30 January 2019
EMA/185142/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal product

Active substance: domperidone

Procedure no.: EMEA/H/N/PSR/J/0015

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Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Motilium 1 mg/ml - Suspension zum Einnehmen	BE/H/0106/003	1-20462	Janssen-Cilag Pharma GmbH	Austria
Motilium 10 mg - Filmtabletten	BE/H/0106/009	17412		
Motilium 10 mg film- coated tablet (Domperidone maleate)	BE/H/0106/001	BE109986	Johnson & Johnson Consumer N.V.	Belgium
Motilium 1 mg/ml oral suspension (ped)	BE/H/0106/002	BE110013		
Motilium 1 mg/ml oral suspension (adults)	BE/H/0106/003	BE190662		
Motilium Instant 10 mg orodispersible tablet	BE/H/0106/008	BE274827		
Motilium 10 mg film-coated tablet (Domperidone base)	BE/H/0106/009	BE272167		
Domperidone Ibidel	n/a	BE445611		
MOTILIUM 10 mg Film-Coated Tablets	Not applicable	20010169	Johnson & Johnson, Prodaja medicinskih in farmacevtskih izdelkov, d.o.o.	Bulgaria

Product Name (in authorisation country)	MRP / DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Motilium® 10 mg Film-coated Tablets	Not applicable	7224	Janssen-Cilag International NV	Cyprus
Motilium 1mg/ml Oral Suspension		7459		
MOTILIUM Potahované tablety	Not applicable	20/813/93-C	Janssen-Cilag s.r.o	Czech Republic
Motilium	BE/H/0106/001	09662	Janssen-Cilag A/S	Denmark
MOTILIUM 1 mg/ml, suspension buvable	BE/H/0106/002-003	34009 323 409-8 9 34009 323 408-1 1	Janssen-Cilag	France
MOTILIUM 10 mg, comprimé pelliculé	BE/H/0106/009	34009 323 411-2 2 34009 336 882-9 5		
Cilroton 1 mg/ml – Oral Suspension	BE/H/0106/003	57067/07/5-11-2008	Johnson & Johnson Hellas Consumer AE	Greece
Cilroton 10 mg – Film coater tablets	BE/H/0106/009	70681/5-11-2008		
MOTILIUM 10 mg filmtabletta	Not applicable	OGYI-T-2223/01	Janssen-Cilag Kft.	Hungary
Motilium Fastmelts 10mg Tablets	N/A	823/51/1 823/51/2	McNeil Healthcare (Ireland) Ltd	Ireland
Motilium 10mg Film-coated Tablets	BE/H/0106/003	823/51/6		
Motilium 1mg/ml Oral Suspension	BE/H/0106/001	823/51/7		
Motilium Rx 10mg Tablets				
MOTILIUM 1 mg/ml sospensione orale	BE/H/0106/003	024953022	Janssen-Cilag SpA	Italy
MOTILIUM 10 mg compresse rivestite con film	BE/H/0106/009	024953034		

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Motilium 10 mg apvalkotās tabletes	Not applicable	98-0800	UAB Johnson & Johnson	Latvia
Motilium 10 mg plėvele dengtos tabletės	Not applicable	LT/1/95/0982/001-002	UAB Johnson & Johnson	Lithuania
Motilium 10 mg film-coated tablet (Domperidone maleate)	BE/H/0106/001	2005028771	Johnson & Johnson Consumer N.V.	Luxembourg
Motilium 1 mg/ml oral suspension (ped)	BE/H/0106/002	2005058774		
Motilium 1 mg/ml oral suspension (adults)	BE/H/0106/003	2005058773		
Motilium Instant 10 mg orodispersible tablet	BE/H/0106/008	2005059999		
Motilium 10 mg film-coated tablet (Domperidone base)	BE/H/0106/009	2008120046		
Domperidone Ibidel	n/a	2014060181		
Motilium 10 mg Film-coated Tablets	Not applicable	018/01605	Janssen-Cilag International NV	Malta
Motilium 1mg/ml Oral Suspension		018/01604		

