	not available	47826/24-05-2017	VERISFIELD (UK) LTD	GR
Lexxema 1 mg/g crema	not available	63.186	ITALFARMACO S.A.	ES
Lexxema 1 mg/g emulsión cutánea	not available	63.904	ITALFARMACO S.A.	ES
Lexxema 1 mg/g pomada	not available	63.187	ITALFARMACO S.A.	ES
Lexxema 1 mg/g ungüento	not available	63.188	ITALFARMACO S.A.	ES
Lexxema 1 mg/ml solución cutánea	not available	63.890	ITALFARMACO S.A.	ES
Lexxema 1 mg/ml solución cutánea	not available	63.890	ITALFARMACO S.A.	ES
Medrol	not available	14643	PFIZER APS	DK
Medrol	not available	13369	PFIZER APS	DK
Medrol	not available	14644	PFIZER APS	DK
Medrol	not available	02457	PFIZER APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MEDROL	not available	44419/09/30-10-2014	PFIZER HELLAS, A.E.	GR
MEDROL	not available	44418/09/03-02-2010	PFIZER HELLAS, A.E.	GR
Medrol 100 mg tabletta	not available	OGYI-T-907/05	PFIZER KFT.	HU
MEDROL 100 mg tablety	not available	56/156/88-D/C	PFIZER, SPOL. S R.O.	CZ
MEDROL 100 mg, comprimé	not available	34009 372 323 6 4	PFIZER HOLDING FRANCE	FR
MEDROL 100 mg, comprimé	not available	34009 372 324 2 5	PFIZER HOLDING FRANCE	FR
MEDROL 16 mg compresse	not available	014159040	PFIZER ITALIA S.R.L.	IT
Medrol 16 mg comprimidos	not available	5790381	LABORATÓRIOS PFIZER, LDA.	PT
Medrol 16 mg comprimidos	not available	8315234	LABORATÓRIOS PFIZER, LDA.	PT
MEDROL 16 mg tablete	not available	UP/I-530-09/12-02/141	PFIZER CROATIA D.O.O.	HR
Medrol 16 mg tablete	not available	H/92/00985/002	PFIZER LUXEMBOURG SARL	SI
Medrol 16 mg tabletes	not available	99-1043	PFIZER EUROPE MA EEIG	LV
MEDROL 16 mg tabletės	not available	LT/1/94/1927/003	PFIZER EUROPE MA EEIG	LT
Medrol 16 mg tabletta	not available	OGYI-T-907/03	PFIZER KFT.	HU
Medrol 16 mg tabletter	not available	6488	PFIZER AS	NO
Medrol 16 mg tabletter	not available	09513	PFIZER AB	SE
Medrol 16 mg tabletter metylprednisolon	not available	7363	PFIZER OY	FI
MEDROL 16 mg tabletti	not available	7363	PFIZER OY	FI
MEDROL 16 mg tablety	not available	56/156/88-B/C	PFIZER, SPOL. S R.O.	CZ
MEDROL 16 mg tablety	not available	56/0156/88-C/S	PFIZER EUROPE MA EEIG	SK
MEDROL 16 mg, comprimé	not available	34009 331 900 9 5	PFIZER HOLDING FRANCE	FR
MEDROL 16 mg, comprimé	not available	34009 331 899 0 7	PFIZER HOLDING FRANCE	FR
MEDROL 16 mg, comprimé	not available	34009 331 897 8 5	PFIZER HOLDING FRANCE	FR
MEDROL 16 mg, comprimé	not available	34009 331 898 4 6	PFIZER HOLDING FRANCE	FR
MEDROL 16 mg, comprimé	not available	34009 556 820 2 8	PFIZER HOLDING FRANCE	FR
MEDROL 32 mg Comprimés (méthylprednisolone)	not available	BE500293	PFIZER S.A. (BELGIUM)	BE
MEDROL 32 mg Comprimés (méthylprednisolone)	not available	BE129832	PFIZER S.A. (BELGIUM)	BE
MEDROL 32 mg Comprimés (méthylprednisolone)	not available	2008019621	PFIZER S.A. (BELGIUM)	LU
MEDROL 32 mg tablete	not available	UP/I-530-09/12-02/142	PFIZER CROATIA D.O.O.	HR
Medrol 32 mg tablete	not available	H/92/00985/003	PFIZER LUXEMBOURG SARL	SI
Medrol 32 mg tabletta	not available	OGYI-T-907/04	PFIZER KFT.	HU

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
Medrol 32 mg Tabletten (Methylprednisolon)	not available	BE500293	PFIZER S.A. (BELGIUM)	BE
Medrol 32 mg Tabletten (Methylprednisolon)	not available	BE129832	PFIZER S.A. (BELGIUM)	BE
MEDROL 32 mg Tabletten (methylprednisolon)	not available	BE500293	PFIZER S.A. (BELGIUM)	BE
MEDROL 32 mg Tabletten (methylprednisolon)	not available	BE129832	PFIZER S.A. (BELGIUM)	BE
Medrol 32 mg Tabletten (Methylprednisolon)	not available	2008019621	PFIZER S.A. (BELGIUM)	LU
Medrol 32 mg tabletter metylprednisolon	not available	9288	PFIZER OY	FI
MEDROL 32 mg tabletti	not available	9288	PFIZER OY	FI
MEDROL 32 mg tablety	not available	56/156/88-C/C	PFIZER, SPOL. S R.O.	CZ
MEDROL 32 mg, comprimate	not available	6955/2006/01	PFIZER EUROPE MA EEIG	RO
MEDROL 4 mg compresse	not available	014159014	PFIZER ITALIA S.R.L.	IT
MEDROL 4 mg compresse	not available	014159026	PFIZER ITALIA S.R.L.	IT
MEDROL 4 mg Comprimés (méthylprednisolone)	not available	BE061214	PFIZER S.A. (BELGIUM)	BE
MEDROL 4 mg Comprimés (méthylprednisolone)	not available	BE500284	PFIZER S.A. (BELGIUM)	BE
MEDROL 4 mg Comprimés (méthylprednisolone)	not available	2008019618	PFIZER S.A. (BELGIUM)	LU
Medrol 4 mg comprimidos	not available	8315226	LABORATÓRIOS PFIZER, LDA.	PT
Medrol 4 mg comprimidos	not available	5790282	LABORATÓRIOS PFIZER, LDA.	PT
Medrol 4 mg comprimidos	not available	8315200	LABORATÓRIOS PFIZER, LDA.	PT
