24 July 2019
EMA/480512/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: azathioprine

Procedure no.: PSUSA/00000275/201812

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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
Azathioprine Tablets 50 mg	not available	PL 13606/0093	STRIDES PHARMA UK LIMITED	UK
Imuran 50 mg	not available	59/0168/13-S	ASPEN PHARMA TRADING LIMITED	SK
Imuran 25 mg	not available	59/0188/70-CS	ASPEN PHARMA TRADING LIMITED	SK
Aza-Q® 50 mg Tabletten	not available	35480.00.00	JUTA PHARMA GMBH	DE
AZATHIOPRINE/FARMASYN, Επικαλυμμένα με λεπτό υμένιο δισκία, 50 mg	not available	29043/5-4-2016	FARMASYN S.A.	GR
Immufalk 75 mg comprimidos recubiertos con pelicula	DE/H/5684/002	78165	DR FALK PHARMA GMBH	ES
Azafalk 75 mg Filmtabletten	DE/H/5684/002	1-31700	DR FALK PHARMA GMBH	AT
Azafalk 75 mg Filmtabletten	DE/H/5684/002	81859.00.00	DR FALK PHARMA GMBH	DE
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/006	DR FALK PHARMA GMBH	LT
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/005	DR FALK PHARMA GMBH	LT
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/001	DR FALK PHARMA GMBH	LT
Azafalk 75 mg filmomhulde tabletten	DE/H/5684/002	RVG 107494	DR FALK PHARMA GMBH	NL
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/002	DR FALK PHARMA GMBH	LT
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/004	DR FALK PHARMA GMBH	LT
Azafalk 75 mg plėvele dengtos tabletės	DE/H/5684/002	LT/1/12/3074/003	DR FALK PHARMA GMBH	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
Azafalk 75mg film-coated tablets	DE/H/5684/002	PL 08637/0023	DR FALK PHARMA GMBH	UK
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/007	DR FALK PHARMA GMBH	SI
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/008	DR FALK PHARMA GMBH	SI
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/009	DR FALK PHARMA GMBH	SI
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/010	DR FALK PHARMA GMBH	SI
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/011	DR FALK PHARMA GMBH	SI
Azafalk 75 mg filmsko obložene tablete	DE/H/5684/002	H/12/00244/012	DR FALK PHARMA GMBH	SI
Azafalk 75 mg comprimidos revestidos por pelicula	DE/H/5684/002	5423876	DR FALK PHARMA GMBH	PT
Azafalk 75 mg comprimidos revestidos por pelicula	DE/H/5684/002	5423868	DR FALK PHARMA GMBH	PT
Azathioprin ”ratiopharm”, filmovertrukne tabletter	DK/H/0164/001	30549	RATIOPHARM GMBH	DK
Azathioprin-ratiopharm® 50 mg Filmtabletten	DK/H/0164/002	47311.01.00	RATIOPHARM GMBH	DE
Azathioprin ”ratiopharm”, filmovertrukne tabletter	DK/H/0164/002	30550	RATIOPHARM GMBH	DK
Azathioprin-ratiopharm® 25 mg Filmtabletten	DK/H/0164/001	47311.00.00	RATIOPHARM GMBH	DE
IMUREL 25 mg, comprimé pelliculé	not available	34009 364 145 5 6	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 25 mg, comprimé pelliculé	not available	34009 364 142 6 6	ASPEN PHARMA	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
			TRADING LIMITED	
IMUREL 25 mg, comprimé pelliculé	not available	34009 364 143 2 7	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 25 mg, comprimé pelliculé	not available	34009 364 144 9 5	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 50 mg, comprimé pelliculé	not available	34009 364 149 0 7	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 50 mg, comprimé pelliculé	not available	34009 364 146 1 7	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 50 mg, comprimé pelliculé	not available	34009 364 147 8 5	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 50 mg, comprimé pelliculé	not available	34009 364 148 4 6	ASPEN PHARMA TRADING LIMITED	FR