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Motilium 10 mg film-coated tablet (Domperidone maleate)	BE/H/0106/001	RVG 07678	Johnson & Johnson Consumer B.V.	Netherlands
Motilium 1 mg/ml oral suspension (adults)	BE/H/0106/003	RVG 07679		
Domperidon JJC	n/a	RVG 20546=07682		
Domperidon JJC	n/a	RVG 20544=07678		
Motilium Comprimido revestido 10 mg	BE/H/106/009	9512822 (60 units); 9512814 (20 units)	Johnson & Johnson Lda.	Portugal
Motilium Suspensão oral 1 mg/ml	BE/H/106/003	9532200 (1 unit 200 ml bottle); 9532234 (1 unit 100 ml bottle)		
Nauzelín 10 mg comprimidos recubiertos con película	BE/H/0106/009	56.679	Janssen-Cilag, S.A.	Spain
Nauzelín 1 mg/ml suspensión oral	BE/H/0106/003	56.670		
Domperidone 10mg Film-Coated Tablets	Not Applicable (National)	PL 29831/0076	Wockhardt UK Limited	United Kingdom
Domperidone 10mg Film-Coated Tablets	Not Applicable (National)	AA154/05801	Wockhardt UK Limited	Malta
Domperidone 1mg/ml Oral Suspension	UK/H/5247/01/DC	PL 29831/0522	Wockhardt UK Limited	United Kingdom
Domperidone 1mg/ml Oral Suspension	UK/H/5247/01/DC	PA 1339/050/001	Wockhardt UK Limited	Ireland
DOMPERIDONE ARISTO	IT/H/247/01/MR	037402017	Aristo Pharma GmbH	Italy
DOMPERIDONE ANGENERICO	IT/H/247/01/MR	037402029	Aristo Pharma GmbH	Italy

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
DOMPERIDONE ARISTO	IT/H/247/01/MR	037402031	Aristo Pharma GmbH	Italy
DOMPERIDONE ARISTO	IT/H/247/01/MR	037402043	Aristo Pharma GmbH	Italy
DOMPERIDONE ARISTO	IT/H/247/01/MR	037402056	Aristo Pharma GmbH	Italy
Domperidone 10mg Tablets	Not applicable	14251/0002	Manx Healthcare	United Kingdom
Domperidone ABC 10 mg compresse	Not applicable	035809019	ABC Farmaceutici S.p.A.	Italy
DOMPERIDONE ALMUS 10 mg, comprimé pelliculé	Not applicable	30258	Biogaran	France
DOMPERIDONE BIOGARAN 10 mg, comprimé pelliculé	Not applicable	28155	Biogaran	France
DOMPERIDONE BIOGARAN 10 mg, comprimé orodispersible	Not applicable	34351	Biogaran	France
Domperidone 10 mg Tablets	Not applicable	PL 17907/0096	Bristol Laboratories Ltd	United Kingdom
Domperidone 10 mg Tablets	Not applicable	PL 17907/0322	Bristol Laboratories Ltd	United Kingdom
Domperidone 10 mg Tablets	Not applicable	PL 17907/0323	Bristol Laboratories Ltd	United Kingdom

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidone 10mg Tablets	Not applicable	PL 20046/0086	Focus Pharmaceuticals Limited	United Kingdom
Cinet 10 mg Comprimido Cinet 10 mg Comprimido Dispersível Cinet 1mg/ml suspensão oral	Not applicable	9514802 9514828 4432597 4432696 5080874 9621714 9621706	Laboratório Medinfar - Produtos Farmacêuticos, S.A.	Portugal
Cinet 1mg/ml Oral Suspension Cinet 10 mg Tablets	Not applicable	531/00202 531/00201	Laboratório Medinfar - Produtos Farmacêuticos, S.A.	Malta
Domperidona GP 10 mg Comprimido	Not applicable	5343991 5344098	GP - Genéricos Portugueses, Lda.	Portugal
RIGES 10 mg compresse	Not applicable	036107011	S.F. Group S.r.l.	Italy
Motilium 10mg film-coated tablets	Not Applicable	PL 17780/0300	Winthrop Pharmaceuticals UK Ltd trading as Zentiva	United Kingdom
Domperidone 1mg/ml oral suspension	Not Applicable	PL 17780/0299	Winthrop Pharmaceuticals UK Ltd trading as Zentiva	United Kingdom
DOMPERIDONE GERDA 10 mg, comprimé pelliculé	Not applicable	NL29209	Laboratoires Gerda	France
Domperidona toLife 10 mg comprimidos	Not applicable	5430996 5431093	Tolife Produtos Farmacêuticos S.A.	Portugal

