10 March 2022
EMA/164942/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: fluticasone propionate / formoterol fumarate dihydrate

Procedure no.: PSUSA/00010339/202107

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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
Flutiform 125 Mikrogramm/5 Mikrogramm pro Sprühstoß Druckgasinhalation	SE/H/1921/002	1-31520	MUNDIPHARMA GES.M.B.H	AT
Flutiform 250 Mikrogramm/10 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	SE/H/1921/003	1-31521	MUNDIPHARMA GES.M.B.H	AT
Flutiform 50 Mikrogramm/5 Mikrogramm pro Sprühstoß Druckgasinhalation	SE/H/1921/001	1-31519	MUNDIPHARMA GES.M.B.H	AT
Flutiform 125 microgram / 5 microgram , aerosol, suspensie.	UK/H/2872/002	BE429414	MUNDIPHARMA PHARMACEUTICALS BV	BE
Flutiform 250 microgrammes/10 microgrammes , suspension pour inhalation en flacon pressurisé.	UK/H/2872/003	BE429423	MUNDIPHARMA PHARMACEUTICALS BV	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
Flutiform 50 microgram / 5 microgram , aërosol, suspensie.	UK/H/2872/001	BE429405	MUNDIPHARMA PHARMACEUTICALS BV	BE
Flutiform k-haler 125 microgram/5 microgram per dosis, aërosol, suspensie.	UK/H/2872/005	BE520951	MUNDIPHARMA PHARMACEUTICALS BV	BE
Flutiform k-haler 50 microgram/5 microgram per dosis, aërosol, suspensie.	UK/H/2872/004	BE520960	MUNDIPHARMA PHARMACEUTICALS BV	BE
Flutiform K-haler 125 microgram/5 microgram per actuation pressurised inhalation, suspension.	SE/H/1921/005	20170370	MUNDIPHARMA GES.M.B.H	BG
Flutiform K-haler 50 microgram/5 microgram per actuation pressurised inhalation, suspension.	SE/H/1921/004	20170369	MUNDIPHARMA GES.M.B.H	BG
ФЛУТИФОРМ 125 микрограма/5 микрограма на впръскване, суспензия под налягане за инхалация	SE/H/1921/002	20120518	MUNDIPHARMA GES.M.B.H	BG

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
ФЛУТИФОРМ 250 микрограма/10 микрограма на впръскване, суспензия под налягане за инхалация	SE/H/1921/003	20120519	MUNDIPHARMA GES.M.B.H	BG
ФЛУТИФОРМ 50 микрограма/5 микрограма на впръскване, суспензия под налягане за инхалация	SE/H/1921/001	20120517	MUNDIPHARMA GES.M.B.H	BG
Flutiform 250 μικρογραμμάρια/10 μικρογραμμάρια ανά ενεργοποίηση, εναιώρημα για εισπνοή υπό πίεση.	SE/H/1921/003	21397	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Flutiform K-haler 125 μικρογραμμάρια/5 μικρογραμμάρια ανά ψεκασμό, εναιώρημα για εισπνοή υπό πίεση.	SE/H/1921/005	022890	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Flutiform K-haler 50 μικρογραμμάρια/5 μικρογραμμάρια ανά ψεκασμό, εναιώρημα για εισπνοή υπό πίεση	SE/H/1921/004	022889	MUNDIPHARMA PHARMACEUTICALS LTD	CY

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
Flutiform125 μικρογραμμάρια/5 μικρογραμμάρια ανά ενεργοποίηση, εναιώρημα για εισπνοή υπό πίεση.	SE/H/1921/002	21396	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Flutiform50 μικρογραμμάρια/5 μικρογραμμάρια ανά ενεργοποίηση, εναιώρημα για εισπνοή υπό πίεση.	SE/H/1921/001	21395	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Flutiform 125 mikrogramů/5 mikrogramů v jedné dávce suspenze k inhalaci v tlakovém obalu	SE/H/1921/002	14/554/12-C	MUNDIPHARMA GES.M.B.H	CZ
Flutiform 250 mikrogramů/10 mikrogramů v jedné dávce suspenze k inhalaci v tlakovém obalu	SE/H/1921/003	14/555/12-C	MUNDIPHARMA GES.M.B.H	CZ
Flutiform 50 mikrogramů/5 mikrogramů v jedné dávce suspenze k inhalaci v tlakovém obalu	SE/H/1921/001	14/553/12-C	MUNDIPHARMA GES.M.B.H	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