IMUREL 50 mg, poudre pour solution injectable (IV)	not available	34009 561 936 5 3	ASPEN PHARMA TRADING LIMITED	FR
Azathioprin “Orifarm", filmovertrukne tabletter	DK/H/0650/001	35547	ORIFARM GENERICS A/S	DK
Azathioprin Orifarm 50 mg filmdragerad tablett	DK/H/0650/001	22455	ORIFARM GENERICS A/S	SE
Azathioprine Teva 25 mg, filmomhulde tabletten	not available	RVG 106268	TEVA NEDERLAND B.V.	NL
IMURAN mitis 25 mg, comprimés pelliculés	not available	BE 151636	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, Filmtabletten	not available	BE 058177	ASPEN PHARMA TRADING LIMITED	BE
IMURAN mitis 25 mg, Filmtabletten	not available	BE 151636	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, Pulver zur Herstellung	not available	BE 095015	ASPEN PHARMA	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
einer Injektionslösung			TRADING LIMITED	
IMURAN mitis 25 mg, filmomhulde tabletten	not available	BE 151636	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, poudre pour solution injectable	not available	BE 095015	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, poeder voor oplossing voor injectie	not available	BE 095015	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, comprimés pelliculés	not available	BE 058177	ASPEN PHARMA TRADING LIMITED	BE
IMURAN 50 mg, filmomhulde tabletten	not available	BE 058177	ASPEN PHARMA TRADING LIMITED	BE
Azathioprin Heumann 50 mg Filmtabletten	DE/H/5059/001	63079.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
AZATHIOPRINE EG 50 MG, COMPRIME PELLICULE	not available	NL33424	EG LABO LABORATOIRES EUROGENERICS	FR
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/007	DR FALK PHARMA GMBH	LT
Immufalk 100mg comprimidos recubiertos con pelicula	DE/H/5684/003	78164	DR FALK PHARMA GMBH	ES
Azafalk 100 mg Filmtabletten	DE/H/5684/003	81860.00.00	DR FALK PHARMA GMBH	DE
Azafalk 100 mg Filmtabletten	DE/H/5684/003	1-31701	DR FALK PHARMA GMBH	AT
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/008	DR FALK PHARMA GMBH	LT
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/009	DR FALK PHARMA GMBH	LT
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/010	DR FALK PHARMA GMBH	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
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/011	DR FALK PHARMA GMBH	LT
Azafalk 100 mg plėvele dengtos tabletės	DE/H/5684/003	LT/1/12/3074/012	DR FALK PHARMA GMBH	LT
Azafalk 100mg film-coated tablets	DE/H/5684/003	PL 08637/0024	DR FALK PHARMA GMBH	UK
Azafalk 100 mg comprimidos revestidos por pelicula	DE/H/5684/003	5423900	DR FALK PHARMA GMBH	PT
Azafalk 100 mg filmomhulde tabletten	DE/H/5684/003	RVG 107495	DR FALK PHARMA GMBH	NL
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/013	DR FALK PHARMA GMBH	SI
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/014	DR FALK PHARMA GMBH	SI
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/015	DR FALK PHARMA GMBH	SI
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/016	DR FALK PHARMA GMBH	SI
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/017	DR FALK PHARMA GMBH	SI