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidona Pensa 10 mg comprimidos EFG	Not applicable	67.523	Pensa Pharma, S.A	Spain
Domperidone pensa 10 mg compresse effervescenti	Not applicable	037233057	Pensa Pharma S.p.A.	Italy
Motilium 10 mg comprimate filmate	N/A	2068/2009/01	Terapia SA	Romania
DOMPERIDONE ZYDUS 10 mg, comprimé pelliculé	National	34009 300 571 4 8 34009 363 949 3 3 34009 363 950 1 5 34009 364 184 0 0 34009 565 178 8 6	Zydus France	France
Domperidon beta 10 mg Tabletten	DE/H/3359/001/MR	73433.00.00	betapharm Arzneimittel GmbH	Germany
Dismotil 10mg Tablets/Domperidone 10mg tablets	NA	13606/0090	Co-pharma Ltd	United Kingdom
DOMPERIDONE DOC Generici	National	036109015	DOC Generici S.r.l.	Italy
DOMPERIDONE SANDOZ 10 mg, comprimé orodispersible		34009 382 336 3 6 34009 382 338 6 5	Sandoz S.A.S.	France
Domperidon Hexal 10 mg Tabletten	DE/H/0818/001	55178.00.00	Hexal AG	Germany
Domerid Relief 10mg Tablets	DE/H/0818/001	PA 711/46/2	Rowex LTD	Ireland
DOMERID 10mg Tablets	DE/H/0818/001	PA 711/46/1	Rowex LTD	Ireland
DOMPERIDONE SANDOZ		036809010	Sandoz S.P.A.	Italy

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
DOMPERIDON SANDOZ 10 MG, TABLETTEN	NL/H/0490/001	RVG 30008	Sandoz B.V.	Netherlands
Domperidon - 1 A Pharma® 10 mg Tabletten	NL/H/0490/001	59671.00.00	1 A Pharma GMBH	Germany
DOMPERIDONE SANDOZ 10 mg, comprimé pelliculé		34009 300 630 2 6 34009 550 228 8 6 34009 550 228 9 3 34009 550 229 1 6 34009 564 398 4 3	Sandoz S.A.S.	France
Domperidon Sandoz 10mg orodispergeerbare tabletten	PT/H/1353	BE502071	SANDOZ N.V.	Belgium
Domperidona Sandoz	PT/H/1353	MA number not assigned	SANDOZ FARMACEUTICA LDA.	Portugal
Domperidone Giuliani 5 mg compresse masticabili	Not applicable	032090021	Giuliani Spa	Italy
Domperidone Giuliani 5 mg granulato effervescente	Not applicable	032090019	Giuliani Spa	Italy
Domperidone Giuliani 10 mg compresse rivestite con film	Not applicable	032090058	Giuliani Spa	Italy
Domperidona Generis 10 mg comprimidos revestidos por película	Not applicable	5104682 5104781 5104880	Generis farmacêutica S.A.	Portugal
Peridon 10 mg compresse rivestite con film	Not applicable	024309039	Italchimici S.p.A.	Italy
Peridon 10 mg granulato effervescente	Not applicable	024309130	Italchimici S.p.A.	Italy