MEDROL 4 mg tablete	not available	UP/I-530-09/12-02/140	PFIZER CROATIA D.O.O.	HR
Medrol 4 mg tablete	not available	H/92/00985/001	PFIZER LUXEMBOURG SARL	SI
Medrol 4 mg tabletes	not available	99-1042	PFIZER EUROPE MA EEIG	LV
MEDROL 4 mg tabletės	not available	LT/1/94/1927/001	PFIZER EUROPE MA EEIG	LT
MEDROL 4 mg tabletės	not available	LT/1/94/1927/005	PFIZER EUROPE MA EEIG	LT
MEDROL 4 mg tabletės	not available	LT/1/94/1927/002	PFIZER EUROPE MA EEIG	LT
MEDROL 4 mg tabletės	not available	LT/1/94/1927/004	PFIZER EUROPE MA EEIG	LT
Medrol 4 mg tabletta	not available	OGYI-T-907/02	PFIZER KFT.	HU

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
Medrol 4 mg tabletta	not available	OGYI-T-907/01	PFIZER KFT.	HU
Medrol 4 mg Tabletten (Methylprednisolon)	not available	BE500284	PFIZER S.A. (BELGIUM)	BE
Medrol 4 mg Tabletten (Methylprednisolon)	not available	BE061214	PFIZER S.A. (BELGIUM)	BE
MEDROL 4 mg Tabletten (methylprednisolon)	not available	BE061214	PFIZER S.A. (BELGIUM)	BE
MEDROL 4 mg Tabletten (methylprednisolon)	not available	BE500284	PFIZER S.A. (BELGIUM)	BE
Medrol 4 mg Tabletten (Methylprednisolon)	not available	2008019618	PFIZER S.A. (BELGIUM)	LU
Medrol 4 mg tabletter	not available	6325	PFIZER AS	NO
Medrol 4 mg tabletter	not available	05862	PFIZER AB	SE
Medrol 4 mg tabletter metylprednisolon	not available	4493	PFIZER OY	FI
MEDROL 4 mg tabletti	not available	4493	PFIZER OY	FI
MEDROL 4 mg tablety	not available	56/156/88-A/C	PFIZER, SPOL. S R.O.	CZ
MEDROL 4 mg tablety	not available	56/0156/88-C/S	PFIZER EUROPE MA EEIG	SK
MEDROL 4 mg, comprimate	not available	6953/2006/01	PFIZER EUROPE MA EEIG	RO
MEDROL 4 mg, comprimé	not available	34009 553 946 5 5	PFIZER HOLDING FRANCE	FR
MEDROL 4 mg, comprimé	not available	34009 306 559 5 5	PFIZER HOLDING FRANCE	FR
MEDROL A 16 mg comprimate	not available	7622/2015/01	PFIZER EUROPE MA EEIG	RO
MEDROL A 16 mg Comprimés (méthylprednisolone)	not available	BE061372	PFIZER S.A. (BELGIUM)	BE
MEDROL A 16 mg Comprimés (méthylprednisolone)	not available	BE500302	PFIZER S.A. (BELGIUM)	BE
MEDROL A 16 mg Comprimés (méthylprednisolone)	not available	2008019620	PFIZER S.A. (BELGIUM)	LU
Medrol A 16 mg Tabletten (Methylprednisolon)	not available	BE500302	PFIZER S.A. (BELGIUM)	BE
Medrol A 16 mg Tabletten (Methylprednisolon)	not available	BE061372	PFIZER S.A. (BELGIUM)	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
MEDROL A 16 mg Tabletten (methylprednisolon)	not available	BE061372	PFIZER S.A. (BELGIUM)	BE
MEDROL A 16 mg Tabletten (methylprednisolon)	not available	BE500302	PFIZER S.A. (BELGIUM)	BE
Medrol A 16 mg Tabletten (Methylprednisolon)	not available	2008019620	PFIZER S.A. (BELGIUM)	LU
MEDROL PAK 4 mg Comprimés (méthylprednisolone)	not available	BE124494	PFIZER S.A. (BELGIUM)	BE
MEDROL PAK 4 mg Comprimés (méthylprednisolone)	not available	2008019619	PFIZER S.A. (BELGIUM)	LU
Medrol PAK 4 mg Tabletten (Methylprednisolon)	not available	BE124494	PFIZER S.A. (BELGIUM)	BE
MEDROL PAK 4 mg Tabletten (methylprednisolon)	not available	BE124494	PFIZER S.A. (BELGIUM)	BE
Medrol PAK 4 mg Tabletten (Methylprednisolon)	not available	2008019619	PFIZER S.A. (BELGIUM)	LU
Medrol, 16 mg tabletid	not available	056394	PFIZER EUROPE MA EEIG	EE
MEDROL, 16 mg, tabletki	not available	R/6832	PFIZER EUROPE MA EEIG	PL
Medrol, 4 mg tabletid	not available	056294	PFIZER EUROPE MA EEIG	EE
MEDROL, 4 mg, tabletki	not available	R/6831	PFIZER EUROPE MA EEIG	PL
Medrone Tablets 100 mg	not available	PL 00057/1011	PFIZER LIMITED	UK
Medrone Tablets 16 mg	not available	PL 00057/1479	PFIZER LIMITED	UK
Medrone Tablets 2 mg	not available	PL 00057/1013	PFIZER LIMITED	UK
Medrone Tablets 4 mg	not available	PL 00057/1014	PFIZER LIMITED	UK
Meprelon, 8 mg, tabletki	not available	17580	SUN-FARM SP. Z.O.O.	PL
Metasol 16 mg Tabletten	not available	1-31775	DERMAPHARM GMBH	AT
Metasol 8 mg Tabletten	not available	1-31774	DERMAPHARM GMBH	AT
Methylprednisolon Hikma 1000 mg Pulver zur Herstellung einer Injektionslösung	PT/H/0946/005	88420.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	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
