EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

11 June 2020
EMA/326863/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: fluticasone / salmeterol (for nationally authorised products)

Procedure no.: PSUSA/00001455/201910

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
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
ЕРФЛУЗАЛ ФОРСПИРО 50 МИКРОГРАМА + 250 МИКРОГРАМА /ДОЗА ПРАХ ЗА ИНХАЛАЦИЯ, ПРЕДВАРИТЕЛНО ДОЗИРАН	SE/H/1323/003	20150137	SANDOZ PHARMACEUTICALS D.D.	BG
AirFluSol Forspiro 50 mikrogramm/250 mikrogramm/adag adagolt inhalációs por	SE/H/1323/003	OGYI-T-22596/05	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/250 mikrogramm/adag adagolt inhalációs por	SE/H/1323/003	OGYI-T-22596/04	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/250 mikrogramm/adag adagolt inhalációs por	SE/H/1323/003	OGYI-T-22596/06	SANDOZ HUNGÁRIA KFT	HU
AirFluSal® Forspiro® 50 microgame/250 microgame/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/01	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgame/250 microgame/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/02	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgame/250 microgame/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/03	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50	SE/H/1323/003	7673/2015/04	S.C. SANDOZ S.R.L.	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
microgram/250 microgram/ doza, pulbere unidoză de inhalat				
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/05	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgram/250 microgram/doză pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/06	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/07	S.C. SANDOZ S.R.L.	RO
Salmeterol/Fluticasone Sandoz 50 mikrogram/250 mikrogram/dos Inhalationspulver, avdelad dos	SE/H/1323/003	50816	SANDOZ A/S	SE
AirFluSal MDI 25 microgram/125 microgram per actuation pressurised inhalation, suspension	NL/H/3708/001	PL 04416/1475	SANDOZ LTD	UK
AirFluSal® MDI 25 microgram/250 microgram per actuation pressurised inhalation,	NL/H/3708/002	PL 04416/1476	SANDOZ 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
suspension				
Salmeterol/Fluticasonpropionaat Sandoz 25/125 microgram, aerosol, suspensie	NL/H/3708/001	RVG 118834	SANDOZ B.V.	NL
Salmeterol/Fluticasonpropionaat Sandoz 25/250 microgram, aerosol, suspensie	NL/H/3708/002	RVG 118835	SANDOZ B.V.	NL
PROPIONATE DE FLUTICASONE/SALMETEROL MYLAN 100 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	not available	NL42622	MYLAN S.A.S	FR
PROPIONATE DE FLUTICASONE/SALMETEROL MYLAN 250 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	not available	NL42623	MYLAN S.A.S	FR
PROPIONATE DE FLUTICASONE/SALMETEROL MYLAN 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	not available	NL42624	MYLAN S.A.S	FR
Samtoral levis 25 Mikrogramm/50 Mikrogramm pro Sprühstoß -	DE/H/5771/001	1-23953	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	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
Druckgasinhalation				
Viani mite Dosier-Aerosol 25 µg/50 µg Druckgasinhalation, Suspension	DE/H/5771/001	50681.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani Evohaler 25 microgram /50 microgram per metered dose pressurised inhalation, suspension	DE/H/5771/001	PL 10949/0341	GLAXO WELLCOME UK LIMITED	UK
Samtoral standard 25 Mikrogramm/125 Mikrogramm pro Sprühstoß - Druckgasinhalation	DE/H/5771/002	1-23954	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	AT
Viani Dosier-Aerosol 25 µg/125 µg Druckgasinhalation, Suspension	DE/H/5771/002	50681.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani Evohaler 25 microgram /125 microgram per metered dose pressurised inhalation, suspension	DE/H/5771/002	PL 10949/0342	GLAXO WELLCOME UK LIMITED	UK
Samtoral forte 25 Mikrogramm/250 Mikrogramm pro Sprühstoß - Druckgasinhalation	DE/H/5771/003	1-23955	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	AT
Viani forte Dosier-Aerosol 25 µg/250 µg Druckgasinhalation, Suspension	DE/H/5771/003	50681.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani Evohaler 25	DE/H/5771/003	PL 10949/0343	GLAXO WELLCOME UK	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
microgram /250 microgram per metered dose pressurised inhalation, suspension			LIMITED	
Airflusal® Forspiro® 50 microgramos/250 microgramos/inhalación, polvo para inhalación (unidosis)	SE/H/1446/001	80248	SANDOZ FARMACÉUTICA, S.A.	ES
Airflusal® Forspiro® 50 microgramos/500 microgramos/inhalación, polvo para inhalación (unidosis)	SE/H/1446/002	80249	SANDOZ FARMACÉUTICA, S.A.	ES
Mesaflu Forspiro	SE/H/1446/002	51635	SANDOZ A/S	SE
Mesaflu Forspiro	SE/H/1446/001	51634	SANDOZ A/S	SE
PROPIONATE DE FLUTICASONE/SALMETEROL GSK 50 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	DE/H/5772/001	NL41205	LABORATOIRE GLAXOSMITHKLINE	FR
PROPIONATE DE FLUTICASONE/SALMETEROL GSK 125 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	DE/H/5772/002	NL41206	LABORATOIRE GLAXOSMITHKLINE	FR
PROPIONATE DE FLUTICASONE/SALMETE	DE/H/5772/003	NL41207	LABORATOIRE GLAXOSMITHKLINE	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
ROL GSK 250 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé				
Salmeterol/Fluticasonpropionat Dosier-Aerosol Cascan 25 Mikrogramm/50 Mikrogramm Druckgasinhalation, Suspension	DE/H/5772/001	85136.00.00	CASCAN GMBH & CO.KG	DE