Azafalk 100 mg filmsko obložene tablete	DE/H/5684/003	H/12/00244/018	DR FALK PHARMA GMBH	SI
Imuran Tablets 25 mg	not available	PL 39699/0004	ASPEN PHARMA TRADING LIMITED	UK
Imuran Tablets 50mg	not available	PL 39699/0005	ASPEN PHARMA TRADING LIMITED	UK
Imuran Injection	not available	PL 39699/0003	ASPEN PHARMA TRADING LIMITED	UK
AZATIOPRINA ASPEN 50 mg compresse rivestite con film	not available	020957039	ASPEN PHARMA TRADING LIMITED	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
IMURAN 50 mg, comprimés pelliculés	not available	2008049750	ASPEN PHARMA TRADING LIMITED	LU
IMURAN 50 mg, Filmtabletten	not available	2008049750	ASPEN PHARMA TRADING LIMITED	LU
IMURAN mitis 25 mg, comprimés pelliculés	not available	2008049749	ASPEN PHARMA TRADING LIMITED	LU
IMURAN mitis 25 mg, Filmtabletten	not available	2008049749	ASPEN PHARMA TRADING LIMITED	LU
IMURAN 50 mg, Pulver zur Herstellung einer Injektionslösung	not available	2008049751	ASPEN PHARMA TRADING LIMITED	LU
IMURAN 50 mg, poudre pour solution injectable	not available	2008049751	ASPEN PHARMA TRADING LIMITED	LU
IMURAN 25 mg comprimidos revestidos por película	not available	8218313	ASPEN PHARMA TRADING LIMITED	PT
IMURAN 50 mg comprimidos revestidos por película	not available	8218305	ASPEN PHARMA TRADING LIMITED	PT
Imuger 25 mg film-coated tablets	DK/H/0146/001	PA0577/032/001	MCDERMOTT LABORATORIES LTD	IE
Azathioprine Mylan 25 mg, filmomhulde tabletten	DK/H/0146/001	RVG 24721	MYLAN B.V.	NL
Azatioprin "Mylan", filmovertrukne tabletter	DK/H/0146/001	13501	MYLAN AB	DK
AZATHIOPRINE MYLAN 50 mg, comprimé pelliculé sécable	DK/H/0146/002	NL 25086	MYLAN S.A.S	FR
Imuger 50 mg film-coated tablets	DK/H/0146/002	PA0577/032/002	MCDERMOTT LABORATORIES LTD	IE
Azathioprine Mylan 50 mg, filmomhulde tabletten	DK/H/0146/002	RVG 24722	MYLAN 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
Azatioprin "Mylan", filmovertrukne tabletter	DK/H/0146/002	12530	MYLAN AB	DK
Azafalk 25mg Filmtabletten	DK/H/0146/001/MR	47309.00.00	DR FALK PHARMA GMBH	DE
Azafalk 50mg Filmtabletten	DK/H/0146/002/MR	47309.01.00	DR FALK PHARMA GMBH	DE
IMURAN 50 mg apvalkotās tabletes	not available	98-0510	ASPEN PHARMA TRADING LIMITED	LV
Azatioprin Mylan 50 mg tabletter	not available	10428	MYLAN AB	SE
Azatioprin Mylan 25 mg tabletter	not available	11073	MYLAN AB	SE
Imuran 50 mg, filmomhulde tablet	not available	RVG 05565	ASPEN PHARMA TRADING LIMITED	NL
Imuran 25 mg, filmomhulde tablet	not available	RVG 12476	ASPEN PHARMA TRADING LIMITED	NL
Imuran Film-coated Tablets 25 mg	not available	PA 1691/3/2	ASPEN PHARMA TRADING LIMITED	IE
Imuran Film-coated Tablets 50 mg	not available	PA 1691/3/3	ASPEN PHARMA TRADING LIMITED	IE
Imuran Powder for Solution for Injection or Infusion 50 mg	not available	PA 1691/3/1	ASPEN PHARMA TRADING LIMITED	IE
Azathioprin - neuraxpharm 25 mg Filmtabletten	not available	97260.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Azathioprin - neuraxpharm 50 mg Filmtabletten	not available	97261.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Azathioprine Tablets 25 mg	not available	PL 04569/0234	GENERICS [UK] LIMITED	UK
Azathioprine Tablets 50 mg	not available	PL 04569/0073	GENERICS [UK] LIMITED	UK
Azathioprine 1A Pharma 25 mg, filmomhulde tabletten	NL/H/0326/001	RVG 24581	1 A PHARMA GMBH	NL
