13 December 2017
EMA/48730/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: glucosamine

Procedure no.: PSUSA/00001539/201703

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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
Alateris 625 mg compresse	SE/H/0560/001	038121063/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg compresse	SE/H/0560/001	038121012/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg compresse	SE/H/0560/001	038121024/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg compresse	SE/H/0560/001	038121036/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg compresse	SE/H/0560/001	038121048/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg compresse	SE/H/0560/001	038121051/M	LABORATOIRES EXPANSCIENCE	IT
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	UK
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	UK
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	UK
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	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
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	UK
Alateris 625 mg tablets	SE/H/0560/001	PL 32383/0001	LABORATOIRES EXPANSCIENCE	UK
ARAFISIO 1250 mg comprimidos efervescentes	not available	71701	ARAFARMA GROUP, S.A	ES
ARAFISIO 625 mg comprimidos efervescentes	not available	71702	ARAFARMA GROUP, S.A	ES
Arthryl 1,5 g jauhe oraaliliuosta varten	not available	12599	MEDA OY	FI
Arthryl 1,5 g milteliai geriamajam tirpalui	not available	LT/1/97/0224/001	MEDA PHARMA SIA	LT
Arthryl 1,5 g pulver till oral lösning	not available	12599	MEDA OY	FI
Arthryl 1,5 g pulveris iekšķīgi lietojama šķīduma pagatavošanai	not available	00-0145	MEDA PHARMA SIA	LV
Arthryl 750 mg apvalkotās tabletes	not available	09-0457	MEDA PHARMA SIA	LV