SLFP Inhaler 25 microgram/50 microgram per metered dose pressurised inhalation, suspension	DE/H/5772/001	PL 10949/0376	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Salmeterol/Fluticasonpropionat Dosier-Aerosol Cascan 25 Mikrogramm/125 Mikrogramm Druckgasinhalation, Suspension	DE/H/5772/02	85137.00.00	CASCAN GMBH & CO.KG	DE
SLFP Inhaler 25 microgram/125 microgram per metered dose pressurised inhalation, suspension	DE/H/5772/002	PL 10949/0377	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Salmeterol/Fluticasonpropionat Dosier-Aerosol Cascan 25 Mikrogramm/250	DE/H/5772/03	85138.00.00	CASCAN GMBH & CO.KG	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
Mikrogramm Druckgasinhalation, Suspension				
SLFP Inhaler 25 microgram/250 microgram per metered dose pressurised inhalation, suspension	DE/H/5772/003	PL 10949/0378	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Salflucombi Easyhaler 50 microgramos/500 microgramos/inhalación, polvo para inhalación	SE/H/1699/002	83799	ORION CORPORATION	ES
Salflucombi Easyhaler, 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1699/002	56292	ORION CORPORATION	SE
Anasma 25 microgramos/125 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	63.924	ALLEN FARMACÉUTICA, S.A	ES
Anasma 25 microgramos/250 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	63.925	ALLEN FARMACÉUTICA, S.A	ES
Anasma 25 microgramos/50 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	63.923	ALLEN FARMACÉUTICA, S.A	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
Salflumix Easyhaler 50 mikrogram/250 mikrogram/dos inhalationspulver	SE/H/1692/001	56277	ORION CORPORATION	SE
Salflumix Easyhaler 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1692/002	56278	ORION CORPORATION	SE
Salflumix Easyhaler 50 mikrogrami/250 mikrogrami/devā inhalācijas pulveris	SE/H/1692/01/DC	18-0072	ORION CORPORATION	LV
Salflumix Easyhaler 50 mikrogrami/500 mikrogrami/devā inhalācijas pulveris	SE/H/1692/01/DC	18-0073	ORION CORPORATION	LV
Safumix Easyhaler 50 mikrogramm/250 mikrogramm/adag, inhalációs por	SE/H/1692/001	OGYI-T-23372/01	ORION CORPORATION	HU
Safumix Easyhaler 50 mikrogramm/500 mikrogramm/adag, inhalációs por	SE/H/1692/001	OGYI-T-23372/02	ORION CORPORATION	HU
Salflumix Easyhaler, inhalationspulver	SE/H/1692/001	59399	ORION CORPORATION	DK
Salflumix Easyhaler, inhalationspulver	SE/H/1692/001	59400	ORION CORPORATION	DK
Salflumix Easyhaler 50 míkróg/250 míkróg/skammt innöndunarduft	SE/H/1692/01/DC	IS/1/18/048/01	ORION CORPORATION	IS
Salflumix Easyhaler 50 míkróg/500	SE/H/1692/001	IS/1/18/048/02	ORION CORPORATION	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
míkróg/skammt innöndunarduft				
Salflumix Easyhaler 50 mikrog/250 mikrog/annos, inhalaatiojauhe	SE/H/1692/001	35164	ORION CORPORATION	FI
Salflumix Easyhaler 50 mikrog/500 mikrog/annos, inhalaatiojauhe	SE/H/1692/002	35165	ORION CORPORATION	FI
Salflumix Easyhaler 50 mikrogram/250 mikrogram/dos, inhalationspulver	SE/H/1692/001	35164	ORION CORPORATION	FI
Salflumix Easyhaler 50 mikrogram/500 mikrogram/dos, inhalationspulver	SE/H/1692/002	35165	ORION CORPORATION	FI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/dávka inhalacný prášok	SE/H/1692/001	14/0155/18-S	ORION CORPORATION	SK
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/dávka inhalačný prášok	SE/H/1692/001	14/0156/18-S	ORION CORPORATION	SK
Salflumix Easyhaler 50 mikrogrammi/250 mikrogrammi annuses, inhalatsioonipulber	SE/H/1692/001	968318	ORION CORPORATION	EE
Salflumix Easyhaler 50 mikrogrammi/500 mikrogrammi annuses, inhalatsioonipulber	SE/H/1692/002	968418	ORION CORPORATION	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
Salflumix Easyhaler 50 mikrogram/250 mikrogram per dose inhalasjonspulver	SE/H/1692/001	17-11669	ORION CORPORATION	NO
Salflumix Easyhaler 50 mikrogram/500 mikrogram per dose inhalasjonspulver	SE/H/1692/002	17-11670	ORION CORPORATION	NO
Flutisamix Easyhaler 50 microgrammes/250 microgrammes/dose, poudre pour inhalation	SE/H/1692/001	BE530675	ORION CORPORATION	BE
Flutisamix Easyhaler 50 microgram/250 microgram/dosis, inhalatiepoeder	SE/H/1692/001	BE530675	ORION CORPORATION	BE
Flutisamix Easyhaler 50 Mikrogramm/250 Mikrogramm/Dosis, Pulver zur Inhalation	SE/H/1692/001	BE530675	ORION CORPORATION	BE
Flutisamix Easyhaler 50 microgrammes/500 microgrammes/dose, poudre pour inhalation	SE/H/1692/002	BE530684	ORION CORPORATION	BE
Flutisamix Easyhaler 50 microgram/500 microgram/dosis, inhalatiepoeder	SE/H/1692/002	BE530684	ORION CORPORATION	BE
Flutisamix Easyhaler 50 Mikrogramm/500 Mikrogramm/Dosis, Pulver zur Inhalation	SE/H/1692/002	BE530684	ORION CORPORATION	BE
Salflumix Easyhaler 500 microgramas/dose + 50	SE/H/1692/002	5747027	ORION CORPORATION	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
microgramas/dose, pó para inalação				
Salflumix Easyhaler 250 microgramas/dose + 50 microgramas/dose, pó para inalação	SE/H/1692/01	5747035	ORION CORPORATION	PT
Salflumix Easyhaler 50 míkróg/500 míkróg/skammt innöndunarduft	SE/H/1692/002	14/151/17-C	ORION CORPORATION	CZ
Salflumix Easyhaler 50 míkróg/250 míkróg/skammt innöndunarduft	SE/H/1692/01	14/150/17-C	ORION CORPORATION	CZ