AZATHIOPRINE 1A Pharma 50 MG, filmomhulde tabletten	NL/H/0326/002	RVG 24582	1 A PHARMA GMBH	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
Azathioprin Hexal 50 mg - Filmtabletten	NL/H/0326/002	1-24656	HEXAL PHARMA GMBH	AT
Azathioprine Sandoz 50 mg filmomhulde tabletten	NL/H/0326/002	BE242584	SANDOZ N.V.	BE
Azathioprine Sandoz 50 mg filmomhulde tabletten	NL/H/0326/002	BE277751	SANDOZ N.V.	BE
Azathioprin HEXAL® 50 mg Filmtabletten	NL/H/0326/002	35481.00.01	HEXAL AG	DE
Azathioprin HEXAL® 25 mg Filmtabletten	NL/H/0326/001	35481.01.01	HEXAL AG	DE
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292011	SANDOZ S.P.A.	IT
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292050	SANDOZ S.P.A.	IT
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292047	SANDOZ S.P.A.	IT
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292062	SANDOZ S.P.A.	IT
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292035	SANDOZ S.P.A.	IT
AZATIOPRINA HEXAL 50 mg compresse rivestite con film	NL/H/0326/002	036292023	SANDOZ S.P.A.	IT
Azathioprine 25mg Tablets	NL/H/0326/001	PL 04416/1247	SANDOZ LTD	UK
Azathioprine 50mg Tablets	NL/H/0326/002	PL 04416/1248	SANDOZ LTD	UK
Imuprin 50 mg επικαλυμμένα με υμένιο δισκία	not available	9030	REMEDICA LTD	CY
Imurel 50 mg tabletter	not available	5080	ASPEN PHARMA TRADING LIMITED	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
IMURAN 50 mg comprimate filmate	not available	10042/2017/01	ASPEN PHARMA TRADING LIMITED	RO
Imurel 25 mg tabletter	not available	7259	ASPEN PHARMA TRADING LIMITED	NO
Immunoprin 50 mg	DK/H/0843/001	44/0155/06-S	SANDOZ PHARMACEUTICALS D.D.	SK
Immunoprin 50 mg - Filmtabletten	DK/H/0843/001	1-26374	EBEWE PHARMA	AT
Azathioprin "Ebewe"	DK/H/0843/001	35919	EBEWE PHARMA	DK
Azathioprin Ebewe 50 mg filmtabletta	DK/H/0843/001	OGYI-T-20277/01	SANDOZ HUNGÁRIA KFT	HU
Azathioprin Ebewe 50 mg filmtabletta	DK/H/0843/001	OGYI-T-20277/02	SANDOZ HUNGÁRIA KFT	HU
Imuran 50 mg plėvele dengtos tabletės	not available	LT/1/94/1856/001	ASPEN PHARMA TRADING LIMITED	LT
Azathioprin - 1 A Pharma® 25 mg Filmtabletten	NL/H/0328/001	54697.00.00	1 A PHARMA GMBH	DE
Azathioprin - 1 A Pharma® 50 mg Filmtabletten	NL/H/0328/002	54697.01.00	1 A PHARMA GMBH	DE
Azathioprine 25 mg, filmomhulde tablet	NL/H/0328/001	RVG 27564	HEXAL AG	NL
Azathioprine 50 mg, filmomhulde tablet	NL/H/0328/002	RVG 27565	HEXAL AG	NL
Azaqvida 25 mg Filmtabletten	not available	97609.00.00	AQVIDA GMBH	DE
Azaqvida 50 mg Filmtabletten	not available	97610.00.00	AQVIDA GMBH	DE
Imurek 25 mg Filmtabletten	not available	6101735.01.00	ASPEN PHARMA TRADING LIMITED	DE
Imurek 50 mg Filmtabletten	not available	6101735.00.00	ASPEN PHARMA TRADING LIMITED	DE
Imurek i.v. 50 mg Pulver zur Herstellung einer Infusions- oder	not available	6101758.00.00	ASPEN PHARMA TRADING LIMITED	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
Injektionslösung				
IMURAN 50 mg comprimate filmate	not available	10042/2017/01	ASPEN PHARMA TRADING LIMITED	RO
ИМУРАН 50 mg филмирани таблетки	not available	20020842	ASPEN PHARMA TRADING LIMITED	BG
Azathioprin Heumann 25 mg Filmtabletten	DE/H/4215/001/DC	93348.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Azathioprin Heumann 75 mg Filmtabletten	DE/H/4215/002/DC	93349.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Azathioprin Heumann 100 mg Filmtabletten	DE/H/4215/003	93350.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
