09 July 2026
EMADOC-1700519818-3326852
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): apomorphine

EURD list No. PSUSA/00000227/202511

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
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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
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	86569/15/08-10-2018	ITF HELLAS S.A.	GR
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	21473	ITF HELLAS S.A.	CY
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	86569/15/08-10-2018	ITF HELLAS S.A.	GR
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	86569/15/08-10-2018	ITF HELLAS S.A.	GR
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	86569/15/08-10-2018	ITF HELLAS S.A.	GR
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	21473	ITF HELLAS S.A.	CY
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	21473	ITF HELLAS S.A.	CY
APO-go Συσκευή τύπου πένας 10 mg/ml Ενέσιμο Διάλυμα	IE/H/0658/001	21473	ITF HELLAS S.A.	CY
APO-go 5 mg/ml solução para perfusão em cartucho	IE/H/0658/004	5809330	STADA ARZNEIMITTEL AG	PT
APO-go 5 mg/ml solução para perfusão em cartucho	IE/H/0658/004	5805031	STADA ARZNEIMITTEL AG	PT
APO-go 5 mg/ml solução para perfusão em cartucho	IE/H/0658/004	5809322	STADA ARZNEIMITTEL AG	PT
APO-go® 5 mg/ml Infusionslösung in einer	IE/H/0658/004	2203345.00.00	STADAPHARM GMBH	DE

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Patrone				
APO-go® 5 mg/ml Infusionslösung in einer Patrone	IE/H/0658/004	2203345.00.00	STADAPHARM GMBH	DE
APO-go® 5 mg/ml Infusionslösung in einer Patrone	IE/H/0658/004	2203345.00.00	STADAPHARM GMBH	DE
Apo-go 5 mg/ml Infusionslösung in einer Patrone	IE/H/0658/004	2203345.00.00	STADAPHARM GMBH	DE
APO-go för POD 5 mg/ml infusionsvätska, lösning i cylinderampull	IE/H/0658/004	58616	STADA ARZNEIMITTEL AG	SE
APO-go® 5 mg/ml Infusionslösung in einer Patrone	IE/H/0658/004	140268	STADA ARZNEIMITTEL AG	AT
APO-go Pod 5 mg/ml oplossing voor infusie in een patroon	IE/H/0658/004	RVG 124244	STADA ARZNEIMITTEL AG	NL
Britaject 5 mg/ml injeksjonsvæske, oppløsning i sylinderampulle	IE/H/0658/004	18-12638	STADA ARZNEIMITTEL AG	NO
APO-go POD, infusionsvæske, opløsning i cylinderampul	IE/H/0658/004	62088	STADA ARZNEIMITTEL AG	DK
APO-go POD 5 mg/ml solution for infusion in cartridge	IE/H/0658/004	PA0593/042/004	STADA ARZNEIMITTEL AG	IE
APO-go 5 mg/ml raztopina za infundiranje v vložku	IE/H/0658/004	H/12/00205/020	STADA ARZNEIMITTEL AG	SI
APO-go 5 mg/ml raztopina za infundiranje	IE/H/0658/004	H/12/00205/021	STADA ARZNEIMITTEL AG	SI

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v vložku				
APO-go 5 mg/ml raztopina za infundiranje v vložku	IE/H/0658/004	H/12/00205/019	STADA ARZNEIMITTEL AG	SI
APO-GO POD 5 MG/ML SOLUCION PARA PERFUSION EN CARTUCHO	IE/H/0658/004	85.464	STADA ARZNEIMITTEL AG	ES
APOKINON 30 mg/3 ml (1 %), solution injectable en stylo pré-rempli	not available	3400933870867	LABORATOIRE AGUETTANT	FR
APOKINON 30 mg/3 ml (1 %), solution injectable en stylo pré-rempli	not available	3400933870577	LABORATOIRE AGUETTANT	FR
APOKINON 30 mg/3 ml (1 %), solution injectable en stylo pré-rempli	not available	3400933870638	LABORATOIRE AGUETTANT	FR
APO-go 5 mg/ml Infusionslösung in einer Fertigspritze	IE/H/0658/003	69752.00.00	STADAPHARM GMBH	DE
APO-go Pumpfill 5 mg/ml infusionsvätska, lösning i förfylld spruta	IE/H/0658/003	25942	STADA ARZNEIMITTEL AG	SE
APO-go 5mg/ml soluţie perfuzabilă în seringă preumplută unidoză	IE/H/0658/003	12115/2019/01	STADA ARZNEIMITTEL AG	RO
Britaject 5 mg/ml infusjonsvæske, oppløsning i ferdigfylt sprøyte	IE/H/0658/003	10-8045	STADA ARZNEIMITTEL AG	NO
Apo-go® 5 mg/ml Solução para perfusão	IE/H/0658/003	5411616	STADA ARZNEIMITTEL AG	PT