Flutiform K-haler 50 mikrogramů/5 mikrogramů v jedné dávce suspenze k inhalaci v tlakovém obalu	SE/H/1921/004	14/808/15-C	MUNDIPHARMA GES.M.B.H	CZ
flutiform® 125 Mikrogramm/5 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	SE/H/1921/002	81898.00.00	MUNDIPHARMA GMBH	DE
flutiform® 250 Mikrogramm/10 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	UK/H/2872/003	81899.00.00	MUNDIPHARMA GMBH	DE
flutiform® 50 Mikrogramm/5 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	UK/H/2872/001	81897.00.00	MUNDIPHARMA GMBH	DE
Flutiform K-haler, inhalationsspray, suspension	SE/H/1921/004	56938	MUNDIPHARMA A/S	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
Flutiform K-haler, inhalationsspray, suspension	SE/H/1921/005	56939	MUNDIPHARMA A/S	DK
Flutiform, inhalationsspray, suspension	SE/H/1921/003	46691	MUNDIPHARMA A/S	DK
Flutiform, inhalationsspray, suspension	UK/H/2872/002	46690	MUNDIPHARMA A/S	DK
Flutiform, inhalationsspray, suspension	SE/H/1921/001	46689	MUNDIPHARMA A/S	DK
Flutiform 125 microgramos/5 microgramos/inhalación, suspensión para inhalación en envase a presión	SE/H/1921/002	78536	MUNDIPHARMA PHARMACEUTICALS SL	ES
Flutiform 250 microgramos/10 microgramos/inhalación, suspensión para inhalación en envase a presión.	SE/H/1921/003	78537	MUNDIPHARMA PHARMACEUTICALS SL	ES
Flutiform 50 microgramos/5 microgramos/inhalación,	SE/H/1921/001	78538	MUNDIPHARMA PHARMACEUTICALS SL	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
suspensión para inhalación en envase a presión				
Flutiform 125 mikrog/5 mikrog/annos inhalaatiosumute, suspensio.	SE/H/1921/002	28589	MUNDIPHARMA OY	FI
Flutiform 250 mikrog/10 mikrog/annos inhalaatiosumute, suspensio.	SE/H/1921/003	28590	MUNDIPHARMA OY	FI
Flutiform 50 mikrog/5 mikrog/annos inhalaatiosumute, suspensio.	SE/H/1921/001	28588	MUNDIPHARMA OY	FI
Flutiform K-haler 125 mikrog/5 mikrog/annos inhalaatiosumute, suspensio.	SE/H/1921/005	33801	MUNDIPHARMA OY	FI
Flutiform K-haler 50 mikrog/5 mikrog/annos inhalaatiosumute, suspensio.	SE/H/1921/004	33800	MUNDIPHARMA OY	FI
FLUTIFORM 125 microgrammes/5 microgrammes par dose, suspension pour	UK/H/2872/002	34009 266 488 5 5	MUNDIPHARMA	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
inhalation en flacon pressurisé				
FLUTIFORM 125 microgrammes/5 microgrammes par dose, suspension pour inhalation en flacon pressurisé	UK/H/2872/002	34009 266 488 5 5	MUNDIPHARMA	FR
FLUTIFORM 50 microgrammes/5 microgrammes par dose, suspension pour inhalation en flacon pressurisé	UK/H/2872/001	34009 266 487 9 4	MUNDIPHARMA	FR
FLUTIFORM 50 microgrammes/5 microgrammes par dose, suspension pour inhalation en flacon pressurisé	UK/H/2872/001	34009 266 487 9 4	MUNDIPHARMA	FR
Flutiform 125 mikrograma/5 mikrograma po potisku, stlačeni inhalat, suspenzija	SE/H/1921/002	HR-H-702532961	MUNDIPHARMA GESELLSCHAFT M.B.H.	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
Flutiform 250 mikrograma/10 mikrograma po potisku, stlačeni inhalat, suspenzija	SE/H/1921/003	HR-H-341378137	MUNDIPHARMA GESELLSCHAFT M.B.H.	HR
Flutiform 50 mikrograma/5 mikrograma po potisku, stlačeni inhalat, suspenzija	SE/H/1921/001	HR-H-452322131	MUNDIPHARMA GESELLSCHAFT M.B.H.	HR
Flutiform 125 mikrogramm/5 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/002	OGYI-T-23691/03	MUNDIPHARMA GES.M.B.H	HU
Flutiform 125 mikrogramm/5 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/002	OGYI-T-23691/04	MUNDIPHARMA GES.M.B.H	HU