AZA-effect pharma 50 mg Filmtabletten	not available	97853.00.00	EFFECT PHARMA GMBH	DE
Imurel	not available	870108	ASPEN PHARMA TRADING LIMITED	IS
Imurel	not available	660887	ASPEN PHARMA TRADING LIMITED	IS
AZATHIOPRINE SANDOZ 25, FILMOMHULDE TABLETTEN 25 MG	NL/H/0327/001	RVG 27562	SANDOZ B.V.	NL
AZATHIOPRINE SANDOZ 50, FILMOMHULDE TABLETTEN 50 MG	NL/H/0327/002	RVG 27563	SANDOZ B.V.	NL
Azathioprine 50 mg Tablets	DK/H/0840/001	PL 30306/0520	ACTAVIS GROUP PTC EHF.	UK
AZAFOR 50mg compresse rivestite con film	IT/H/0273/001	037534017	SOFAR S.P.A.	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
AZAFOR 50mg compresse rivestite con film	IT/H/0273/001	037534029	SOFAR S.P.A.	IT
Imuran 25 mg	not available	59/188/70-A/C	ASPEN PHARMA TRADING LIMITED	CZ
Imuran 50 mg	not available	59/188/70-B/C	ASPEN PHARMA TRADING LIMITED	CZ
AZAMUN® 25 mg, filmdragerade tabletter	not available	11021	TAKEDA OY	FI
AZAMUN® 50 mg, filmdragerade tabletter	not available	10104	TAKEDA OY	FI
Azamun 50 mg -tabletti, kalvopäällysteinen	not available	10104	TAKEDA OY	FI
Azamun 25 mg -tabletti, kalvopäällysteinen	not available	11021	TAKEDA OY	FI
Azathioprin beta 50 mg Filmtabletten	DE/H/3128/001	54699.01.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Imuprin 50 mg film-coated tablets	not available	MA084/01101	REMEDICA LTD	MT
Azathioprine 50mg Film-coated Tablets	not available	PL 00530/0647	NORTON HEALTHCARE LTD T/A IVAX PHARMACEUTICALS UK	UK
Azathioprine 25mg Film-coated Tablets	not available	PL 00530/0646	NORTON HEALTHCARE LTD T/A IVAX PHARMACEUTICALS UK	UK
Azathioprin ”Actavis”, filmovertrukne tabletter	DK/H/0655/001	35598	ACTAVIS GROUP PTC EHF.	DK
Azathioprin Actavis 50 mg filmdragerade tabletter	DK/H/0655/001	22463	ACTAVIS NORDIC A/S	SE
Azathioprin AL 25 mg Filmtabletten	not available	91115.00.00	ALIUD PHARMA 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
Azathioprin AL 50 mg Filmtabletten	not available	91116.00.00	ALIUD PHARMA GMBH	DE
Imurek-50 mg Filmtabletten	not available	13.362	ASPEN PHARMA TRADING LIMITED	AT
Azatioprina medac 25 mg comprimidos revestidos por película	DE/H/3039/001	5589619	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Azatioprina medac 50 mg comprimidos revestidos por película	DE/H/3039/002	5589627	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Azathioprine medac 50 mg filmdragerade tabletter	DE/H/3039/002	45353	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SE
Azamedac 25 mg Filmtabletten	DE/H/3039/001	83625.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Azamedac 50 mg Filmtabletten	DE/H/3039/002	83626.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Azathioprin medac 25 mg potahované tablety	DE/H/3039/001	59/242/13-C	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	CZ
Azathioprin medac 50 mg potahované tablety	DE/H/3039/002	59/243/13-C	MEDAC GESELLSCHAFT FÜR KLINISCHE	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
			SPEZIALPRÄPARATE MBH (WEDEL)	
Azatioprin medac 50 mg tabletter, filmdrasjerte	DE/H/3039/002	10-8007	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	NO
Azathioprin medac 50 mg tabletit, kalvopäällysteiset	DE/H/3039/002	29294	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Azathioprin medac 25 mg tabletit, kalvopäällysteiset	DE/H/3039/001	29293	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Azathioprin medac 25 mg filmom obalené tablety	DE/H/3039/001	59/0250/13-S	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SK