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/001-012	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/001-012	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/007	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/001	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/002	ORION CORPORATION	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
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/003	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/004	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/005	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/001	H/18/02477/006	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/008	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/009	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/0010	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje	SE/H/1692/002	H/18/02477/0011	ORION CORPORATION	SI
Salflumix Easyhaler 50 mikrogramov/500	SE/H/1692/002	H/18/02477/0012	ORION CORPORATION	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
mikrogramov/odmerek prašek za inhaliranje				
Salflumix Easyhaler, (50 mikrogramów + 500 mikrogramów)/dawkę odmierzoną, proszek do inhalacji	SE/H/1692/002	24929	ORION CORPORATION	PL
Salflumix Easyhaler, (50 mikrogramów + 250 mikrogramów)/dawkę odmierzoną, proszek do inhalacji	SE/H/1692/001	24928	ORION CORPORATION	PL
Flutisamix Easyhaler 50 microgrammes/250 microgrammes/dose, poudre pour inhalation	SE/H/1692/001	2018110336	ORION CORPORATION	LU
Flutisamix Easyhaler 50 microgrammes/500 microgrammes/dose, poudre pour inhalation	SE/H/1692/002	2018110337	ORION CORPORATION	LU
Flutisamix Easyhaler 50 Mikrogramm/250 Mikrogramm/Dosis, Pulver zur Inhalation	SE/H/1692/001	2018110336	ORION CORPORATION	LU
Flutisamix Easyhaler 50 Mikrogramm/500 Mikrogramm/Dosis, Pulver zur Inhalation	SE/H/1692/002	2018110337	ORION CORPORATION	LU
Fusamix Easyhaler 50 mikrograma + 500 mikrograma u jednoj dozi, prašak inhalata	SE/H/1692/002	HR-H-554692844	ORION CORPORATION	HR
Fusamix Easyhaler 50 mikrograma + 250	SE/H/1692/001	HR-H-684471832	ORION CORPORATION	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
mikrograma u jednoj dozi, prašak inhalata				
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/001	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/002	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/003	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/004	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/005	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/250 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/001	LT/1/18/4265/006	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/001	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/002	ORION CORPORATION	LT

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Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/003	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/004	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/005	ORION CORPORATION	LT
Fusamix Easyhaler 50 mikrogramų/500 mikrogramų/dozėje įkvepiamieji milteliai	SE/H/1692/002	LT/1/18/4266/006	ORION CORPORATION	LT
Safumix 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1692/001	045960010	ORION CORPORATION	IT
Safumix 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1692/001	045960022	ORION CORPORATION	IT
Safumix 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1692/001	045960034	ORION CORPORATION	IT
Safumix 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1692/001	045960046	ORION CORPORATION	IT
Safumix 50 microgrammi/250	SE/H/1692/001	045960059	ORION CORPORATION	IT

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microgrammi/dose, polvere per inalazione				
Safumix 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1692/001	045960061	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960073	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960085	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960097	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960109	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960111	ORION CORPORATION	IT
Safumix 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1692/002	045960123	ORION CORPORATION	IT
Saflutin 50/100 mikrogrami/devā inhalācijas pulveris, dozēts	not available	17-0175	UAB INTELI GENERICS NORD	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
Saflutin 50/250 mikrogrami/devā inhalācijas pulveris, dozēts	not available	17-0176	UAB INTELI GENERICS NORD	LV
Saflutin 50/500 mikrogrami/devā inhalācijas pulveris, dozēts	not available	17-0177	UAB INTELI GENERICS NORD	LV
Ventonorm Forspiro, 50 microgram/250 microgram/dose, inhalation powder, predispensed	SE/H/1412/001	50818	SANDOZ A/S	SE
Ventonorm Forspiro, 50 microgram/500 microgram/dose, inhalation powder, predispensed	SE/H/1412/002	50819	SANDOZ A/S	SE
FLUSALIO 50 microgrammi/250 microgrammi/dose polvere per inalazione in contenitore monodose	DE/H/4378/001	041893013	ELPEN PHARMACEUTICAL CO. INC.	IT
FLUSALIO 50 microgrammi/500 microgrammi/dose polvere per inalazione in contenitore monodose	DE/H/4378/002	041893025	ELPEN PHARMACEUTICAL CO. INC.	IT
Flusalio 50 Mikrogramm/250 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	DE/H/4378/001	84254.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Flusalio 50	DE/H/4378/002	84255.00.00	ELPEN PHARMACEUTICAL	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
Mikrogramm/500 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation			CO. INC.	