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em seringa pré-cheia				
APO-go 5 mg/ml raztopina za infundiranje v napolnjeni injekcijski brizgi	IE/H/0658/003	H/12/00205/016	STADA ARZNEIMITTEL AG	SI
APO-go PFS 5mg/ml Solution for Infusion in Pre-filled Syringe	IE/H/0658/003	PA0593/043/003	STADA ARZNEIMITTEL AG	IE
APO-go 5mg/ml soluţie perfuzabilă în seringă preumplută unidoză	IE/H/0658/003	12115/2019/02	STADA ARZNEIMITTEL AG	RO
APO-go 5mg/ml soluţie perfuzabilă în seringă preumplută unidoză	IE/H/0658/003	12115/2019/03	STADA ARZNEIMITTEL AG	RO
АПО-го ПНС 5 mg/ml инфузионен разтвор в предварително напълнена спринцовка	IE/H/0658/003	20110740	STADA ARZNEIMITTEL AG	BG
APO-go 5 mg/ml raztopina za infundiranje v napolnjeni injekcijski brizgi	IE/H/0658/003	H/12/00205/018	STADA ARZNEIMITTEL AG	SI
APO-go 5 mg/ml raztopina za infundiranje v napolnjeni injekcijski brizgi	IE/H/0658/003	H/12/00205/017	STADA ARZNEIMITTEL AG	SI
APO-go Pumpfill, infusionsvæske, opløsning i fyldt injektionssprøjte	IE/H/0658/003	41287	STADA ARZNEIMITTEL AG	DK
APO-go 5 mg/ml Infusionslösung in einer Fertigspritze	IE/H/0658/003	1-27540	STADA ARZNEIMITTEL AG	AT
APO-go® 5 mg/ml oplossing voor infusie in	IE/H/0658/003	RVG 35295	STADA ARZNEIMITTEL AG	NL

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een voorgevulde spuit				
APO-go® PFS 5 mg/ml Solution for Infusion in Pre-filled Syringe	IE/H/0658/003	MA 830/00301	STADA ARZNEIMITTEL AG	MT
APO-go PFS 5 mg/ml Solución para Perfusión en Jeringa Precargada	IE/H/0658/003	74.401	STADA ARZNEIMITTEL AG	ES
APOFIN 50 mg/5 ml soluzione iniettabile per infusione sottocutanea	not available	033403015	CHIESI ITALIA S.P.A.	IT
APOFIN 30 mg/3 ml soluzione iniettabile per uso sottocutaneo	not available	033403041	CHIESI ITALIA S.P.A.	IT
APOFIN 30 mg/3 ml soluzione iniettabile per uso sottocutaneo	not available	033403039	CHIESI ITALIA S.P.A.	IT
KYNMOBI 10 mg, film sublingual	PT/H/2739/001	34009 302 644 1 6	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 10 mg, film sublingual	PT/H/2739/001	NL53905	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 15 mg, film sublingual	PT/H/2739/002	34009 302 644 2 3	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 15 mg, film sublingual	PT/H/2739/002	NL53906	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 20 mg, film sublingual	PT/H/2739/003	NL53907	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 20 mg, film sublingual	PT/H/2739/003	34009 302 644 3 0	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 25 mg, film sublingual	PT/H/2739/004	34009 302 644 4 7	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 25 mg, film sublingual	PT/H/2739/004	NL53908	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 30 mg, film sublingual	PT/H/2739/005	NL53909	BIAL - PORTELA & Cª , SA	FR
KYNMOBI 30 mg, film	PT/H/2739/005	34009 302 644 5 4	BIAL - PORTELA & Cª , SA	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
sublingual				
Pack d'initiation de traitement KYNMOBI 10 mg, film sublingual KYNMOBI 15 mg, film sublingual KYNMOBI 20 mg, film sublingual KYNMOBI 25 mg, film sublingual KYNMOBI 30 mg, film sublingual	PT/H/2739/006	34009 302 644 6 1	BIAL - PORTELA & Cª , SA	FR
Kynmobi 10 mg película sublingual	PT/H/2739/001	5847546	BIAL - PORTELA & Cª , SA	PT
Kynmobi 10 mg película sublingual	PT/H/2739/001	PT/H/2739/001	BIAL - PORTELA & Cª , SA	PT
Kynmobi 15 mg película sublingual	PT/H/2739/002	5847553	BIAL - PORTELA & Cª , SA	PT
Kynmobi 15 mg película sublingual	PT/H/2739/002	PT/H/2739/002	BIAL - PORTELA & Cª , SA	PT
Kynmobi 20 mg película sublingual	PT/H/2739/003	5847561	BIAL - PORTELA & Cª , SA	PT
Kynmobi 20 mg película sublingual	PT/H/2739/003	PT/H/2739/003	BIAL - PORTELA & Cª , SA	PT
Kynmobi 25 mg película sublingual	PT/H/2739/004	5847579	BIAL - PORTELA & Cª , SA	PT