Azathioprin medac 50 mg filmom obalené tablety	DE/H/3039/002	59/0251/13-S	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SK
Azatioprina medac 25 mg comprimidos revestidos por película	DE/H/3039/001	DE/H/3039/001	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Azatioprina medac 50 mg comprimidos revestidos por película	DE/H/3039/002	DE/H/3039/002	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH	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
			(WEDEL)	
Azathioprin medac 50 mg filmdragerade tabletter	DE/H/3039/002	29294	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Imuran 25 mg filmtabletta	not available	OGYI-T-665/02	ASPEN PHARMA TRADING LIMITED	HU
Imuran 50 mg filmtabletta	not available	OGYI-T-665/01	ASPEN PHARMA TRADING LIMITED	HU
IMURAN, 50 mg õhukese polümeerikattega tabletid	not available	101495	ASPEN PHARMA TRADING LIMITED	EE
Azapress Tablets 50 mg	not available	PL 40147/0009	ENNOGEN PHARMA LIMITED	UK
Azathioprine Tablets 50 mg	not available	PL 40147/0009	ENNOGEN PHARMA LIMITED	UK
Azaqvida 50 mg Filmtabletten	not available	98242.00.00	AQVIDA GMBH	DE
Imuran 50 mg filmom obložene tablete	not available	HR-H-628872163	ASPEN PHARMA TRADING LIMITED	HR
Imuran Film-coated Tablets 25 mg	not available	PA 1691/3/2	ASPEN PHARMA TRADING LIMITED	IE
Imuran Film-coated Tablets 50 mg	not available	PA 1691/3/3	ASPEN PHARMA TRADING LIMITED	IE
Imuran Powder for Solution for Injection or Infusion 50 mg	not available	PA 1691/3/1	ASPEN PHARMA TRADING LIMITED	IE
IMURAN 50 mg filmsko obložene tablete	not available	H/93/00772/001	ASPEN PHARMA TRADING LIMITED	SI
Azathioprine 25 mg film-coated tablets	not available	PL 11311/0475	TILLOMED LABORATORIES LTD	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
Azathioprine 50 mg film-coated tablets	not available	PL 11311/0476	TILLOMED LABORATORIES LTD	UK
Imurel 50 mg comprimidos recubiertos con película	not available	50.043	UCB PHARMA S.A. (MADRID ES)	ES
Imurel 50 mg polvo para solución inyectable	not available	51.003	UCB PHARMA S.A. (MADRID ES)	ES
Azathioprine 50 PCH, tabletten 50 mg.	not available	RVG 10467	PHARMACHEMIE B.V	NL
Imurel 50 mg filmdragerad tablett	not available	8204	ASPEN PHARMA TRADING LIMITED	SE
Imurel 25 mg filmdragerad tablett	not available	10577	ASPEN PHARMA TRADING LIMITED	SE
Imurel, tabletter, filmovertrukne	not available	12943	ASPEN PHARMA TRADING LIMITED	DK
Imurel, tabletter, filmovertrukne	not available	05209	ASPEN PHARMA TRADING LIMITED	DK
Azathioprine Teva 25 mg, filmomhulde tabletten	not available	RVG 106268	TEVA NEDERLAND B.V.	NL
Imuran, 25 mg, tabletki powlekane	not available	R/1442	ASPEN PHARMA TRADING LIMITED	PL
Imuran, 50 mg, tabletki powlekane	not available	R/2778	ASPEN PHARMA TRADING LIMITED	PL
AZATHIOPRINE/PHARMACHEMIE 50 mg δισκία	not available	88576/15/30	CHEMIPHARM S.G. DE TCHAVES & CIE E.E	GR
Immunoprin 75 mg õhukese polümeerikattega tabletid	AT/H/0270/001	710310	SANDOZ PHARMACEUTICALS D.D.	EE
Immunoprin 100 mg õhukese polümeerikattega tabletid	AT/H/0270/002	710510	SANDOZ PHARMACEUTICALS D.D.	EE
Immunoprin 100 mg, filmom obalené	AT/H/0270/002	59/0714/10-S	SANDOZ	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
tablety			PHARMACEUTICALS D.D.	