Flutiform 250 mikrogramm/10 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/003	OGYI-T-23691/06	MUNDIPHARMA GES.M.B.H	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
Flutiform 250 mikrogramm/10 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/003	OGYI-T-23691/05	MUNDIPHARMA GES.M.B.H	HU
Flutiform 50 mikrogramm/5 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/001	OGYI-T-23691/01	MUNDIPHARMA GES.M.B.H	HU
Flutiform 50 mikrogramm/5 mikrogramm /adag túlnyomásos inhalációs szuszpenzió	SE/H/1921/001	OGYI-T-23691/02	MUNDIPHARMA GES.M.B.H	HU
Flutiform 50 microgram/5 microgram per metered dose pressurised inhalation, suspension.	UK/H/2872/001	PA 1688/13/1	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Flutiform 125 microgram/5 microgram per metered dose pressurised inhalation, suspension.	UK/H/2872/002	PA 1688/13/2	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Flutiform 250 microgram/10 microgram per metered dose	UK/H/2872/003	PA 1688/13/3	MUNDIPHARMA PHARMACEUTICALS LIMITED	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
pressurised inhalation, suspension.				
Flutiform K-haler 125 microgram/5 microgram per actuation pressurised inhalation, suspension.	SE/H/1921/005	PA 1688/013/005	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Flutiform K-haler 50 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/004	PA 1688/013/004	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Flutiform 125 míkrógrömm/5 míkrógrömm í úðaskammti innúðalyf, dreifa.	UK/H/2872/002	IS/1/12/096/02	MUNDIPHARMA A/S	IS
Flutiform 250 míkrógrömm/10 míkrógrömm í úðaskammti innúðalyf, dreifa.	UK/H/2872/003	IS/1/12/096/03	MUNDIPHARMA A/S	IS
Flutiform 50 míkrógrömm/5 míkrógrömm í úðaskammti innúðalyf, dreifa.	UK/H/2872/001	IS/1/12/096/01	MUNDIPHARMA A/S	IS

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
Abriff 125 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	UK/H/4379/002	042292021/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Abriff 250 microgrammi/10 microgrammi per erogazione, sospensione pressurizzata per inalazione	UK/H/4379/003	042292033/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Abriff 50 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione	UK/H/4379/001	042292019/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Affera 125 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	UK/H/4378/002	042293023/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Affera 250 microgrammi/10 microgrammi per erogazione, sospensione pressurizzata per inalazione.	UK/H/4378/003	042293035/M	MUNDIPHARMA PHARMACEUTICALS SRL	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
Affera 50 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	UK/H/4378/001	042293011/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiformo 125 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	SE/H/1921/002	042294025	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiformo 125 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	SE/H/1921/002	042294052	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiformo 250 microgrammi/10 microgrammi per erogazione, sospensione pressurizzata per inalazione.	SE/H/1921/003	042294064	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiformo 250 microgrammi/10 microgrammi per erogazione, sospensione	SE/H/1921/003	042294037	MUNDIPHARMA PHARMACEUTICALS SRL	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
pressurizzata per inalazione.				