Methylprednisolon Hikma 1000 mg Pulver zur Herstellung einer Injektionslösung	PT/H/0946/005	88420.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Methylprednisolon Hikma 250 mg Pulver zur Herstellung einer Injektionslösung	PT/H/0946/003	88419.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Methylprednisolon Hikma 250 mg Pulver zur Herstellung einer Injektionslösung	PT/H/0946/003	88419.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Methylprednisolone Hikma 1000 mg Prášek pro injekční roztok	PT/H/0946/005	56/476/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 1000 mg Prášek pro injekční roztok	PT/H/0946/005	56/476/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 125 mg Prášek pro injekční roztok	PT/H/0946/002	56/473/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 125 mg Prášek pro injekční roztok	PT/H/0946/002	56/473/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 250 mg Prášek pro injekční roztok	PT/H/0946/003	56/474/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 250 mg Prášek pro injekční roztok	PT/H/0946/003	56/474/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 40 mg Prášek pro injekcní roztok	PT/H/0946/001	56/472/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 40 mg Prášek pro injekční roztok	PT/H/0946/001	56/472/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ

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
Methylprednisolone Hikma 500 mg Prášek pro injekční roztok	PT/H/0946/004	56/475/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
Methylprednisolone Hikma 500 mg Prášek pro injekční roztok	PT/H/0946/004	56/475/13-C	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	CZ
methylprednisolone sodium succinate for injection	not available	PL 00057/1045	PFIZER LIMITED	UK
methylprednisolone sodium succinate for injection	not available	PL 00057/1047	PFIZER LIMITED	UK
methylprednisolone sodium succinate for injection	not available	PL 00057/1049	PFIZER LIMITED	UK
methylprednisolone sodium succinate for injection	not available	PL 00057/1048	PFIZER LIMITED	UK
methylprednisolone sodium succinate for injection	not available	PL 00057/1046	PFIZER LIMITED	UK
Metypred, 1000 mg, proszek do sporządzania roztworu do wstrzykiwań	FI/H/0797/004	22149	ORION CORPORATION	PL
Metypred, 125 mg, proszek do sporządzania roztworu do wstrzykiwań	FI/H/0797/001	22146	ORION CORPORATION	PL
Metypred, 250 mg, proszek do sporządzania roztworu do wstrzykiwań	FI/H/0797/002	22147	ORION CORPORATION	PL
Metypred, 500 mg, proszek do sporządzania roztworu do wstrzykiwań	FI/H/0797/003	22148	ORION CORPORATION	PL
Solomet depot 40 mg/ml injektioneste, suspensio	not available	9408	ORION OYJ	FI
SOLU MEDROL 1000 mg/16 ml polvere e solvente per soluzione iniettabile	not available	023202068	PFIZER ITALIA S.R.L.	IT
SOLU MEDROL 125 mg/2 ml polvere e solvente per soluzione iniettabile	not available	023202043	PFIZER 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
SOLU MEDROL 40 mg/ ml polvere e solvente per soluzione iniettabile	not available	023202017	PFIZER ITALIA S.R.L.	IT
SOLU MEDROL 500 mg/8 ml polvere e solvente per soluzione iniettabile	not available	023202056	PFIZER ITALIA S.R.L.	IT
SOLU-MEDROL	not available	41159/10/15-2-2012	PFIZER HELLAS, A.E.	GR
SOLU-MEDROL	not available	41158/10/15-2-2012	PFIZER HELLAS, A.E.	GR
SOLU-MEDROL	not available	41165/10/15-2-2012	PFIZER HELLAS, A.E.	GR
SOLU-MEDROL	not available	41163/10/15-2-2012	PFIZER HELLAS, A.E.	GR
SOLU-MEDROL 1 g injektiokuiva-aine ja liuotin, liuosta varten	not available	6397	PFIZER OY	FI
SOLU-MEDROL 1 g Prášok a rozpúšťadlo na injekčný roztok	not available	56/0045/75-S	PFIZER EUROPE MA EEIG	SK
Solu-Medrol 1 g pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	6397	PFIZER OY	FI
Solu-Medrol 1 g pulver och vätska till injektionsvätska, lösning	not available	09123-2	PFIZER AB	SE
Solu-Medrol 1 g pulver og væske til injeksjonsvæske, oppløsning	not available	06-4067	PFIZER AS	NO
SOLUMEDROL 1 g, poudre et solvant pour solution injectable	not available	34009 386 774 5 4	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 1 g, poudre et solvant pour solution injectable	not available	34009 386 773 9 3	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 1 g, poudre et solvant pour solution injectable	not available	34009 386 772 2 5	PFIZER HOLDING FRANCE	FR
SOLU-MEDROL 1000 mg milteliai ir tirpiklis injekciniam tirpalui	not available	LT/1/94/1936/004	PFIZER EUROPE MA EEIG	LT

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
Solu-Medrol 1000 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE062002	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 1000 mg por és oldószer oldatos injekcióhoz	not available	OGYI-T-2245/05	PFIZER KFT.	HU