Salmeterol/Fluticasonpropionaat Elpen 50 microgram/250 microgram/dosis inhalatiepoeder, voorverdeeld	SE/H/1190/002	RVG 110377	ELPEN PHARMACEUTICAL CO. INC.	NL
Salmeterol/Fluticasonpropionaat Elpen 50 microgram/500 microgram/dosis inhalatiepoeder, voorverdeeld	SE/H/1190/003	RVG 110378	ELPEN PHARMACEUTICAL CO. INC.	NL
Salmeson 50 Mikrogramm/250 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	SE/H/1190/002	1-31952	ELPEN PHARMACEUTICAL CO. INC.	AT
Salmeson 50 Mikrogramm/500 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	SE/H/1190/003	1-31953	ELPEN PHARMACEUTICAL CO. INC.	AT
Flusalio 50 microgrammes /250 microgrammes/dose, poudre pour inhalation en récipient unidose.	SE/H/1190/002	BE437622	ELPEN PHARMACEUTICAL CO. INC.	BE
Flusalio 50 microgrammes /500 microgrammes /dose,	SE/H/1190/003	BE437656	ELPEN PHARMACEUTICAL CO. INC.	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
poudre pour inhalation en récipient unidose.				
Flamerio 50 microgram/250 microgram/dose inhalation powder, pre-dispensed.	SE/H/1190/002	PA1879/001/001	ELPEN PHARMACEUTICAL CO. INC.	IE
Flamerio 50 microgram/500 microgram/dose inhalation powder, pre-dispensed.	SE/H/1190/003	PA1879/001/002	ELPEN PHARMACEUTICAL CO. INC.	IE
Flusaterol, inhalationspulver, afdelt	SE/H/1190/002	49572	ELPEN PHARMACEUTICAL CO. INC.	DK
Flusaterol, inhalationspulver, afdelt	SE/H/1190/003	49573	ELPEN PHARMACEUTICAL CO. INC.	DK
Flusaterol 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1190/002	46637	ELPEN PHARMACEUTICAL CO. INC.	SE
Flusaterol 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1190/003	46638	ELPEN PHARMACEUTICAL CO. INC.	SE
Sereflo 25 microgram/125 microgram per actuation pressurised inhalation, suspension.	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	IE
Sereflo 25 microgram/250 microgram per actuation	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	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.				