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
Arthryl 750 mg filmdragerade tabletter	not available	21203	MEDA OY	FI
Arthryl 750 mg plėvele dengtos tabletės	not available	LT/1/97/0224/002	MEDA PHARMA SIA	LT
Arthryl 750 mg plėvele dengtos tabletės	not available	LT/1/97/0224/003	MEDA PHARMA SIA	LT
Arthryl 750 mg tabletti, kalvopäällysteinen	not available	21203	MEDA OY	FI
Arthryl por belsőleges oldathoz tasakban	not available	OGYI-T-9015/02	ROTTAPHARM LIMITED	HU
Arthryl por belsőleges oldathoz tasakban	not available	OGYI-T-9015/01	ROTTAPHARM LIMITED	HU
Arthryl, 1,2 g suukaudse lahuse pulber	not available	334900	MEDA PHARMA SIA	EE
Arthryl, 1500 mg, proszek do sporządzania roztworu doustnego	not available	7727	MEDA PHARMA GMBH & CO. KG	PL
ARTHRYL, 589 mg õhukese polümeerikattega tabletid	not available	584608	MEDA PHARMA SIA	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
CARTISORB 1500 mg polvo para solución oral	not available	64.546	BIOIBÉRICA S.A.	ES
CODEROL 1.500 mg polvo para solución oral	not available	65.278	ALMIRALL, S.A.	ES
DONA	not available	29/110/00-C	MEDA PHARMA S.R.O.	CZ
DONA	not available	29/110/00-C	MEDA PHARMA S.R.O.	CZ
DONA	not available	29/110/00-C	MEDA PHARMA S.R.O.	CZ
DONA	not available	29/110/00-C	MEDA PHARMA S.R.O.	CZ
DONA 1178 mg prašek za peroralno raztopino	not available	HN/04/01828/001	MEDA PHARMA GMBH & CO. KG	SI
DONA 1500 mg polvere per soluzione orale	not available	026023061	MEDA PHARMA S.P.A.	IT
DONA 1500 mg polvere per soluzione orale	not available	026023061	MEDA PHARMA 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
Dona 1500 mg por belsőleges oldathoz	not available	OGYI-T- 4701/05	MEDA PHARMA HUNGARY KFT.	HU
Dona 1500 mg por belsőleges oldathoz	not available	OGYI-T-4701/04	MEDA PHARMA HUNGARY KFT.	HU
DONA 1500 mg pulbere pentru solutie orala	not available	1420/2009/01	ROTTAPHARM S.P.A.	RO
Dona 1500mg Powder for Oral Solution	not available	PA1332/54/1	MEDA HEALTH SALES IRELAND LIMITED	IE
DONA 250 mg capsule rigide	not available	026023010	MEDA PHARMA S.P.A.	IT
DONA 250 mg capsule rigide	not available	026023010	MEDA PHARMA S.P.A.	IT
DONA 250 mg compresse rivestite	not available	026023046	MEDA PHARMA S.P.A.	IT
DONA 250 mg compresse rivestite	not available	026023046	MEDA PHARMA S.P.A.	IT
Dona 250 mg kemény kapszula	not available	OGYI-T- 4701/01	MEDA PHARMA HUNGARY 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
dona 250 mg überzogene Tabletten	not available	6592176.00.00	MEDA PHARMA GMBH & CO. KG	DE
DONA 400 mg/3 ml concentrato per soluzione iniettabile e solvente per uso intramuscolare con lidocaina	not available	026023059	MEDA PHARMA S.P.A.	IT
DONA 400 mg/3 ml concentrato per soluzione iniettabile e solvente per uso intramuscolare con lidocaina	not available	026023059	MEDA PHARMA S.P.A.	IT
Dona 500mg Capsules	not available	PA1332/54/2	MEDA HEALTH SALES IRELAND LIMITED	IE
DONA 750 mg compresse rivestite con film	not available	026023085	MEDA PHARMA S.P.A.	IT
DONA 750 mg compresse rivestite con film	not available	026023097	MEDA PHARMA S.P.A.	IT
DONA 750 mg compresse rivestite con film	not available	026023085	MEDA PHARMA S.P.A.	IT
DONA 750 mg compresse rivestite con film	not available	026023097	MEDA PHARMA 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
Dona 750 mg filmtabletta	not available	OGYI-T-4701/07	MEDA PHARMA HUNGARY KFT.	HU
Dona 750 mg filmtabletta	not available	OGYI-T-4701/02	MEDA PHARMA HUNGARY KFT.	HU
Dona 750 mg filmtabletta	not available	OGYI-T-4701/03	MEDA PHARMA HUNGARY KFT.	HU
dona 750 mg Filmtabletten	not available	65462.00.00	MEDA PHARMA GMBH & CO. KG	DE
Dona Arthro 400 mg oldatos injekció	not available	OGYI-T- 4701/06	MEDA PHARMA HUNGARY KFT.	HU
DONA prášek pro přípravu perorálního roztoku	not available	29/118/97-C	MEDA PHARMA S.R.O.	CZ
DONA prášek pro přípravu perorálního roztoku	not available	29/118/97-C	MEDA PHARMA S.R.O.	CZ
DONA prášek pro přípravu perorálního roztoku	not available	29/118/97-C	MEDA PHARMA S.R.O.	CZ
DONA prášek pro přípravu perorálního roztoku	not available	29/118/97-C	MEDA PHARMA 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
dona® 750 mg Filmtabletten Wirkstoff: Glucosaminhemisulfat	not available	2009110012	MEDA PHARMA GMBH & CO. KG	LU
DONAROT 1500 mg κόνις μιας δόσης για πόσιμο υγρό	not available	70440/25-11-2016	MEDA PHARMA GMBH & CO. KG	GR
DONAROT 750 mg δισκία επικαλυμμένα με λεπτό υμένιο	not available	70439/25-11-2016	MEDA PHARMA GMBH & CO. KG	GR
donasan 750 mg Filmtabletten	not available	70067.00.00	MEDA PHARMA GMBH & CO. KG	DE
donasol 750 mg Filmtabletten	not available	70066.00.00	MEDA PHARMA GMBH & CO. KG	DE
Doppelherz® GLUCOSAMIN-HYDROCHLORID 750 mg TABLETTEN	SE/H/0886/001	76484.00.00	LABORATOIRES EXPANSCIENCE	DE
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 388 816 7 7	LABORATOIRES EXPANSCIENCE	FR
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 380 533 6 4	LABORATOIRES EXPANSCIENCE	FR
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 380 534 2 5	LABORATOIRES EXPANSCIENCE	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
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 380 535 9 3	LABORATOIRES EXPANSCIENCE	FR
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 380 536 5 4	LABORATOIRES EXPANSCIENCE	FR
FLEXEA 625 mg, comprimé	SE/H/0560/001	34009 380 537 1 5	LABORATOIRES EXPANSCIENCE	FR
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE
Flexove 625 mg tablets	SE/H/0560/001	PA 1616/1/1	LABORATOIRES EXPANSCIENCE	IE