Solu-Medrol 1000 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE062002	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 1000 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069834	PFIZER S.A. (BELGIUM)	LU
SOLU-MEDROL 1000 mg prašak i otapalo za otopinu za injekciju	not available	UP/I-530-09/12-02/42	PFIZER CROATIA D.O.O.	HR
Solu-Medrol 1000 mg prašek in vehikel za raztopino za injiciranje ali infundiranje	not available	H/93/01441/003	PFIZER LUXEMBOURG SARL	SI
Solu-Medrol 1000 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE062002	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 1000 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069834	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol 1000 mg pulveris un šķīdinātājs injekciju šķīduma pagatavošanai	not available	03-0513	PFIZER EUROPE MA EEIG	LV
Solu-Medrol 1000 mg/15,6 ml Pó e solvente para solução injetável	not available	8303834	LABORATÓRIOS PFIZER, 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
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 554 420 7 3	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 651 3 1	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 315 528 1 9	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 653 6 0	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 550198 8 6	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 120 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 337 477 0 1	PFIZER HOLDING FRANCE	FR
SOLU-MEDROL 125 mg injektiokuiva-aine ja liuotin, liuosta varten	not available	6397	PFIZER OY	FI
Solu-Medrol 125 mg por és oldószer oldatos injekcióhoz	not available	OGYI-T-2245/02	PFIZER KFT.	HU
SOLU-MEDROL 125 mg prašak i otapalo za otopinu za injekciju	not available	UP/I-530-09/12-02/39	PFIZER CROATIA D.O.O.	HR
Solu-Medrol 125 mg prašek in vehikel za raztopino za injiciranje ali infundiranje	not available	H/93/01441/001	PFIZER LUXEMBOURG SARL	SI
SOLU-MEDROL 125 mg Prášok a rozpúšťadlo na injekčný roztok	not available	56/0045/75-S	PFIZER EUROPE MA EEIG	SK
Solu-Medrol 125 mg pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	6397	PFIZER OY	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
Solu-Medrol 125 mg pulver och vätska till injektionsvätska, lösning	not available	09122	PFIZER AB	SE
Solu-Medrol 125 mg pulver og væske til injeksjonsvæske, oppløsning	not available	06-4065	PFIZER AS	NO
Solu-Medrol 125 mg pulveris un šķīdinātājs injekciju šķīduma pagatavošanai	not available	03-0510	PFIZER EUROPE MA EEIG	LV
Solu-Medrol 125 mg stungulyfsstofn og leysir, lausn	not available	731651	PFIZER APS	IS
Solu-Medrol 125 mg/2 ml Pó e solvente para solução injetável	not available	8303867	LABORATÓRIOS PFIZER, LDA.	PT
Solu-Medrol 125 mg/2 ml Pó e solvente para solução injetável	not available	8303818	LABORATÓRIOS PFIZER, LDA.	PT
SOLU-MEDROL 2 g injektiokuiva-aine ja liuotin, liuosta varten	not available	6397	PFIZER OY	FI
Solu-Medrol 2 g pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	6397	PFIZER OY	FI
Solu-Medrol 2 g pulver och vätska till injektionsvätska, lösning	not available	09123-3	PFIZER AB	SE
Solu-Medrol 2 g pulver og væske til injeksjonsvæske, oppløsning	not available	06-4068	PFIZER AS	NO
SOLUMEDROL 20 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 309 717 0 3	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 20 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 338 338 4 8	PFIZER HOLDING 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
SOLUMEDROL 20 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 553 945 9 4	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 20 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 647 6 9	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 20 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 648 2 0	PFIZER HOLDING FRANCE	FR
Solu-Medrol 2000 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE114362	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 2000 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE114362	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 2000 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE114362	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 2000 mg/31,2 ml Pó e solvente para solução injetável	not available	8303842	LABORATÓRIOS PFIZER, LDA.	PT
SOLU-MEDROL 250 mg injektiokuiva-aine ja liuotin, liuosta varten	not available	6397	PFIZER OY	FI