Kynmobi 25 mg película sublingual	PT/H/2739/004	PT/H/2739/004	BIAL - PORTELA & Cª , SA	PT
Kynmobi 30 mg película sublingual	PT/H/2739/005	5847603	BIAL - PORTELA & Cª , SA	PT
Kynmobi 30 mg película sublingual	PT/H/2739/005	PT/H/2739/005	BIAL - PORTELA & Cª , SA	PT
Kynmobi 10 mg, 15 mg, 20 mg, 25 mg e 30 mg	PT/H/2739/006	5847611	BIAL - PORTELA & Cª , SA	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
película sublingual Embalagem de início de tratamento Kynmobi 10 mg película sublingual Kynmobi 15 mg película sublingual Kynmobi 20 mg película sublingual Kynmobi 25 mg película sublingual Kynmobi 30 mg película sublingual				
Kynmobi 10 mg film sublinguale	PT/H/2739/001	050407028	BIAL - PORTELA & Cª , SA	IT
Kymnobi 10 mg film sublinguale	PT/H/2739/001	050407016	BIAL - PORTELA & Cª , SA	IT
Kynmobi 15 mg film sublinguale	PT/H/2739/002	050407042	BIAL - PORTELA & Cª , SA	IT
Kynmobi 15 mg film sublinguale	PT/H/2739/002	050407030	BIAL - PORTELA & Cª , SA	IT
Kynmobi 20 mg film sublinguale	PT/H/2739/003	050407067	BIAL - PORTELA & Cª , SA	IT
Kynmobi 20 mg film sublinguale	PT/H/2739/003	050407055	BIAL - PORTELA & Cª , SA	IT
Kynmobi 25 mg film sublinguale	PT/H/2739/004	050407081	BIAL - PORTELA & Cª , SA	IT
Kynmobi 25 mg film sublinguale	PT/H/2739/004	050407079	BIAL - PORTELA & Cª , SA	IT
Kynmobi 30 mg film sublinguale	PT/H/2739/005	050407105	BIAL - PORTELA & Cª , SA	IT
Kynmobi 30 mg film sublinguale	PT/H/2739/005	050407093	BIAL - PORTELA & Cª , SA	IT
Confezione di inizio trattamento	PT/H/2739/006	050407117	BIAL - PORTELA & Cª , SA	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
Kynmobi 10 mg film sublinguale Kynmobi 15 mg film sublinguale Kynmobi 20 mg film sublinguale Kynmobi 25 mg film sublinguale Kynmobi 30 mg film sublinguale				
Kynmobi 10 mg Sublingualfilm	PT/H/2739/001	7007533.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 15 mg Sublingualfilm	PT/H/2739/002	7007534.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 20 mg Sublingualfilm	PT/H/2739/003	7007535.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 30 mg Sublingualfilm	PT/H/2739/005	7007537.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 10 mg, 15 mg, 20 mg, 25 mg, 30 mg Sublingualfilm Packung zur Einleitung der Behandlung	PT/H/2739/006	7012700.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 25 mg Sublingualfilm	PT/H/2739/004	7007536.00.00	BIAL - PORTELA & Cª , SA	DE
Kynmobi 10 mg película sublingual	PT/H/2739/001	88976-PT/H/2739/001/DC	BIAL - PORTELA & Cª , SA	ES
Kynmobi 15 mg película sublingual	PT/H/2739/002	88977-PT/H/2739/002/DC	BIAL - PORTELA & Cª , SA	ES
Kynmobi 20 mg película sublingual	PT/H/2739/003	88978-PT/H/2739/003/DC	BIAL - PORTELA & Cª , SA	ES
Kynmobi 25 mg película sublingual	PT/H/2739/004	88979-PT/H/2739/004/DC	BIAL - PORTELA & Cª , SA	ES
Kynmobi 30 mg película sublingual	PT/H/2739/005	88980-PT/H/2739/005/DC	BIAL - PORTELA & Cª , SA	ES

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Envase para el inicio del tratamiento Kynmobi 10 mg + 15 mg + 20 mg + 25 mg + 30 mg película sublingual	PT/H/2739/006	88975-PT/H/2739/006/DC	BIAL - PORTELA & Cª , SA	ES
APO-go PFS 5 mg/ml Διάλυμα για Ἐγχυση σε Προγεμισμένη Σύριγγα	IE/H/0658/003	21474	ITF HELLAS S.A.	CY
APO-go PFS 5 mg/ml Διάλυμα για Ἐγχυση σε Προγεμισμένη Σύριγγα	IE/H/0658/003	17119/17/08-10-2018	ITF HELLAS S.A.	GR
APOMORPHINE VIATRIS 5 mg/ml, solution injectable	not available	34009 301 693 6 0	VIATRIS SANTE	FR
APOMORPHINE VIATRIS 5 mg/ml, solution injectable	not available	34009 301 693 7 7	VIATRIS SANTE	FR
APOMORPHINE VIATRIS 5 mg/ml, solution injectable	not available	34009 301 693 4 6	VIATRIS SANTE	FR
APOKINON 5 mg/ml, solution pour perfusion	not available	3400949740642	LABORATOIRE AGUETTANT	FR