IMMUNOPRIN 100 mg filmsko obložene tablete	AT/H/0270/002	H/11/00771/003	EBEWE PHARMA	SI
IMMUNOPRIN 100 mg filmsko obložene tablete	AT/H/0270/002	H/11/00771/004	EBEWE PHARMA	SI
IMMUNOPRIN 75 mg filmsko obložene tablete	AT/H/0270/001	H/11/00771/001	EBEWE PHARMA	SI
IMMUNOPRIN 75 mg filmsko obložene tablete	AT/H/0270/001	H/11/00771/002	EBEWE PHARMA	SI
Immunoprin 100 mg Filmtabletten	AT/H/0270/002	1-29708	EBEWE PHARMA	AT
Immunoprin 75 mg Filmtabletten	AT/H/0270/001	1-29707	EBEWE PHARMA	AT
Azathioprin Sandoz 100mg filmomhulde tabletten	AT/H/0270/002	BE383887	SANDOZ N.V.	BE
Azathioprin HEXAL 75 mg Filmtabletten	AT/H/0270/001	76504.00.00	HEXAL AG	DE
Azathioprin HEXAL 100 mg Filmtabletten	AT/H/0270/002	76505.00.00	HEXAL AG	DE
Immunoprin 75 mg filmdragerad tablett	AT/H/0270/001	41673	EBEWE PHARMA	SE
Immunoprin 100 mg filmdragerad tablett	AT/H/0270/002	41674	EBEWE PHARMA	SE
ИМУНОПРИН 100 MG ФИЛМИРАНИ ТАБЛЕТКИ	AT/H/0270/002	20120356	EBEWE PHARMA	BG
ИМУНОПРИН 75 MG ФИЛМИРАНИ ТАБЛЕТКИ	AT/H/0270/001	20120355	EBEWE PHARMA	BG
AZATHIOPRINE VIS, 50 mg, tabletki	not available	R/2328	ZAKŁADY CHEMICZNO-FARMACEUTYCZNE "VIS" SPÓŁKA Z O. O.	PL

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
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/013	DR FALK PHARMA GMBH	LT
Azafalk 50 mg Filmtabletten	DE/H/5684/004	135590	DR FALK PHARMA GMBH	AT
Azafalk 50 mg Filmtabletten	DE/H/5684/004	90364.00.00	DR FALK PHARMA GMBH	DE
Azafalk 50mg film-coated tablets	DE/H/5684/004	PL 08637/0026	DR FALK PHARMA GMBH	UK
Azalfalk 50 mg comprimidos revestidos por Película	DE/H/5684/004	5603550	DR FALK PHARMA GMBH	PT
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/014	DR FALK PHARMA GMBH	LT
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/015	DR FALK PHARMA GMBH	LT
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/016	DR FALK PHARMA GMBH	LT
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/017	DR FALK PHARMA GMBH	LT
Azafalk 50 mg plėvele dengtos tabletės	DE/H/5684/004	LT/1/12/3074/018	DR FALK PHARMA GMBH	LT
Azafalk 50 mg filmomhulde tabletten	DE/H/5684/004	RVG 113802	DR FALK PHARMA GMBH	NL
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/001	DR FALK PHARMA GMBH	SI
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/002	DR FALK PHARMA GMBH	SI
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/003	DR FALK PHARMA GMBH	SI
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/004	DR FALK PHARMA GMBH	SI
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/005	DR FALK PHARMA GMBH	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
Azafalk 50 mg filmsko obložene tablete	DE/H/5684/004	H/12/00244/006	DR FALK PHARMA GMBH	SI
Azalfalk 50 mg comprimidos revestidos por Película	DE/H/5684/004	5603543	DR FALK PHARMA GMBH	PT
Azathioprine 50mg film coated tablets	UK/H/0934/001	PL 20395/0066	RELON CHEM LIMITED	UK
Azathioprine 50mg film coated tablets	UK/H/0934/001	PA 1128/4/1	RELON CHEM LIMITED	IE
Azathioprin STADA® 50 mg Filmtabletten	DE/H/3036/002	83193.00.00	STADAPHARM GMBH	DE
Azathioprin STADA® 25 mg Filmtabletten	DE/H/3036/001	83192.00.00	STADAPHARM GMBH	DE
Azathioprine CF 50 mg, filmomhulde tabletten	DE/H/3036/002	RVG 108784	CENTRAFARM B.V.	NL
Azathioprine CF 25 mg, filmomhulde tabletten	DE/H/3036/001	RVG 108782	CENTRAFARM B.V.	NL
Imuran Tablets 25 mg	not available	PL 39699/0004	ASPEN PHARMA TRADING LIMITED	UK
Imuran Tablets 50mg	not available	PL 39699/0005	ASPEN PHARMA TRADING LIMITED	UK
Imuran Injection	not available	PL 39699/0003	ASPEN PHARMA TRADING LIMITED	UK