SOLU-MEDROL 250 mg milteliai ir tirpiklis injekciniam tirpalui	not available	LT/1/94/1936/002	PFIZER EUROPE MA EEIG	LT
Solu-Medrol 250 mg por és oldószer oldatos injekcióhoz	not available	OGYI-T-2245/03	PFIZER KFT.	HU
SOLU-MEDROL 250 mg prašak i otapalo za otopinu za injekciju	not available	UP/I-530-09/12-02/40	PFIZER CROATIA D.O.O.	HR

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
SOLU-MEDROL 250 mg Prášok a rozpúšťadlo na injekčný roztok	not available	56/0045/75-S	PFIZER EUROPE MA EEIG	SK
Solu-Medrol 250 mg pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	6397	PFIZER OY	FI
Solu-Medrol 250 mg pulveris un šķīdinātājs injekciju šķīduma pagatavošanai	not available	03-0511	PFIZER EUROPE MA EEIG	LV
SOLU-MEDROL 40 mg injektiokuiva-aine ja liuotin, liuosta varten	not available	5658	PFIZER OY	FI
SOLU-MEDROL 40 mg milteliai ir tirpiklis injekciniam tirpalui	not available	LT/1/94/1936/001	PFIZER EUROPE MA EEIG	LT
Solu-Medrol 40 mg por és oldószer oldatos injekcióhoz	not available	OGYI-T-2245/01	PFIZER KFT.	HU
SOLU-MEDROL 40 mg prašak i otapalo za otopinu za injekciju	not available	UP/I-530-09/12-02/38	PFIZER CROATIA D.O.O.	HR
Solu-Medrol 40 mg prašek in vehikel za raztopino za injiciranje ali infundiranje	not available	H/93/01441/004	PFIZER LUXEMBOURG SARL	SI
SOLU-MEDROL 40 mg Prášok a rozpúšťadlo na injekčný roztok	not available	56/0045/75-S	PFIZER EUROPE MA EEIG	SK
Solu-Medrol 40 mg pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	5658	PFIZER OY	FI
Solu-Medrol 40 mg pulver och vätska till injektionsvätska, lösning	not available	09408	PFIZER AB	SE
Solu-Medrol 40 mg pulver og væske til injeksjonsvæske, oppløsning	not available	5796	PFIZER AS	NO

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
Solu-Medrol 40 mg pulveris un šķīdinātājs injekciju šķīduma pagatavošanai	not available	03-0509	PFIZER EUROPE MA EEIG	LV
Solu-Medrol 40 mg stungulyfsstofn og leysir, lausn	not available	739001	PFIZER APS	IS
Solu-Medrol 40 mg/1 ml Pó e solvente para solução injetável	not available	8303859	LABORATÓRIOS PFIZER, LDA.	PT
Solu-Medrol 40 mg/1 ml Pó e solvente para solução injetável	not available	8303800	LABORATÓRIOS PFIZER, LDA.	PT
SOLUMEDROL 40 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 649 9 8	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 40 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 309 718 7 1	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 40 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 337 476 4 0	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 40 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 558 650 7 0	PFIZER HOLDING FRANCE	FR
SOLUMEDROL 40 mg/2 ml, lyophilisat et solution pour usage parentéral	not available	34009 553 947 1 6	PFIZER HOLDING FRANCE	FR
Solu-Medrol 40 mg/ml Prášek a rozpouštědlo pro injekční roztok	not available	56/045/75-A/C	PFIZER, SPOL. S R.O.	CZ
Solu-Medrol 40 mg/ml, poeder en oplosmiddel voor oplossing voor injectie	not available	RVG 05302	PFIZER B.V.	NL
SOLU-MEDROL 500 mg injektiokuiva-aine ja liuotin, liuosta varten	not available	6397	PFIZER OY	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
SOLU-MEDROL 500 mg milteliai ir tirpiklis injekciniam tirpalui	not available	LT/1/94/1936/003	PFIZER EUROPE MA EEIG	LT
Solu-Medrol 500 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE061993	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 500 mg por és oldószer oldatos injekcióhoz	not available	OGYI-T-2245/04	PFIZER KFT.	HU
Solu-Medrol 500 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE061993	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 500 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069833	PFIZER S.A. (BELGIUM)	LU
SOLU-MEDROL 500 mg prašak i otapalo za otopinu za injekciju	not available	UP/I-530-09/12-02/41	PFIZER CROATIA D.O.O.	HR
Solu-Medrol 500 mg prašek in vehikel za raztopino za injiciranje ali infundiranje	not available	H/93/01441/002	PFIZER LUXEMBOURG SARL	SI
SOLU-MEDROL 500 mg Prášok a rozpúšťadlo na injekčný roztok	not available	56/0045/75-S	PFIZER EUROPE MA EEIG	SK
Solu-Medrol 500 mg pulver och vätska till injektionsvätska, lösning metylprednisolon	not available	6397	PFIZER OY	FI
Solu-Medrol 500 mg pulver och vätska till injektionsvätska, lösning	not available	09123-1	PFIZER AB	SE
Solu-Medrol 500 mg pulver og væske til injeksjonsvæske, oppløsning	not available	06-4066	PFIZER AS	NO