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Peridon 1 mg/ml sospensione orale	Not applicable	024309142	Italchimici S.p.A.	Italy
Peridon 30 mg supposte	Not applicable	024309066	Italchimici S.p.A.	Italy
Zilium 10 mg, tabletten	NL/H/241/1	BE221392	Kela Pharma	Belgium
Domperidona Labesfal 10 mg comprimidos	Not applicable	5764287 5764386	Generis Farmacêutica S.A.	Portugal
DOMPERIDONE MYLAN 10 mg, comprimé pelliculé	n/a	NL 27909	Mylan SAS	France
Domperidona Mylan	n/a	5104989,5105085, 5105184	Mylan Lda	Portugal
Domperidone Mylan Generics	n/a	035810011	Mylan SpA	Italy
Domperidone Mylan 10 mg, comprimé orodispersible	n/a	NL 34350	Mylan SAS	France
Domperidon Mylan 10 mg, tabletten	n/a	RVG 22830	Mylan BV	The Netherlands
Domperidon Mylan 10 mg, filmomhulde tabletten	n/a	BE208722	Mylan BVBA/SPRL	Belgium
OROPERIDYS 10 mg, ORODISPERSIBLE TABLET	FR/H/0335/01/MR	BE329357	Pierre Fabre Medicament	Belgium
OROPERIDYS ORODISPERSIBLE TABLETS 10MG	FR/H/0335/01/MR	20499	Pierre Fabre Medicament	Cyprus
OROPERIDYS 10 mg	FR/H/0335/01/MR	20/652/08-C	Pierre Fabre Medicament	Czech Republic

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
OROPERIDYS 10 mg	FR/H/0335/01/MR	607408	Pierre Fabre Medicament	Estonia
PERIDYS 10 mg, comprimé pelliculé	Not applicable	34009 328 641 6 4	Pierre Fabre Medicament	France
OROPERIDYS 10 mg, comprimé orodispersible	Not applicable	34009 374 778 0 2	Pierre Fabre Medicament	France
DOMPERIDONE PIERRE FABRE MEDICAMENT 10 mg, comprimé orodispersible	Not applicable	34009 374 514 3 7 34009 416 553 1 9	Pierre Fabre Medicament	France
DOMPERIDONE PIERRE FABRE 10 mg, comprimé orodispersible	FR/H/0335/01/MR	34009 398 031 2 8 34009 379 252 7 3	Pierre Fabre Medicament	France
PERIDYS 1 mg/ml, suspension buvable	Not applicable	34009 328 642 2 5	Pierre Fabre Medicament	France
OROPERIDYS 10 mg, ORODISPERSIBLE TABLETS	FR/H/0335/01/MR	51705/08/13-2-2009	Pierre Fabre Medicines SA	Greece
NEOPERIDYS 10 mg, ORODISPERSIBLE TABLET	FR/H/0335/01/MR	039985027 039985015	Pierre Fabre Pharma S.R.L.	Italy
OROPERIDYS 10 mg, ORODISPERSIBLE TABLETS	FR/H/0335/01/MR	08-0269	Pierre Fabre Medicament	Latvia

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OROPERIDYS 10 mg, ORODISPERSIBLE TABLETS	FR/H/0335/01/MR	LT/1/08/1317/001 LT/1/08/1317/002	Pierre Fabre Medicament	Lithuania
OROPERIDYS 10 mg, ORODISPERSIBLE TABLET	FR/H/0335/01/MR	2009030033	Pierre Fabre Medicament	Luxembourg
OROPERIDYS 10 mg, ORODISPERSIBLE TABLET	FR/H/0335/01/MR	15213	Pierre Fabre Médicament	Poland
OROPERIDYS 10 mg, ORODISPERSIBLE TABLETS	FR/H/0335/01/MR	4203/2012/01 4203/2012/02	Pierre Fabre Médicament	Romania
OROPERIDYS 10 mg, TABLET	FR/H/0335/01/MR	20/0503/08-S	Pierre Fabre Médicament	Slovak Republic
Domperidone Instant EG 10 mg orodispergeerbare tabletten	FR/H/0364/001	BE342946	Eurogenerics N.V./S.A.	Belgium
Domperidone EG 10 mg tabletten	National	BE249496		
DOMPERIDONE EG 10 mg, comprimé orodispersible	FR/H/0364/001	NL 34352	EG LABO Laboratoires EuroGenerics	France
DOMPERIDONE EG 10 mg, comprimé pelliculé	National	NL 28350		
Domperidon AL 10 mg Tabletten	DE/H/1214/001	55220.00.00	Aliud Pharma GmbH Stada Pharm GmbH	Germany
Domperidon STADA 10 mg Tabletten	NL/H/0196/001	54070.00.00		
Domperidone EG 10 mg compresse	National	035812015	EG S.p.A.	Italy
RAXAR 10 mg compresse orodispersibili	FR/H/0364/001	039200/M	Crinos S.p.A.	