APOKINON 5 mg/ml, solution pour perfusion	not available	3400949740703	LABORATOIRE AGUETTANT	FR
APOKINON 5 mg/ml, solution pour perfusion	not available	3400949740871	LABORATOIRE AGUETTANT	FR
APOKINON 5 mg/ml, solution pour perfusion	not available	3400930202524	LABORATOIRE AGUETTANT	FR
APO-go 10 mg/ml soluţie injectabilă în pen multidoză	IE/H/0658/001	4185/2012/01	STADA ARZNEIMITTEL AG	RO
APO-go POD 5 mg/ml διάλυμα για ἐγχυση σε φυσἰγγιο	IE/H/0658/004	20654/05-03-2021	ITF HELLAS S.A.	GR
APO-go POD 5 mg/ml	IE/H/0658/004	20654/05-03-2021	ITF HELLAS S.A.	GR

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διάλυμα για έγχυση σε φυσίγγιο				
APO-go POD 5 mg/ml διάλυμα για έγχυση σε φυσίγγιο	IE/H/0658/004	20654/05-03-2021	ITF HELLAS S.A.	GR
APO-GO-PEN 10 mg/ml Solution injectable	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml Injektionslösung	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml oplossing voor injectie	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml oplossing voor injectie	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml oplossing voor injectie	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml Injektionslösung	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml Injektionslösung	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml Solution injectable	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-GO-PEN 10 mg/ml Solution injectable	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-go® PEN oplossing voor injectie 10 mg/ml	IE/H/0658/001	RVG 108873	STADA ARZNEIMITTEL AG	NL
APO-GO-PEN 10 mg/ml oplossing voor injectie	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
Britaject 10 mg/ml injeksjonsvæske, oppløsning i ferdigfylt penn.	IE/H/0658/001	10-8044	STADA ARZNEIMITTEL AG	NO
Apo-go® Pen 10 mg/ml solução injetável	IE/H/0658/001	3579786	STADA ARZNEIMITTEL AG	PT
APO-go 10 mg/ml raztopina za injiciranje v	IE/H/0658/001	H/12/00205/001	STADA ARZNEIMITTEL AG	SI

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peresniku				
APO-GO-PEN 10 mg/ml Solution injectable	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
APO-go PEN 10 mg/ml Solution for Injection	IE/H/0658/001	PA 0593/043/001	STADA ARZNEIMITTEL AG	IE
Britaject 10 mg/ml injekcinis tirpalas	IE/H/0658/001	LT/1/12/2917/008	STADA ARZNEIMITTEL AG	LT
Britaject 10 mg/ml injekcinis tirpalas	IE/H/0658/001	LT/1/12/2917/009	STADA ARZNEIMITTEL AG	LT
Britaject 10 mg/ml injekcinis tirpalas	IE/H/0658/001	LT/1/12/2917/007	STADA ARZNEIMITTEL AG	LT
APO-GO-PEN 10 mg/ml Injektionslösung	IE/H/0658/001	BE430692	EUROGENERICS N.V./S.A.	BE
Apo-go Pen 10 mg/ml solução injetável	IE/H/0658/001	3579588	STADA ARZNEIMITTEL AG	PT
Apo-go® Pen 10 mg/ml solução injetável	IE/H/0658/001	3579687	STADA ARZNEIMITTEL AG	PT
Apogo PEN 10 mg/ml injektioneste, liuos	IE/H/0658/001	29340	STADA ARZNEIMITTEL AG	FI
APO-go, 10 mg/ml süstelahus pen-süstlis*	IE/H/0658/001	763011	STADA ARZNEIMITTEL AG	EE
APO-go Pen, injektionsvæske, opløsning	IE/H/0658/001	31715	STADA ARZNEIMITTEL AG	DK
АПО-го ПИСАЛКА 10 mg/ml инжекционен разтвор	IE/H/0658/001	20110739	STADA ARZNEIMITTEL AG	BG
APO-go PEN 10 mg/ml Injektionslösung	IE/H/0658/001	50275.00.00	STADAPHARM GMBH	DE
Britaject PEN 10 mg/ml injekční roztok	IE/H/0658/001	27/787/11-C	STADA ARZNEIMITTEL AG	CZ
APO-go® PEN 10 mg/ml Injektionslösung	IE/H/0658/001	1-24115	STADA ARZNEIMITTEL AG	AT
APO-go PEN 10 mg/ml Solution for Injection	IE/H/0658/001	2015120165	EUROGENERICS N.V./S.A.	LU

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APO-go 10 mg/ml raztopina za injiciranje v peresniku	IE/H/0658/001	H/12/00205/002	STADA ARZNEIMITTEL AG	SI
APO-go 10 mg/ml raztopina za injiciranje v peresniku	IE/H/0658/001	H/12/00205/003	STADA ARZNEIMITTEL AG	SI
APO-go PEN 10 mg/ml injektionsvätska, lösning	IE/H/0658/001	16547	STADA ARZNEIMITTEL AG	SE