Sereflo 25 microgram/125 microgram per actuation pressurised inhalation, suspension.	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	IE
Sereflo 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	IE
Seretide Evohaler 25 mikrogram / 50 mikrogram/dos inhalationsspray, suspension	DE/H/5770/001	16181	GLAXOSMITHKLINE OY	FI
Seretide Evohaler 25 mikrogram / 125 mikrogram/dos inhalationsspray, suspension	DE/H/5770/002	16182	GLAXOSMITHKLINE OY	FI
Seretide Evohaler 25 mikrogram / 250 mikrogram/dos inhalationsspray, suspension	DE/H/5770/003	16183	GLAXOSMITHKLINE OY	FI
Seretide 25 microgram/50 microgram per afgemeten dosis aërosol, suspensie.	DE/H/5770/001	BE220683	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide 25 Mikrogramm/50	DE/H/5770/001	BE220683	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension				
Seretide 25 microgram/125 microgram per afgemeten dosis aërosol, suspensie.	DE/H/5770/002	BE220692	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide 25 Mikrogramm/125 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5770/002	BE220692	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide 25 microgram/250 microgram per afgemeten dosis aërosol, suspensie.	DE/H/5770/003	BE220701	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide 25 Mikrogramm/250 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5770/003	BE220701	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide 25 Mikrogramm/125 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5770/002	260/11/01/0949	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide 25 Mikrogramm/250	DE/H/5770/003	260/11/01/0950	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension				
Seretide 25 Mikrogramm/50 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5770/001	260/11/01/0954	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide levis 25 Mikrogramm/50 Mikrogramm pro Sprühstoß - Druckgasinhalation	DE/H/5770/001	1-23942	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seretide 25 microgrammes/50 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5770/001	BE220683	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide	DE/H/5770/001	31803	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Evohaler 25 mikrog/50 mikrog/annos inhalaatiosumute, suspensio	DE/H/5770/001	16181	GLAXOSMITHKLINE OY	FI
SERETIDE 50 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé avec valve doseuse	DE/H/5770/001	NL26158	LABORATOIRE GLAXOSMITHKLINE	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
Seretide Inhaler 25 µ??????aµµ???a /50 µ??????aµµ???a a?? ?a????sµ??? d?s? e?a????µat?? ??a e?sp??? ?p? p?es?.	DE/H/5770/001	2439204	GLAXOSMITHKLINE AEBE	GR
Seretide 25 míkróg/50 míkróg í afmældum skammti af innúðalyfi, dreifu	DE/H/5770/001	IS/1/00/035/01	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide 50 Evohaler 25 microgram/50 microgram per metered dose pressurised inhalation, suspension	DE/H/5770/001	PA 1077/46/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seretide 25 microgrammi/50 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5770/001	034371106	GLAXOSMITHKLINE S.P.A.	IT
Seretide 25 microgrammes/50 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5770/001	260/11/01/0954	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide 25/50 Inhalator CFK-vrij, aërosol, suspensie 25 microgram/50 microgram/dosis	DE/H/5770/001	RVG 25865	GLAXOSMITHKLINE B.V.	NL
Seretaide Inalador, 25 microgramas/50	DE/H/5770/001	3512688	GLAXO WELLCOME 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
microgramas por dose calibrada, suspensão pressurizada para inalação				
Seretide 25 microgramos/50 microgramos/inhalación, suspensión para inhalación en envase a presión	DE/H/5770/001	63.796	GLAXOSMITHKLINE S.A.	ES
Seretide Evohaler mite 25 mikrogram/50 mikrogram/ dos inhalationsspray, suspension	DE/H/5770/001	19165	GLAXOSMITHKLINE AB	SE
Seretide Evohaler 25 microgram /50 microgram per metered dose pressurised inhalation, suspension	DE/H/5770/001	PL 10949/0337	GLAXO WELLCOME UK LIMITED	UK
Seretide standard 25 Mikrogramm/125 Mikrogramm pro Sprühstoß - Druckgasinhalation	DE/H/5770/002	1-23943	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seretide 25 microgrammes/125 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5770/002	BE220692	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide	DE/H/5770/002	31804	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Evohaler 25	DE/H/5770/002	16182	GLAXOSMITHKLINE OY	FI

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
mikrog/125 mikrog/annos inhalaatiosumute, suspensio				
SERETIDE 125 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé avec valve doseuse	DE/H/5770/002	NL26159	LABORATOIRE GLAXOSMITHKLINE	FR
Seretide Inhaler 25 µ??????aµµ???a /125 µ??????aµµ???a a?? ?a????sµ??? d?s? e?a????µat?? ??a e?sp??? ?p? p?es?.	DE/H/5770/002	2439205	GLAXOSMITHKLINE AEBE	GR
Seretide 25 míkróg/125 míkróg í afmældum skammti af innúðalyfi, dreifu	DE/H/5770/002	IS/1/00/035/02	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide 125 Evohaler 25 microgram/125 microgram per metered dose pressurised inhalation, suspension.	DE/H/5770/002	PA 1077/46/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seretide 25 microgrammi/125 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5770/002	034371118	GLAXOSMITHKLINE S.P.A.	IT
Seretide 25 microgrammes/125	DE/H/5770/002	260/11/01/0949	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé				
Seretide 25/125 Inhalator CFK-vrij, aërosol, suspensie 25 microgram/125 microgram/dosis	DE/H/5770/002	RVG 25866	GLAXOSMITHKLINE B.V.	NL
Seretaide Inalador, 25 microgramas/125 microgramas por dose calibrada, suspensão pressurizada para inalação	DE/H/5770/002	3512787	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide 25 microgramos/125 microgramos/inhalación, suspensión para inhalación en envase a presión	DE/H/5770/002	63.797	GLAXOSMITHKLINE S.A.	ES
Seretide Evohaler 25 mikrogram/125 mikrogram/dos inhalationsspray, suspension	DE/H/5770/002	19166	GLAXOSMITHKLINE AB	SE
Seretide Evohaler 25 microgram /125 microgram per metered dose pressurised inhalation, suspension	DE/H/5770/002	PL 10949/0338	GLAXO WELLCOME UK LIMITED	UK
Seretide forte 25 Mikrogramm/250 Mikrogramm pro	DE/H/5770/003	1-23944	GLAXOSMITHKLINE PHARMA GMBH.	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
Sprühstoß - Druckgasinhalation				
Seretide 25 microgrammes/250 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5770/003	BE220701	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide	DE/H/5770/003	31805	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Evohaler 25 mikrog/250 mikrog/annos inhalaatiosumute, suspensio	DE/H/5770/003	16183	GLAXOSMITHKLINE OY	FI
SERETIDE 250 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé avec valve doseuse	DE/H/5770/003	NL26160	LABORATOIRE GLAXOSMITHKLINE	FR
Seretide Inhaler 25 µ??????aµµ???a /250 µ??????aµµ???a a?? ?a????sµ??? d?s? e?a????µat?? ??a e?sp??? ?p? p?es?.	DE/H/5770/003	2439206	GLAXOSMITHKLINE AEBE	GR
Seretide 25 míkróg/250 míkróg í afmældum skammti af innúðalyfi, dreifu	DE/H/5770/003	IS/1/00/035/03	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide 250 Evohaler 25 microgram/250	DE/H/5770/003	PA 1077/46/6	GLAXOSMITHKLINE (IRELAND) 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
microgram per metered dose pressurised inhalation, suspension.				