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidone Instant EG 10 mg comprimés orodispersibles	FR/H/0364/001 National	2010050043 2007060035	Eurogenerics N.V./S.A.	Luxembourg
Domperidone EG 10 mg comprimés				
Domperidon STADA 10 mg tabletten	NL/H/0196/001	RVG 25168	STADA Arzneimittel AG	Netherlands
Domperidon CF 10 mg, tabletten	National	RVG 26086=24277	Centrafarm B.V	
Domperidon HTP 10 mg, tabletten	National	RVG 24277	Healthypharm B.V.	
DOMPERIDONE STADA 10 mg	BE/H/0249/001	BE501582	STADA Arzneimittel AG	Belgium
DOMPERIDONE 10 mg, orodispersible tablets	BE/H/0249/001	PA0126/291/001	Clonmel Healthcare Ltd.	Ireland
Domperidona Ciclum	National	PTmutiple	Ciclum Farma Unipessoal Lda.	Portugal
DOMPERIDONE Stada 10 mg, raspadljive tablete za usta	BE/H/0249/001	HR-H-083826960	STADA d.o.o.	Croatia
Domperidona Flas STADA 10 mg comprimidos bucodispersables EFG	BE/H/0249/001	81.379	Laboratorio STADA, S.L.	Spain
Domperidona Azevedos 10mg comprimidos	Not applicable	5586581 5586680 5811088	Laboratórios Azevedos-Indústria Farmacêutica, S.A.	Portugal
Remotil 10 mg comprimidos	Not applicable	9514604 9514612	Laboratórios Azevedos-Indústria Farmacêutica, S.A.	Portugal
Motilium Tabletten Motilium Tropfen 10 mg/ml Suspension	DE/H/0415/001 DE/H/0415/002	43.00.00 43.00.02	Takeda GmbH	Germany

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidone	NL/H/0195/001/MR	RVG 24213	Accord Healthcare Limited, UK	The Netherlands
Domperidon Actavis 10 mg tabletes	NL/H/0830/001	LT/1/06/0615/001-005	Actavis Group hf.	Lithuania
Domperidon Actavis 10 mg tabletes	NL/H/0830/001	06-0269	Actavis Group PTC ehf.	Latvia
Domperidon Actavis	NL/H/0830/001	523706	Actavis Group PTC ehf.	Estonia
DOMPERIDONE ZENTIVA 10 mg, comprimé pelliculé	Not applicable	CIS : 6 241 755 2	Sanofi Aventis France	France
DOMPERIDONE ZENTIVA 10 mg, comprimé orodispersible	Not applicable	CIS : 6 506 501 7	Sanofi Aventis France	France
DOMPERIDONA GAMIR 1mg/ml Suspensión oral	NATIONAL	57151	Meda S.L.U.	Spain
DOMPERIDONA GAMIR 10 mg, Cápsulas duras	NATIONAL	57096	Meda S.L.U.	Spain
Domperidon-TEVA 10 mg Filmtabletten	NL-H-0208-01	50103.00.00	Teva GmbH; Ulm	Germany
Domperidon TEVA 10 mg filmomhulde tabletten	NL-H-0208-01	BE372066	Teva Pharma Belgium NV; Laarstraat 16, Wilrijk 2610, Belgium	Belgium