APO-go PEN 10 mg/ml Solution for Injection*	IE/H/0658/001	MA 957/00102	STADA ARZNEIMITTEL AG	MT
APO-go PEN 10 mg/ml šķīdums injekcijām* * Tekstā nosaukums saīsināts: APO-go PEN	IE/H/0658/001	12-0060	STADA ARZNEIMITTEL AG	LV
APOMORPHINE HCl STEROP 5 mg/1ml solution injectable	not available	BE223395	LABORATOIRES STEROP	BE
APOMORPHINE HCl STEROP 5 mg/1ml oplossing voor injectie	not available	BE223395	LABORATOIRES STEROP	BE
APOMORPHINE HCl STEROP 5mg/1ml Infusionslösung	not available	BE223395	LABORATOIRES STEROP	BE
APO-go 5 mg/ml soluție perfuzabilă în cartuș	IE/H/0658/004	15997/2025/01	STADA ARZNEIMITTEL AG	RO
APO-go 5 mg/ml soluție perfuzabilă în cartuș	IE/H/0658/004	15997/2025/02	STADA ARZNEIMITTEL AG	RO
APO-go 5 mg/ml soluție perfuzabilă în cartuș	IE/H/0658/004	15997/2025/03	STADA ARZNEIMITTEL AG	RO
APOMORPHINE BIOGARAN 5 mg/ml, solution injectable	not available	3400930169391	BIOGARAN	FR
APOMORPHINE BIOGARAN 5 mg/ml, solution injectable	not available	3400930169384	BIOGARAN	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
APOMORPHINE BIOGARAN 5 mg/ml, solution injectable	not available	3400930169407	BIOGARAN	FR
Dacepton 10 mg/ml Injektionslösung in einer Patrone	AT/H/0524/001	94086.00.00	EVER NEURO PHARMA GMBH	DE
Dacepton® 10 mg/ml Injektionslösung in einer Patrone	AT/H/0524/001	136927	EVER NEURO PHARMA GMBH	AT
Dacepton 10 mg/ml oplossing voor injectie in een patroon	AT/H/0524/001	BE498417	EVER NEURO PHARMA GMBH	BE
Dacepton®	AT/H/0524/001	55009	EVER NEURO PHARMA GMBH	DK
Dacepton® 10 mg/ml injektionsvätska, lösning i cylinderampull	AT/H/0524/001	52041	EVER NEURO PHARMA GMBH	SE
Dacepton® 10 mg/ml oplossing voor injectie in een patroon	AT/H/0524/001	RVG 116466	EVER NEURO PHARMA GMBH	NL
Dacepton® 10 mg/ml injektioneste, liuos, sylinteriampulli	AT/H/0524/001	32822	EVER NEURO PHARMA GMBH	FI
Dacepton 10 mg/ml injekčný roztok v náhradnej náplni	AT/H/0524/001	27/0320/16-S	EVER NEURO PHARMA GMBH	SK
Dopaceptin 10 mg/ml solution injectable/pour perfusion	AT/H/0524/001	34009 300 719 2 2	EVER NEURO PHARMA GMBH	FR
Dopaceptin 10 mg/ml solution injectable/pour perfusion	AT/H/0524/001	34009 300 719 3 9	EVER NEURO PHARMA GMBH	FR
Dopaceptin 10 mg/ml solution injectable/pour perfusion	AT/H/0524/001	34009 300 719 4 6	EVER NEURO PHARMA GMBH	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
Dopaceptin 10 mg/ml solution injectable/pour perfusion	AT/H/0524/001	34009 300 719 5 3	EVER NEURO PHARMA GMBH	FR
Dopaceptin 10 mg/ml solution injectable/pour perfusion	AT/H/0524/001	34009 300 719 6 0	EVER NEURO PHARMA GMBH	FR
Dacepton 10 mg/ml solución inyectable en cartucho EFG	AT/H/0524/001	80998	EVER NEURO PHARMA GMBH	ES
Dacepton® 10 mg/ml otopina za injekciju u ulošku	AT/H/0524/001	HR-H-428746321	EVER NEURO PHARMA GMBH	HR
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/11	EVER NEURO PHARMA GMBH	HU
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/12	EVER NEURO PHARMA GMBH	HU
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/13	EVER NEURO PHARMA GMBH	HU
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/14	EVER NEURO PHARMA GMBH	HU
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/15	EVER NEURO PHARMA GMBH	HU
Dacepton 10 mg/ml oldatos injekció patronban	AT/H/0524/001	OGYI-T-22316/16	EVER NEURO PHARMA GMBH	HU
Dopaceptin® 10 mg/ml ενέσιμο διάλυμα σε φυσίγγιο	AT/H/0524/001	71553/10-08-2017	EVER NEURO PHARMA GMBH	GR
Dopaceptin 10 mg/ml ενέσιμο διάλυμα σε	AT/H/0524/001	71553	EVER NEURO PHARMA GMBH	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
φυσίγγιο				
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/001	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/004	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/003	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/005	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/002	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml raztopina za injiciranje v vložku	AT/H/0524/001	H/17/02416/006	EVER NEURO PHARMA GMBH	SI