Seretide 25 microgrammi/250 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5770/003	034371120	GLAXOSMITHKLINE S.P.A.	IT
Seretide 25 microgrammes/250 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5770/003	260/11/01/0950	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide 25/250 Inhalator CFK-vrij, aërosol, suspensie 25 microgram/250 microgram/dosis.	DE/H/5770/003	RVG 25867	GLAXOSMITHKLINE B.V.	NL
Seretaide Inalador, 25 microgramas/250 microgramas por dose calibrada, suspensão pressurizada para inalação	DE/H/5770/003	3512886	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide 25 microgramos/250 microgramos/inhalación, suspensión para inhalación en envase a presión	DE/H/5770/003	63.798	GLAXOSMITHKLINE S.A.	ES
Seretide Evohaler forte 25 mikrogram/250	DE/H/5770/003	19167	GLAXOSMITHKLINE 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
mikrogram/dos inhalationsspray, suspension				
Seretide Evohaler 25 microgram /250 microgram per metered dose pressurised inhalation, suspension	DE/H/5770/003	PL 10949/0339	GLAXO WELLCOME UK LIMITED	UK
atmadisc mite Dosier-Aerosol 25 µg/50 µg Druckgasinhalation, Suspension	DE/H/5770/001	50678.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
atmadisc Dosier-Aerosol 25 µg/125 µg Druckgasinhalation, Suspension	DE/H/5770/002	50678.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
atmadisc® forte Dosier-Aerosol 25 µg/250 µg Druckgasinhalation, Suspension (-)	DE/H/5770/003	50678.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Comboterol, (25 µg + 125 µg)/dawke inhalacyjna, aerozol inhalacyjny, zawiesina	not available	23600	PRZEDSIĘBIORSTWO FARMACEUTYCZNE LEK-AM SP. Z O.O.	PL
Comboterol, (25 µg + 250 µg)/dawke inhalacyjna, aerozol inhalacyjny, zawiesina	not available	23601	PRZEDSIĘBIORSTWO FARMACEUTYCZNE LEK-AM SP. Z O.O.	PL
AirFluSol Sprayhaler 25 mikrogramm/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	NL/H/3707/001	OGYI-T-22596/08	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Sprayhaler 25	NL/H/3707/001	OGYI-T-22596/07	SANDOZ HUNGÁRIA 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
mikrogramm/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió				
AirFluSol Sprayhaler 25 mikrogramm/250 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	NL/H/3707/002	OGYI-T-22596/11	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Sprayhaler 25 mikrogramm/250 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	NL/H/3707/002	OGYI-T-22596/10	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Sprayhaler 25 mikrogramm/250 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	NL/H/3707/002	OGYI-T-22596/12	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Sprayhaler 25 mikrogramm/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	NL/H/3707/001	OGYI-T-22596/09	SANDOZ HUNGÁRIA KFT	HU
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/011	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/012	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje	NL/H/3707/002	LT/1/17/4070/013	SANDOZ PHARMACEUTICALS D.D.	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
suslėgtoji įkvepiamoji suspensija				
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/014	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/015	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/016	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/017	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/018	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 250 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/019	SANDOZ PHARMACEUTICALS D.D.	LT
AirFluSal 25 / 250 mikrogramų / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/002	LT/1/17/4070/020	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/006	SANDOZ PHARMACEUTICALS D.D.	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
AirFluSal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/001	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/010	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/003	SANDOZ PHARMACEUTICALS D.D.	LT
AirFluSal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/005	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/002	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/008	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/009	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje suslėgtoji įkvepiamoji suspensija	NL/H/3707/001	LT/1/17/4070/004	SANDOZ PHARMACEUTICALS D.D.	LT
Airflusal 25 / 125 mikrogramai / dozėje	NL/H/3707/001	LT/1/17/4070/007	SANDOZ PHARMACEUTICALS D.D.	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
suslėgtoji įkvepiamoji suspensija				
Airflusal Dosieraerosol 25 Mikrogramm/250 Mikrogramm pro Dosis - Druckgasinhalation, Suspension	NL/H/3707/002	137581	SANDOZ GMBH	AT