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
Flexove 625 mg tabletta	SE/H/0560/001	OGYI-T- 20642/01	LABORATOIRES EXPANSCIENCE	HU
Flexove 625 mg tabletta	SE/H/0560/001	OGYI-T- 20642/02	LABORATOIRES EXPANSCIENCE	HU
Flexove 625 mg tabletta	SE/H/0560/001	OGYI-T- 20642/03	LABORATOIRES EXPANSCIENCE	HU
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	AT
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	AT
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	AT
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	AT
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	AT
Flexove 625 mg Tabletten	SE/H/0560/001	1-26831	LABORATOIRES EXPANSCIENCE	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
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tabletter	SE/H/0560/001	05-3642	LABORATOIRES EXPANSCIENCE	NO
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	CZ
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	CZ
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	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
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	CZ
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	CZ
Flexove 625 mg tablety	SE/H/0560/001	29/186/07-C	LABORATOIRES EXPANSCIENCE	CZ
Flexove, tabletki 625 mg	SE/H/0560/001	14425	LABORATOIRES EXPANSCIENCE	PL
Flexove, tabletki 625 mg	SE/H/0560/001	14425	LABORATOIRES EXPANSCIENCE	PL
Flexove, tabletki 625 mg	SE/H/0560/001	14425	LABORATOIRES EXPANSCIENCE	PL
Flexove, tabletki 625 mg	SE/H/0560/001	14425	LABORATOIRES EXPANSCIENCE	PL
Glucomed 625 mg comprimidos	SE/H/0560/001	5027446	LABORATOIRES EXPANSCIENCE	PT
Glucomed 625 mg comprimidos	SE/H/0560/001	5820386	LABORATOIRES EXPANSCIENCE	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
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletten	SE/H/0560/001	RVG 33081	LABORATOIRES EXPANSCIENCE	NL
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE

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
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE
Glucomed 625 mg tabletter	SE/H/0560/001	20971	LABORATOIRES EXPANSCIENCE	SE
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	SK
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	SK
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	SK
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	SK
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	SK
GLUCOMED 625 mg tablety	SE/H/0560/001	29/0007/07-S	LABORATOIRES EXPANSCIENCE	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
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed 625 mg töflur	SE/H/0560/001	IS/1/05/056/01	LABORATOIRES EXPANSCIENCE	IS
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	DK
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	DK
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	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
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	DK
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	DK
Glucomed, tabletter	SE/H/0560/001	38436	LABORATOIRES EXPANSCIENCE	DK
GLUCOSAMIL, πυκνό διάλυμα και διαλύτης για την παρασκευή ενέσιμου διαλύματος, 400 mg/amp	not available	92287/15/18-1-2016	VERISFIELD (UK) LTD	GR
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE

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
Glucosamine Navamedic 625 mg tabletter	SE/H/0886/001	41572	LABORATOIRES EXPANSCIENCE	SE
HESPERCORBIN 1500MG POLVO PARA SOLUCIÓN ORAL	not available	61105	ROTTAPHARM LIMITED	ES
VIARTRIL	not available	43342/22-09-2016	MEDA PHARMACEUTICALS S.A.	GR
VIARTRIL 1500mg Powder for Oral Solution	not available	PA 868/5/1	ROTTAPHARM LIMITED	IE
VIARTRIL 750 mg δισκία επικαλυμμένα με λεπτό υμένιο	not available	69755/22-09-2016	MEDA PHARMACEUTICALS S.A.	GR
VIARTRIL-S	not available	MA116/00201	ROTTAPHARM S.P.A.	MT
Viartril-S 250mg Cápsula	not available	8499913	LABORATÓRIOS DELTA, S.A.	PT
Viartril-S 250mg Cápsula	not available	8499921	LABORATÓRIOS DELTA, S.A.	PT
VIARTRIL-S, 400 mg/3 ml ,Solução injectável	not available	8535807	LABORATÓRIOS DELTA, S.A.	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
ДОНА 1500 mg Прах за перорален разтвор	not available	9600092	ROTTAPHARM S.P.A.	BG