Dacepton® 10 mg/ml solution for injection in cartridge	AT/H/0524/001	PA 1774/001/003	EVER NEURO PHARMA GMBH	IE
Дацептон® 10 mg/ml инжекционен разтвор в патрон	AT/H/0524/001	20170209	EVER NEURO PHARMA GMBH	BG
Dacepton 10 mg/ml solution injectable en cartouche	AT/H/0524/001	2017070196	EVER NEURO PHARMA GMBH	LU
Dacepton 10 mg/ml injekční roztok v zásobní vložce	AT/H/0524/001	27/950/16-C	EVER NEURO PHARMA GMBH	CZ
Dacepton 10 mg/ml injeksjonsvæske, oppløsning i sylinderampulle	AT/H/0524/001	16-11441	EVER NEURO PHARMA GMBH	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
Dacepton 10 mg/ml solução injetável em cartucho	AT/H/0524/001	5713961	EVER NEURO PHARMA GMBH	PT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	045369016	EVER NEURO PHARMA GMBH	IT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	45369028	EVER NEURO PHARMA GMBH	IT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	45369030	EVER NEURO PHARMA GMBH	IT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	45369042	EVER NEURO PHARMA GMBH	IT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	45369055	EVER NEURO PHARMA GMBH	IT
Damine 10mg/ml soluzione iniettabile in cartuccia	AT/H/0524/001	45369067	EVER NEURO PHARMA GMBH	IT
APO-go 10 mg/ml injektions-/infusionsvätska, lösning	IE/H/0658/002	18333	STADA ARZNEIMITTEL AG	SE
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/01	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/03	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/02	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/09	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/07	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie	IE/H/0658/002	4186/2012/08	STADA ARZNEIMITTEL AG	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
injectabilă/perfuzabilă				
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/05	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/12	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/11	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/06	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/04	STADA ARZNEIMITTEL AG	RO
APO-go 10 mg/ml soluţie injectabilă/perfuzabilă	IE/H/0658/002	4186/2012/10	STADA ARZNEIMITTEL AG	RO
APO-go Ampullen 10 mg/ml Injektions-/Infusionslösung	IE/H/0658/002	1-24700	STADA ARZNEIMITTEL AG	AT
APO-go Ampullen 10 mg/ml Injektions-/Infusionslösung	IE/H/0658/002	54504.00.00	STADAPHARM GMBH	DE
Britaject 10mg/ml injekcinis/infuzinis tirpalas	IE/H/0658/002	LT/1/12/2917/006	STADA ARZNEIMITTEL AG	LT
Apo-go® 10 mg/ml Solução injetável ou para perfusão	IE/H/0658/002	5411624	STADA ARZNEIMITTEL AG	PT
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/005	STADA ARZNEIMITTEL AG	SI
APO-go AMPOULES 10 mg/ml Solution for Injection or Infusion	IE/H/0658/002	PA0593/043/002	STADA ARZNEIMITTEL AG	IE
Britaject 10 mg/ml injekcinis/infuzinis tirpalas	IE/H/0658/002	LT/1/12/2917/005	STADA ARZNEIMITTEL AG	LT
Britaject 10 mg/ml	IE/H/0658/002	LT/1/12/2917/004	STADA ARZNEIMITTEL AG	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
injekcinis/infuzinis tirpalas				
APO-GO-AMP 10mg/ml Lösung zur Injektion oder Infusion	IE/H/0658/002	BE430701	EUROGENERICS N.V./S.A.	BE
Apo-go® 10 mg/ml Solução injetável ou para perfusão	IE/H/0658/002	5411640	STADA ARZNEIMITTEL AG	PT
Apo-go® 10 mg/ml Solução injetável ou para perfusão	IE/H/0658/002	5411665	STADA ARZNEIMITTEL AG	PT
Apo-go® 10 mg/ml Solução injetável ou para perfusão * Abreviado para Apo-go no texto	IE/H/0658/002	5411632	STADA ARZNEIMITTEL AG	PT
Apo-go® 10 mg/ml Solução injetável ou para perfusão	IE/H/0658/002	5411657	STADA ARZNEIMITTEL AG	PT