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
Solu-Medrol 500 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE061993	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol 500 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069833	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol 500 mg pulveris un šķīdinātājs injekciju šķīduma pagatavošanai	not available	03-0512	PFIZER EUROPE MA EEIG	LV
Solu-Medrol 500 mg stungulyfsstofn og leysir, lausn	not available	731650	PFIZER APS	IS
SOLUMEDROL 500 mg, poudre pour solution injectable	not available	34009 386 777 4 4	PFIZER HOLDING FRANCE	FR
Solu-Medrol 500 mg/7,8 ml Pó e solvente para solução injetável	not available	8303826	LABORATÓRIOS PFIZER, LDA.	PT
Solu-Medrol 62,5 mg/ml Prášek a rozpouštědlo pro injekční roztok	not available	56/045/75-B/C	PFIZER, SPOL. S R.O.	CZ
Solu-Medrol 62,5 mg/ml Prášek a rozpouštědlo pro injekční roztok	not available	56/045/75-B/C	PFIZER, SPOL. S R.O.	CZ
Solu-Medrol 62,5 mg/ml Prášek a rozpouštědlo pro injekční roztok	not available	56/045/75-B/C	PFIZER, SPOL. S R.O.	CZ
Solu-Medrol 62,5 mg/ml Prášek a rozpouštědlo pro injekční roztok	not available	56/045/75-B/C	PFIZER, SPOL. S R.O.	CZ
Solu-Medrol 62,5 mg/ml, poeder en oplosmiddel voor oplossing voor injectie	not available	RVG 07041	PFIZER 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
Solu-Medrol Act-O-Vial 250 mg, liofilizat şi solvent pentru soluţie injectabilă	not available	7170/2006/01	PFIZER EUROPE MA EEIG	RO
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 1000 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE145232	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 1000 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE145232	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 125 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE133847	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 125 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE133847	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 125 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069836	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 2000 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE145223	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 2000 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE145223	PFIZER S.A. (BELGIUM)	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
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 40 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE133761	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 40 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE133761	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 40 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069835	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 500 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE145214	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) 500 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE145214	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 125 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE061747	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 125 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE061747	PFIZER S.A. (BELGIUM)	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
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 125 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069831	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 250 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE145205	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 250 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE145205	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 250 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069832	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 40 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	BE061582	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 40 mg Poeder en oplosmiddel voor oplossing voor injectie (methylprednisolon)	not available	BE061582	PFIZER S.A. (BELGIUM)	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