Airflusan Sprayhaler 25 mikrogramů/125 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	NL/H/3707/001	14/524/16-C	SANDOZ S.R.O.	CZ
Airflusan Sprayhaler 25 mikrogramů/250 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	NL/H/3707/002	14/525/16-C	SANDOZ S.R.O.	CZ
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125	NL/H/3707/001	H/17/02320/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
mikrogramov/vpih inhalacijska suspenzija pod tlakom				
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25	NL/H/3707/002	H/17/02320/019	LEK PHARMACEUTICALS	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
mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom			D.D. LJUBLJANA	
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/001	H/17/02320/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	NL/H/3707/002	H/17/02320/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
AirFluSal Sprayhaler 25 micrograms+125 micrograms/dose pressurised inhalation, suspension	NL/H/3707/001	20170144	SANDOZ PHARMACEUTICALS D.D.	BG
AirFluSal Sprayhaler 25 micrograms+250 micrograms/dose pressurised inhalation, suspension	NL/H/3707/002	20170145	SANDOZ PHARMACEUTICALS D.D.	BG
Airflusal 25 mikrogramov/125 mikrogramov, ihalačná suspenzia v tlakovom obale	NL/H/3707/001	14/0101/17-S	SANDOZ PHARMACEUTICALS D.D.	SK
Airflusal 25 mikrogramov/250 mikrogramov, ihalačná suspenzia v tlakovom	NL/H/3707/002	14/0102/17-S	SANDOZ PHARMACEUTICALS D.D.	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
obale				
Airflusal Sprayhaler 25 mikrog/125 mikrog/annos inhalaatiosumute, suspensio	NL/H/3707/001	34010	SANDOZ A/S	FI
Airflusal Sprayhaler 25 mikrog/250 mikrog/annos inhalaatiosumute, suspensio	NL/H/3707/002	34011	SANDOZ A/S	FI
Airflusal Dosieraerosol 25 Mikrogramm/125 Mikrogramm pro Dosis - Druckgasinhalation, Suspension	NL/H/3707/001	137580	SANDOZ GMBH	AT
AirFluSal 25/125 mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija	NL/H/3707/001	17-0090	SANDOZ PHARMACEUTICALS D.D.	LV
AirFluSal 25/250 mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija	NL/H/3707/002	17-0091	SANDOZ PHARMACEUTICALS D.D.	LV
AirFluSal,(25 mikrogramów + 250 mikrogramów)/ dawkę odmierzoną, aerozol inhalacyjny	NL/H/3707/002	24116	SANDOZ GMBH	PL
AirFluSal 25 micrograme/250 micrograme suspensie de	NL/H/3707/002	10027/2017/04	S.C. SANDOZ S.R.L.	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
inhalat presurizată				
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/05	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/06	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/07	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/08	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/09	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/10	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/01	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/02	S.C. SANDOZ S.R.L.	RO
AirFluSal 25	NL/H/3707/001	10026/2017/03	S.C. SANDOZ S.R.L.	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
microgram/125 microgram suspensie de inhalat presurizată				
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/04	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/05	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/06	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/07	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/08	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/09	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/125 microgram suspensie de inhalat presurizată	NL/H/3707/001	10026/2017/10	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de	NL/H/3707/002	10027/2017/01	S.C. SANDOZ S.R.L.	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
inhalat presurizată				
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/02	S.C. SANDOZ S.R.L.	RO
AirFluSal 25 microgram/250 microgram suspensie de inhalat presurizată	NL/H/3707/002	10027/2017/03	S.C. SANDOZ S.R.L.	RO
Airflusal Sprayhaler 25 microgram/125 microgram per afgemeten dosis aërosol, suspensie	NL/H/3707/001	BE514480	SANDOZ N.V.	BE
Airflusal Sprayhaler 25 microgram/250 microgram per afgemeten dosis aërosol, suspensie	NL/H/3707/002	BE514542	SANDOZ N.V.	BE
Salmeterol/Fluticasonpropionaat Sandoz 25/125 microgram, aerosol, suspensie	NL/H/3707/001	RVG 118829	SANDOZ B.V.	NL
Salmeterol/Fluticasonpropionaat Sandoz 25/250 microgram, aerosol, suspensie	NL/H/3707/002	RVG 118833	SANDOZ B.V.	NL
Airflusal® Dosieraerosol 25 Mikrogramm/125 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	NL/H/3707/001	97091.00.00	HEXAL AG	DE
Airflusal Dosieraerosol 25	NL/H/3707/002	97092.00.00	HEXAL AG	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
Mikrogramm/250 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension				