Apo-go® 10 mg/ml Solução injetável ou para perfusão	IE/H/0658/002	5411673	STADA ARZNEIMITTEL AG	PT
APO-go® Ampullen 10 mg/ml oplossing voor injectie of infusie	IE/H/0658/002	RVG 109039	STADA ARZNEIMITTEL AG	NL
APO-go, 10 mg/ml süste-/infusioonilahus* * Tekstis lühendatult APO-go	IE/H/0658/002	763111	STADA ARZNEIMITTEL AG	EE
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/004	STADA ARZNEIMITTEL AG	SI
APO-go AMPOULES 10 mg/ml Solution for Injection or Infusion	IE/H/0658/002	2015120166	EUROGENERICS N.V./S.A.	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
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/012	STADA ARZNEIMITTEL AG	SI
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/008	STADA ARZNEIMITTEL AG	SI
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/014	STADA ARZNEIMITTEL AG	SI
APO-go 10 mg/ml raztopina za injiciranje ali infundiranje v ampuli	IE/H/0658/002	H/12/00205/010	STADA ARZNEIMITTEL AG	SI
APO-go 10 mg/ml Solución Inyectable o para Perfusión en Ampollas	IE/H/0658/002	74.400	STADA ARZNEIMITTEL AG	ES
Apodev 5 mg/ml, solución para perfusión EFG	SE/H/1272/001	88396	EVOLAN PHARMA AB	ES
Apomorfine hydrochloride PharmSwed 5 mg/ml, oplossing voor infusie	SE/H/1272/001	RVG 112181	EVOLAN PHARMA AB	NL
Apomorfin ”PharmSwed”	SE/H/1272/001	50982	EVOLAN PHARMA AB	DK
Apomorphine hydrochloride 5 mg/ml, solution for infusion	SE/H/1272/001	PL 44616/0004	EVOLAN PHARMA AB	XI
Apomorphinhydrochlorid PharmSwed 5 mg/ml Infusionslösung	SE/H/1272/001	88102.00.00	EVOLAN PHARMA AB	DE
Apomorfin PharmSwed 5 mg/ml, infusionsvätska, lösning	SE/H/1272/001	42703	EVOLAN PHARMA AB	SE
Apomorfin PharmSwed 5 mg/ml infusjonsvæske,	SE/H/1272/001	12-9111	EVOLAN PHARMA AB	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
oppløsning				
Dacepton 5 mg/ml Infusionslösung	AT/H/0364/002	135446	EVER NEURO PHARMA GMBH	AT
Dacepton® 5 mg /ml oplossing voor infusie	AT/H/0364/002	BE456702	EVER NEURO PHARMA GMBH	BE
Dacepton® 5 mg/ml Solution pour perfusion	AT/H/0364/002	BE456702	EVER NEURO PHARMA GMBH	BE
Dacepton 5 mg/ml infuzní roztok	AT/H/0364/002	27/130/14-C	EVER NEURO PHARMA GMBH	CZ
Dacepton® 5 mg/ml infusionsvæske, opløsning	AT/H/0364/002	28148	EVER NEURO PHARMA GMBH	DK
Dacepton 5 mg/ml infusioonilahus	AT/H/0364/002	839014	EVER NEURO PHARMA GMBH	EE
Dopaceptin® 5 mg/ml Solution pour perfusion	AT/H/0364/002	34009 278 186 9 1	EVER NEURO PHARMA GMBH	FR
Dopaceptin® 5 mg/ml Solution pour perfusion	AT/H/0364/002	34009 278 187 5 2	EVER NEURO PHARMA GMBH	FR
Dacepton 5 mg/ml oldatos infúzió	AT/H/0364/002	OGYI-T-22316/04	EVER NEURO PHARMA GMBH	HU
Dacepton 5 mg/ml oldatos infúzió	AT/H/0364/002	OGYI-T-22316/05	EVER NEURO PHARMA GMBH	HU
Dacepton 5 mg/ml oldatos infúzió	AT/H/0364/002	OGYI-T-22316/06	EVER NEURO PHARMA GMBH	HU
Dacepton 5 mg/ml oldatos infúzió	AT/H/0364/002	OGYI-T-22316/07	EVER NEURO PHARMA GMBH	HU
Dacepton 5 mg/ml oldatos infúzió	AT/H/0364/002	OGYI-T-22316/08	EVER NEURO PHARMA GMBH	HU
Dacepton 5 mg/ml Infusionslösung	AT/H/0364/002	88898.00.00	EVER NEURO PHARMA GMBH	DE
Dacepton 5 mg/ml infúzny roztok	AT/H/0364/002	27/0099/14-S	EVER NEURO PHARMA GMBH	SK
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/008	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml	AT/H/0364/002	LT/1/11/2759/009	EVER NEURO PHARMA	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
infuzinis tirpalas			GMBH	
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/010	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/011	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/012	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/013	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml infuzinis tirpalas	AT/H/0364/002	LT/1/11/2759/014	EVER NEURO PHARMA GMBH	LT