Product Name (in authorisation country)	MRP / DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061011	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061023	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061035	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061047	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061050	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061062	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061074	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidone Teva 10 mg compresse rivestite con film	NL-H-0208-01	035061086	Teva Italia S.r.l.; Via Messina, 38 - 20154 Milano, Italy	Italy
Domperidon 10 PCH, filmomhulde tabletten 10 mg	NL-H-0208-01	RVG 23949	Pharmachemie B.V.; Swensweg 5, P.O. Box 552, Haarlem	Netherlands
Domperidon AbZ 10 mg Filmtabletten	DE-H-1885-001	53898.00.00	AbZ-Pharma GmbH; Graf-Arco-Str. 3, 89079 Ulm, Germany	Germany

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidon-CT 10 mg Filmtabletten	DE-H-1886-001	53897.00.00	AbZ-Pharma GmbH; Graf-Arco-Str. 3, 89079 Ulm, Germany	Germany
Domperidon-ratiopharm® 10 mg Filmtabletten	DE-H-1887-001	53896.00.00	ratiopharm GmbH; Graf-Arco-Str. 3, 89079 Ulm, Germany	Germany
Domperidone Teva 10 mg, comprimé pelliculé	FR NL27592	NL27592	Teva Santé SAS	France
Domperidone Teva 10 mg, comprimé orodispersible	FR NL38642	NL38642	Teva Santé SAS	France
Domperidon 10 mg PCH, tabletten	RVG 104463	RVG 104463	Pharmachemie B.V.; Swensweg 5, P.O. Box 552, Haarlem	Netherlands
Domperidone Maleate Film-coated tablet	NL-H-0208-01	BE220981	Teva Pharma Belgium NV; Laarstraat 16, Wilrijk 2610, Belgium	Belgium
Domperidone Instant Teva 10 mg Orodispergeerbare tabletten	BE-H-0211-001-DC	BE445225	Teva Pharma Belgium NV; Laarstraat 16, Wilrijk 2610, Belgium	Belgium
Domperidon Smelttablet Teva 10 mg, Orodispergeerbare tabletten	BE-H-0211-001-DC	RVG 112357	Teva Nederland B.V.	Netherlands
DOMPERIDONA ratiopharm 10 mg comprimidos	04/H/0179/001	5362181	Ratiopharm Lda; Portugal	Portugal

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
DOMPERIDONA ratiopharm 10 mg comprimidos	04/H/0179/001	5362280	Ratiopharm Lda; Portugal	Portugal
Motilium 10mg comprimidos recubiertos con película	Not applicable	55.410	Laboratorios del Dr. Esteve, S.A.	Spain
Motilium 1mg/ml suspensión oral	Not applicable	55.411	Laboratorios del Dr. Esteve, S.A.	Spain
Domperidone 10mg Tablets	Not Applicable	PL 16363/0106	Milpharm Limited	United Kingdom
Domperidona Aurovitas 10 mg Comprimido	06/H/0244/001	5053459; 5053467; 5053475; 5053509; 5053517; 5053525; 5053533; 5053541	Aurovitas Unipessoal, Lda.	Portugal
Domperidon Alternova	Not Applicable	36353	Alternova A/S	Denmark
DOMPERIDONE ARROW 10 mg, comprimé orodispersible	Not Applicable	35565	Arrow Génériques	France
DOMPERIDONE ARROW 10 mg, comprimé pelliculé	Not Applicable	27475	Arrow Génériques	France
DOMPERIDON SOFAR, 10mg, tablet	NL/H/0491/001	RVG30209	SOFAR S.p.A.	Netherlands

Product Name (in authorisation country)	MRP/DCP Authorisation number	Authorisation Number of product in the member state	MAH of product in the member state	Member State where product is authorised
Domperidon Pharmacin 10 mg, tabletten	NL/H/0241/001	RVG 23565	Pharmacin BV	Netherlands
Domperidon Aurobindo 10 mg, omhulde tabletten	Not Applicable	RVG 26065	Aurobindo Pharma B.V.	Netherlands
PERMOTIL 10mg, compresse	NL/H/0491/001	036670014; 036670026; 036670038; 036670040; 036670053.	SOFAR S.p.A.	Italy