Solu-Medrol S.A.B. (= Sine Alcohol Benzylicus) Act-O-Vial 40 mg Poudre et solvant pour solution injectable (méthylprednisolone)	not available	2008069830	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 1000 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE145232	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE133847	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069836	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 2000 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE145223	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 40 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE133761	PFIZER S.A. (BELGIUM)	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
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 40 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069835	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) 500 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE145214	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE061747	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069831	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 250 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE145205	PFIZER S.A. (BELGIUM)	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
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 250 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069832	PFIZER S.A. (BELGIUM)	LU
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 40 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	BE061582	PFIZER S.A. (BELGIUM)	BE
Solu-Medrol S.A.B. (Sine Alcohol Benzylicus) Act-O-Vial 40 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung Methylprednisolon	not available	2008069830	PFIZER S.A. (BELGIUM)	LU
SOLU-MEDROL, 1000 mg süstelahuse pulber ja lahusti	not available	056894	PFIZER EUROPE MA EEIG	EE
SOLU-MEDROL, 1000 mg, proszek i rozpuszczalnik do sporządzania roztworu do wstrzykiwań	not available	R/2368	PFIZER EUROPE MA EEIG	PL
SOLU-MEDROL, 125 mg süstelahuse pulber ja lahusti	not available	056594	PFIZER EUROPE MA EEIG	EE
SOLU-MEDROL, 125 mg, proszek i rozpuszczalnik do sporządzania roztworu do wstrzykiwań	not available	R/2366	PFIZER EUROPE MA EEIG	PL
SOLU-MEDROL, 250 mg süstelahuse pulber ja lahusti	not available	056694	PFIZER EUROPE MA EEIG	EE

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
SOLU-MEDROL, 250 mg, proszek i rozpuszczalnik do sporządzania roztworu do wstrzykiwań	not available	R/3116	PFIZER EUROPE MA EEIG	PL
SOLU-MEDROL, 40 mg süstelahuse pulber ja lahusti	not available	056494	PFIZER EUROPE MA EEIG	EE
SOLU-MEDROL, 40 mg, proszek i rozpuszczalnik do sporządzania roztworu do wstrzykiwań	not available	R/1553	PFIZER EUROPE MA EEIG	PL
SOLU-MEDROL, 500 mg süstelahuse pulber ja lahusti	not available	056794	PFIZER EUROPE MA EEIG	EE
SOLU-MEDROL, 500 mg, proszek i rozpuszczalnik do sporządzania roztworu do wstrzykiwań	not available	R/2367	PFIZER EUROPE MA EEIG	PL
Solu-Medrol, poeder voor oplossing voor infusie, 1000 mg	not available	RVG 06664	PFIZER B.V.	NL
Solu-Medrol, poeder voor oplossing voor infusie, 500 mg	not available	RVG 06664	PFIZER B.V.	NL
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	6132	PFIZER APS	DK
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	6613	PFIZER APS	DK
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	6133	PFIZER APS	DK
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	5496	PFIZER APS	DK
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	5496	PFIZER APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solu-Medrol, pulver og solvens til injektionsvæske, opløsning	not available	6132	PFIZER APS	DK
Solu-Medrol® 1000 mg – Trockenstechampulle mit Lösungsmittel	not available	16.218	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Solu-Medrol® 500 mg – Trockenstechampulle mit Lösungsmittel	not available	16.217	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Solu-Medrone 1 gram	not available	PL 00057/1048	PFIZER LIMITED	UK
Solu-Medrone 125 mg	not available	PL 00057/1046	PFIZER LIMITED	UK
Solu-Medrone 2 gram	not available	PL 00057/1049	PFIZER LIMITED	UK
Solu-Medrone 40 mg	not available	PL 00057/1045	PFIZER LIMITED	UK
Solu-Medrone 500 mg	not available	PL 00057/1047	PFIZER LIMITED	UK
Solu-Medrone powder and solvent for solution for injection or concentrate for solution for infusion 40 mg/vial	not available	PA 0822/136/001	PFIZER HEALTHCARE IRELAND	IE
Solu-Medrone powder and solvent for solution for injection or concentrate for solution for infusion 500 mg/vial	not available	PA 0822/136/003	PFIZER HEALTHCARE IRELAND	IE
Solu-Medrone powder and solvent for solution for injection or concentrate for solution for infusion 1000 mg/vial	not available	PA 0822/136/004	PFIZER HEALTHCARE IRELAND	IE