Flutiformo 50 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	SE/H/1921/001	042294013	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiformo 50 microgrammi/5 microgrammi per erogazione, sospensione pressurizzata per inalazione.	SE/H/1921/001	042294049	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Flutiform 125 microgrammes/5 microgrammes , suspension pour inhalation en flacon pressurisé.	SE/H/1921/002	2013040097	MUNDIPHARMA PHARMACEUTICALS BV	LU
Flutiform 250 microgrammes/10 microgrammes , suspension pour inhalation en flacon pressurisé.	SE/H/1921/003	2013040098	MUNDIPHARMA PHARMACEUTICALS BV	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
Flutiform 50 microgrammes/5 microgrammes , suspension pour inhalation en flacon pressurisé.	SE/H/1921/001	2013040096	MUNDIPHARMA PHARMACEUTICALS BV	LU
Flutiform k-haler 125 microgrammes/5 microgrammes par dose, suspension pour inhalation en flacon pressurisé	SE/H/1921/005	2018060172	MUNDIPHARMA PHARMACEUTICALS BV	LU
Flutiform k-haler 50 microgrammes/5 microgrammes par dose, suspension pour inhalation en flacon pressurisé	SE/H/1921/005	2018060171	MUNDIPHARMA PHARMACEUTICALS BV	LU
Flutiform 125 microgram / 5 microgram / dosis aërosol, suspensie.	UK/H/2872/002	RVG 107242	MUNDIPHARMA PHARMACEUTICALS BV	NL
Flutiform 250 microgram / 10 microgram / dosis aërosol, suspensie.	UK/H/2872/003	RVG 107243	MUNDIPHARMA PHARMACEUTICALS BV	NL
Flutiform 50 microgram / 5 microgram / dosis aërosol, suspensie.	UK/H/2872/001	RVG 107222	MUNDIPHARMA PHARMACEUTICALS BV	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
Flutiform 125 mikrogram/ 5 mikrogram per dose, inhalasjonsaerosol, suspensjon.	UK/H/2872/002	10-7475	MUNDIPHARMA AS	NO
Flutiform 250 mikrogram/ 10 mikrogram per dose, inhalasjonsaerosol, suspensjon.	UK/H/2872/003	10-7476	MUNDIPHARMA AS	NO
Flutiform 50 mikrogram/ 5 mikrogram per dose, inhalasjonsaerosol, suspensjon.	UK/H/2872/001	10-7474	MUNDIPHARMA AS	NO
Flutiform K-haler 125 mikrogram/5 mikrogram per dose inhalasjonsaerosol, suspensjon	SE/H/1921/005	15-10966	MUNDIPHARMA AS	NO
Flutiform K-haler 50 mikrogram/5 mikrogram per dose inhalasjonsaerosol, suspensjon	SE/H/1921/004	15-10965	MUNDIPHARMA AS	NO
Flutiform 125 microgramas/5 microgramas suspensão pressurizada para inalação.	UK/H/2872/002	5488234	MUNDIPHARMA FARMACÊUTICA, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Flutiform 250 microgramas/10 microgramas suspensão pressurizada para inalação.	UK/H/2872/003	5488242	MUNDIPHARMA FARMACÊUTICA, LDA.	PT
Flutiform 50 microgramas/5 microgramas suspensão pressurizada para inalação.	UK/H/2872/001	5488226	MUNDIPHARMA FARMACÊUTICA, LDA.	PT
Flutiform K-haler 125 microgramas/5 microgramas suspensão pressurizada para inalação.	SE/H/1921/005	5733357	MUNDIPHARMA FARMACÊUTICA, LDA.	PT
Flutiform K-haler 50 microgramas/5 microgramas suspensão pressurizada para inalação.	SE/H/1921/004	5733365	MUNDIPHARMA FARMACÊUTICA, LDA.	PT
Flutiform 125 mcg/5mcg suspensie de inhalat presurizata	SE/H/1921/002	10612/2018/01-02	MUNDIPHARMA GES.M.B.H	RO
Flutiform 250 microgram/10 microgram suspensie de inhalat presurizată.	SE/H/1921/003	10613/2018/01	MUNDIPHARMA GES.M.B.H	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
Flutiform 250 micrograme/10 micrograme suspensie de inhalat presurizată.	SE/H/1921/003	10613/2018/02	MUNDIPHARMA GES.M.B.H	RO
Flutiform 50 mcg/5mcg suspensie de inhalat presurizata	SE/H/1921/001	10611/2018/01-02	MUNDIPHARMA GES.M.B.H	RO
Abriff 125 microgram/5 microgram per actuation pressurised inhalation, suspension	UK/H/4379/02/E/02	58823	MUNDIPHARMA AB	SE