Azathioprin AqVida 50 mg Filmtabletten	DE/H/3037/002	83195.00.00	AQVIDA GMBH	DE
Azathioprin AqVida 25 mg Filmtabletten	DE/H/3037/001	83194.00.00	AQVIDA GMBH	DE
Imasup 25 mg potahované tablety	DE/H/3037/001	59/500/15-C	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	CZ
Imasup 50 mg potahované tablety	DE/H/3037/002	59/501/15-C	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 25 mg filmtabletta	DE/H/3037/001	OGYI-T-22927/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/07	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/08	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/09	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/10	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/11	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atioprin 50 mg filmtabletta	DE/H/3037/002	OGYI-T-22927/12	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Atsimutin 25 mg filmom obložene tablete	DE/H/3037/001	HR-H-555114709	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HR
Atsimutin 50 mg filmom obložene tablete	DE/H/3037/002	HR-H-120877050	PHARMASWISS ČESKÁ REPUBLIKA S.R.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
Atsimutin 50 mg apvalkotās tabletes	DE/H/3037/002	15-0273	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LV
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/007	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/008	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/009	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/010	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/011	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
Atsimutin 50 mg plėvele dengtos tabletės	DE/H/3037/002	LT/1/15/3834/012	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	LT
ATSIMUTIN, 25 mg õhukese polümeerikattega tabletid	DE/H/3037/001	891615	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
ATSIMUTIN, 50 mg õhukese polümeerikattega tabletid	DE/H/3037/001	891715	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	EE
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO

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
Atsimutin 50 mg comprimate filmate	DE/H/3037/002	8474/2015/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Atsimutin 25 mg comprimate filmate	DE/H/3037/001	8473/2015/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	RO
Imasup 25 mg filmom obalené tablety	DE/H/3037/001	59/0078/16-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK
Imasup 50 mg filmom obalené tablety	DE/H/3037/002	59/0079/16-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK
Atioprin, 25 mg, tabletki powlekane	DE/H/3037/001	23092	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Atioprin, 50 mg, tabletki powlekane	DE/H/3037/002	23093	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Azathioprin dura N 25 mg Filmtabletten Wirkstoff: Azathioprin	DE/H/5614/001	47315.00.00	MYLAN DURA GMBH	DE
Azathioprin dura N 50 mg Filmtabletten Wirkstoff: Azathioprin	DE/H/5614/002	47315.01.00	MYLAN DURA GMBH	DE
AZATHIOPRINE TEVA 50 mg, comprimé pelliculé	not available	NL29846	TEVA SANTÉ	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
IMUREL 25 mg kalvopäällysteinen tabletti	not available	9809	ASPEN PHARMA TRADING LIMITED	FI
Imurel® 50 mg filmdragerade tabletter	not available	5204	ASPEN PHARMA TRADING LIMITED	FI
Imurel® 25 mg filmdragerade tabletter	not available	9809	ASPEN PHARMA TRADING LIMITED	FI
IMUREL 50 mg kalvopäällysteinen tabletti	not available	5204	ASPEN PHARMA TRADING LIMITED	FI