Solu-Medrone powder and solvent for solution for injection or concentrate for solution for infusion 125 mg/vial	not available	PA 822/136/002	PFIZER HEALTHCARE IRELAND	IE
Solu-Moderín 1 g polvo y disolvente para solución inyectable	not available	53.203	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
Solu-Moderín 125 mg polvo y disolvente para solución inyectable	not available	49.768	PFIZER, S.L.	ES
Solu-Moderín 40 mg polvo y disolvente para solución inyectable	not available	49.767	PFIZER, S.L.	ES
Solu-Moderín 500 mg polvo y disolvente para solución inyectable	not available	53.202	PFIZER, S.L.	ES
URBASON	not available	024001012	SANOFI SPA	IT
URBASON 16 MG COMPRIMIDOS	not available	59123	SANOFI-AVENTIS, S.A.	ES
URBASON 16 MG COMPRIMIDOS	not available	59123	SANOFI-AVENTIS, S.A.	ES
Urbason 20 mg polvo y disolvente para solución inyectable	not available	34022	SANOFI-AVENTIS, S.A.	ES
Urbason 20 mg polvo y disolvente para solución inyectable	not available	34022	SANOFI-AVENTIS, S.A.	ES
Urbason 250 mg polvo y disolvente para solución inyectable o para perfusión	not available	50537	SANOFI-AVENTIS, S.A.	ES
Urbason 250 mg polvo y disolvente para solución inyectable o para perfusión	not available	50537	SANOFI-AVENTIS, S.A.	ES
Urbason 250 mg polvo y disolvente para solución inyectable o para perfusión	not available	50537	SANOFI-AVENTIS, S.A.	ES
URBASON 4 MG COMPRIMIDOS	not available	32729	SANOFI-AVENTIS, S.A.	ES
URBASON 4 MG COMPRIMIDOS	not available	32729	SANOFI-AVENTIS, S.A.	ES
URBASON 4 MG TABLETTEN	not available	10.807	SANOFI-AVENTIS GMBH OSTERREICH	AT
URBASON 4 MG TABLETTEN	not available	10.807	SANOFI-AVENTIS GMBH OSTERREICH	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
URBASON 40 MG COMPRIMIDOS	not available	59124	SANOFI-AVENTIS, S.A.	ES
URBASON 40 MG POLVO Y DISOLVENTE PARA SOLUCION INYECTABLE	not available	34023	SANOFI-AVENTIS, S.A.	ES
URBASON 40 MG POLVO Y DISOLVENTE PARA SOLUCION INYECTABLE	not available	34023	SANOFI-AVENTIS, S.A.	ES
URBASON 40 MG TABLETTEN	not available	12.208	SANOFI-AVENTIS GMBH OSTERREICH	AT
URBASON 40 MG TABLETTEN	not available	12.208	SANOFI-AVENTIS GMBH OSTERREICH	AT
Urbason 8 mg polvo y disolvente para solución inyectable	not available	34021	SANOFI-AVENTIS, S.A.	ES
Urbason 8 mg polvo y disolvente para solución inyectable	not available	34021	SANOFI-AVENTIS, S.A.	ES
URBASON SOLUBILE	not available	018259022	SANOFI SPA	IT
URBASON SOLUBILE	not available	018259034	SANOFI SPA	IT
URBASON SOLUBILE	not available	018259059	SANOFI SPA	IT
Urbason solubile 1000 mg Trockenstechampulle mit Lösungsmittel	not available	16.419	SANOFI-AVENTIS GMBH OSTERREICH	AT
URBASON SOLUBILE 16 MG TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	11.404	SANOFI-AVENTIS GMBH OSTERREICH	AT
URBASON SOLUBILE 16 MG TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	11.404	SANOFI-AVENTIS GMBH OSTERREICH	AT
Urbason solubile 250 mg Trockenampullen mit Lösungsmittel	not available	15.040	SANOFI-AVENTIS GMBH OSTERREICH	AT
URBASON SOLUBILE 32 MG TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	11.405	SANOFI-AVENTIS GMBH OSTERREICH	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
URBASON SOLUBILE 32 MG TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	11.405	SANOFI-AVENTIS GMBH OSTERREICH	AT
Адвантан 0,1 % крем	not available	20010606	BAYER AG	BG
Адвантан 0,1 % маз	not available	20010603	BAYER AG	BG
Адвантан Мляко 0,1% емулсия за кожа	not available	20040539	BAYER AG	BG
Депо-Медрол 40 mg/ml инжекционна суспензия	not available	20040391	PFIZER ENTERPRISES SARL	BG
Медрол 4 mg таблетки	not available	20050452	PFIZER ENTERPRISES SARL	BG
Солу-Медрол 1000 mg прах и разтворител за инжекционен разтвор	not available	20040386	PFIZER ENTERPRISES SARL	BG
Солу-Медрол 125 mg прах и разтворител за инжекционен разтвор	not available	20040383	PFIZER ENTERPRISES SARL	BG
Солу-Медрол 2000 mg прах и разтворител за инжекционен разтвор	not available	20040387	PFIZER ENTERPRISES SARL	BG
Солу-Медрол 40 mg прах и разтворител за инжекционен разтвор	not available	20040382	PFIZER ENTERPRISES SARL	BG
Metilprednisolona Reig Jofre 1 mg/g emulsão cutânea	not available	5703772	LABORATORIO REIG JOFRE, S.A.	PT
Metilprednisolona Reig Jofre 1mg/g crème	not available	5703830	LABORATORIO REIG JOFRE, S.A.	PT
Metilprednisolona Reig Jofre 1mg/g creme	not available	5703822	LABORATORIO REIG JOFRE, S.A.	PT