Abriff 50 microgram/5 microgram per actuation pressurised inhalation, suspension	UK/H/4379/01/E/02	58822	MUNDIPHARMA AB	SE
Abriff, 250 mikrogram/10 mikrogram/puff inhalationsspray, suspension	UK/H/4379/03/E/02	58824	MUNDIPHARMA AB	SE
Affera 125 microgram/5 microgram per actuation pressurised inhalation, suspension	UK/H/4378/02/E/02	58826	MUNDIPHARMA AB	SE
Affera 50 microgram/5 microgram per actuation pressurised inhalation, suspension	UK/H/4378/01/E/02	58825	MUNDIPHARMA AB	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
Affera, 250 mikrogram/10 mikrogram/puff inhalationsspray, suspension	UK/H/4378/01/E/02	58827	MUNDIPHARMA AB	SE
Flutiform 125 mikrogram/5 mikrogram per puff, inhalationsspray, suspension.	UK/H/2872/002	44065	MUNDIPHARMA AB	SE
Flutiform 250 mikrogram/10 mikrogram per puff, inhalationsspray, suspension.	UK/H/2872/003	44066	MUNDIPHARMA AB	SE
Flutiform 50 mikrogram/5 mikrogram per puff, inhalationsspray, suspension.	UK/H/2872/001	44064	MUNDIPHARMA AB	SE
Flutiform K-haler 125 mikrogram/5 mikrogram per puff, inhalationsspray, suspension.	SE/H/1921/005	53719	MUNDIPHARMA AB	SE
Flutiform K-haler 50 mikrogram/5 mikrogram per puff, inhalationsspray, suspension.	SE/H/1921/004	53718	MUNDIPHARMA AB	SE
flutiform 125 mikrogramov/5 mikrogramov/potisk	SE/H/1921/002	H/20/02747/003	MUNDIPHARMA GESELLSCHAFT M.B.H.	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
inhalacijska suspenzija pod tlakom				
flutiform 125 mikrogramov/5 mikrogramov/potisk inhalacijska suspenzija pod tlakom	SE/H/1921/002	H/20/02747/004	MUNDIPHARMA GESELLSCHAFT M.B.H.	SI
flutiform 250 mikrogramov/10 mikrogramov/potisk inhalacijska suspenzija pod tlakom	SE/H/1921/003	H/20/02747/005	MUNDIPHARMA GESELLSCHAFT M.B.H.	SI
flutiform 250 mikrogramov/10 mikrogramov/potisk inhalacijska suspenzija pod tlakom	SE/H/1921/003	H/20/02747/006	MUNDIPHARMA GESELLSCHAFT M.B.H.	SI
flutiform 50 mikrogramov/5 mikrogramov/potisk inhalacijska suspenzija pod tlakom	SE/H/1921/001	H/20/02747/001	MUNDIPHARMA GESELLSCHAFT M.B.H.	SI
flutiform 50 mikrogramov/5 mikrogramov/potisk inhalacijska suspenzija	SE/H/1921/001	H/20/02747/002	MUNDIPHARMA GESELLSCHAFT M.B.H.	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
pod tlakom				
Flutiform 125 microgram/5 microgram	UK/H/2872/002	14/0357/12-S	MUNDIPHARMA GES.M.B.H	SK
Flutiform 250 mikrogramov/10 mikrogramov	UK/H/2872/003	14/0358/12-S	MUNDIPHARMA GES.M.B.H	SK
Flutiform 50 mikrogramov/5 mikrogramov	UK/H/2872/001	14/0356/12-S	MUNDIPHARMA GES.M.B.H	SK
Affera® 125 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/4378/002	PL 16950/0186	NAPP PHARMACEUTICALS LTD	XI
Affera® 250 microgram/10 microgram per actuation pressurised inhalation, suspension.	UK/H/4378/003	PL 16950/0187	NAPP PHARMACEUTICALS LTD	XI
Affera® 50 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/4378/001	PL 16950/0185	NAPP PHARMACEUTICALS LTD	XI

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
Flutiform K-haler 125 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/005	PL 16950/0339	NAPP PHARMACEUTICALS LTD	XI
Flutiform K-haler 50 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/004	PL 16950/0338	NAPP PHARMACEUTICALS LTD	XI
Flutiform® 125 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/002	PL 16950/0168	NAPP PHARMACEUTICALS LTD	XI
Flutiform® 250 microgram/10 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/003	PL 16950/0169	NAPP PHARMACEUTICALS LTD	XI
Flutiform® 50 microgram/5 microgram per actuation pressurised inhalation, suspension.	UK/H/2872/001	PL 16950/0167	NAPP PHARMACEUTICALS LTD	XI