Dacepton 5 mg/ml solution for infusion	AT/H/0364/002	PA1774/001/002	EVER NEURO PHARMA GMBH	IE
Dacepton 5 mg/ml šķīdums infūzijām	AT/H/0364/002	14-0066	EVER NEURO PHARMA GMBH	LV
Dacepton 5 mg/ml oplossing voor infusie	AT/H/0364/002	RVG 112999	EVER NEURO PHARMA GMBH	NL
Dacepton 5 mg/ml infusjonsvæske, oppløsning	AT/H/0364/002	12-9355	EVER NEURO PHARMA GMBH	NO
Dacepton 5 mg/ml infusionsvätska, lösning	AT/H/0364/002	48853	EVER NEURO PHARMA GMBH	SE
Dacepton 5 mg/ml raztopina za infundiranje	AT/H/0364/002	5363-I-1007/14	EVER NEURO PHARMA GMBH	SI
Dacepton 5 mg/ml raztopina za infundiranje	AT/H/0364/002	5363-I-1008/14	EVER NEURO PHARMA GMBH	SI
Dacepton 5 mg/ml raztopina za infundiranje	AT/H/0364/002	5363-I-1009/14	EVER NEURO PHARMA GMBH	SI
Dacepton 5 mg/ml raztopina za infundiranje	AT/H/0364/002	5363-I-1010/14	EVER NEURO PHARMA GMBH	SI
Dacepton 5 mg/ml raztopina za infundiranje	AT/H/0364/002	5363-I-1011/14	EVER NEURO PHARMA GMBH	SI
Дацептон 5 mg/ml инфузионен разтвор	AT/H/0364/002	20140074	EVER NEURO PHARMA GMBH	BG
Dacepton 10 mg/ml,	AT/H/0364/001	789412	EVER NEURO PHARMA	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
süste- või infusioonilahus			GMBH	
Dacepton 10 mg /ml oplossing voor injectie of infusie	AT/H/0364/001	RVG 111059	EVER NEURO PHARMA GMBH	NL
Dacepton 10 mg/ml injeksjons-/infusjonsvæske, oppløsning	AT/H/0364/001	11-8741	EVER NEURO PHARMA GMBH	NO
Dacepton 5 mg/ml solución para perfusión EFG	AT/H/0364/002	79174	EVER NEURO PHARMA GMBH	ES
Dacepton® 5 mg /ml infuusioneste, liuos	AT/H/0364/002	31216	EVER NEURO PHARMA GMBH	FI
Dacepton, 5 mg/ml, roztwór do infuzji	AT/H/0364/002	22789	EVER NEURO PHARMA GMBH	PL
Dopaceptin 5 mg/ml Διάλυμα για έγχυση	AT/H/0364/002	49653/13.06.2014	EVER NEURO PHARMA GMBH	GR
Dopaceptin 10 mg/ml Διάλυμα για ένεση ή έγχυση	AT/H/0364/001	31746/10.06.2014	EVER NEURO PHARMA GMBH	GR
Dacepton 5 mg/ml solução para perfusão	AT/H/0364/002	5672167	EVER NEURO PHARMA GMBH	PT
Dacepton 5 mg/ml solução para perfusão	AT/H/0364/002	5672175	EVER NEURO PHARMA GMBH	PT
Dacepton, 10 mg/ml, roztwór do wstrzykiwań we wkładzie	AT/H/0364/001	23778	EVER NEURO PHARMA GMBH	PL
Dacepton 10 mg/ml Injektions-/Infusionslösung	AT/H/0364/001	1-30904	EVER NEURO PHARMA GMBH	AT
Dacepton 10 mg/ml šķīdums injekcijām vai infūzijām	AT/H/0364/001	11-0308	EVER NEURO PHARMA GMBH	LV
Dacepton 10 mg/ml	AT/H/0364/001	5403779	EVER NEURO PHARMA	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
Solução injetável ou para perfusão			GMBH	
Dacepton 10 mg/ml Solução injetável ou para perfusão	AT/H/0364/001	5403761	EVER NEURO PHARMA GMBH	PT
Dacepton 10 mg/ml Solução injetável ou para perfusão	AT/H/0364/001	5403753	EVER NEURO PHARMA GMBH	PT
Dacepton 10 mg/ml Injekčný a infúzny roztok	AT/H/0364/001	27/0360/11-S	EVER NEURO PHARMA GMBH	SK
Дацептон 10 mg/ml Инжекционен/инфузионен разтвор	AT/H/0364/001	20110560	EVER NEURO PHARMA GMBH	BG
Dacepton, injektions- og infusionsvæske, opløsning	AT/H/0364/001	49907	EVER NEURO PHARMA GMBH	DK
Dacepton 5 mg/ml otopina za infuziju	AT/H/0364/002	HR-H-643764746	EVER NEURO PHARMA GMBH	HR
Dacepton 5 mg/ml solution pour perfusion	AT/H/0364/002	2020020039	EVER NEURO PHARMA GMBH	LU
Dacepton 5mg/ml soluzione per infusione	AT/H/0364/002	042035042	EVER NEURO PHARMA GMBH	IT
Dacepton 5mg/ml soluzione per infusione	AT/H/0364/002	042035067	EVER NEURO PHARMA GMBH	IT
Dacepton 5mg/ml soluzione per infusione	AT/H/0364/002	042035055	EVER NEURO PHARMA GMBH	IT
Dacepton 10 mg/ml soluzione iniettabile o per infusione	AT/H/0364/001	042035016	EVER NEURO PHARMA GMBH	IT
Dacepton 10 mg/ml soluzione iniettabile o per infusione	AT/H/0364/001	042035028	EVER NEURO PHARMA GMBH	IT
Dacepton 10 mg/ml soluzione iniettabile o	AT/H/0364/001	042035030	EVER NEURO PHARMA GMBH	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
per infusione				