Airflusal Sprayhaler	NL/H/3707/001	045152028	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152030	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152042	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152055	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152079	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152156	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152168	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152170	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152067	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/001	045152016	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152081	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152093	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152105	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152117	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152129	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152131	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152143	SANDOZ S.P.A.	IT
Airflusal Sprayhaler 25 microgrammi/250 microgrammi/dose sospensione pressurizzata per inalazione	NL/H/3707/002	045152182	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152194	SANDOZ S.P.A.	IT
Airflusal Sprayhaler	NL/H/3707/002	045152206	SANDOZ S.P.A.	IT
AirFluSal, (25 mikrogramów + 125 mikrogramów)/dawkę	NL/H/3707/001	24115	SANDOZ GMBH	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
odmierzoną, aerozol inhalacyjny, zawiesina				
AirFluSal MDI 25 microgram/125 microgram/dose pressurised inhalation, suspension	NL/H/3707/001	PA0711/270/001	ROWEX LTD	IE
AirFluSal MDI 25 microgram/250 microgram/dose pressurised inhalation suspension	NL/H/3707/002	PA0711/270/002	ROWEX LTD	IE
Salmex 50 míkrógrömm/100 míkrógrömm/skammt innöndunarduft, afmældir skammtar	SE/H/1590/001	IS/1/18/001/01	GLENMARK PHARMACEUTICALS NORDIC AB	IS
Salmex 50 míkrógrömm/250 míkrógrömm/skammt innöndunarduft, afmældir skammtar	SE/H/1590/002	IS/1/18/001/02	GLENMARK PHARMACEUTICALS NORDIC AB	IS
Salmex 50 míkrógrömm/500 míkrógrömm/skammt innöndunarduft, afmældir skammtar	SE/H/1590/003	IS/1/18/001/03	GLENMARK PHARMACEUTICALS NORDIC AB	IS
Salmex 50 mikrogram/100 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1590/001	53815	GLENMARK PHARMACEUTICALS NORDIC AB	SE
Salmex, afdelt inhalationspulver	SE/H/1590/003	57099	GLENMARK PHARMACEUTICALS NORDIC	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
			AB	
Salmex 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1590/002	53816	GLENMARK PHARMACEUTICALS NORDIC AB	SE
Salmex 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1590/003	53817	GLENMARK PHARMACEUTICALS NORDIC AB	SE
Salmex, afdelt inhalationspulver	SE/H/1590/002	57098	GLENMARK PHARMACEUTICALS NORDIC AB	DK
Salmex, afdelt inhalationspulver	SE/H/1590/001	57097	GLENMARK PHARMACEUTICALS NORDIC AB	DK
Salmex 50 mikrog / 100 mikrog / annos, inhalaatiojauhe, annosteltu	SE/H/1590/001	33885	GLENMARK PHARMACEUTICALS EUROPE LIMITED	FI
Salmex 50 mikrog / 100 mikrog / dos inhalationspulver, avdelad dos	SE/H/1590/001	33885	GLENMARK PHARMACEUTICALS EUROPE LIMITED	FI
Salmex 50 mikrog / 250 mikrog / dos inhalationspulver, avdelad dos	SE/H/1590/002	33886	GLENMARK PHARMACEUTICALS EUROPE LIMITED	FI
Salmex 50 mikrog / 250 mikrog / annos, inhalaatiojauhe, annosteltu	SE/H/1590/002	33886	GLENMARK PHARMACEUTICALS EUROPE LIMITED	FI
Salmex 50 mikrog / 500	SE/H/1590/003	33887	GLENMARK	FI

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
mikrog / annos, inhalaatiojauhe, annosteltu			PHARMACEUTICALS EUROPE LIMITED	
Salmex 50 mikrog /500 mikrog / dos inhalationspulver, avdelad dos	SE/H/1590/003	33887	GLENMARK PHARMACEUTICALS EUROPE LIMITED	FI
Salmex 50 mikrogram/dose / 100 mikrogram/dose inhalasjonspulver, dosedispensert	SE/H/1590/001	15-10998	GLENMARK PHARMACEUTICALS NORDIC AB	NO
Salmex 50 mikrogram/dose / 500 mikrogram/dose inhalasjonspulver, dosedispensert	SE/H/1590/003	15-11000	GLENMARK PHARMACEUTICALS NORDIC AB	NO
Salmex 50 mikrogram/ dose / 250 mikrogram/dose inhalasjonspulver, dosedispensert	SE/H/1590/002	15-10999	GLENMARK PHARMACEUTICALS NORDIC AB	NO
Asthmex 50 mikrogramov/100 mikrogramov dávkovaný inhalačný prášok	SE/H/1590/001	14/0335/18-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Asthmex 50 mikrogramov/250 mikrogramov dávkovaný inhalačný prášok	SE/H/1590/002	14/0336/18-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Asthmex 50 mikrogramov/500 mikrogramov dávkovaný inhalačný prášok	SE/H/1590/003	14/0337/18-S	GLENMARK PHARMACEUTICALS S.R.O.	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
Asthmex 50 mikrogramů/250 mikrogramů dávkovaný prášek k inhalaci	SE/H/1590/002	14/279/18-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Asthmex 50 mikrogramů/500 mikrogramů dávkovaný prášek k inhalaci	SE/H/1590/003	14/280/18-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Asthmex 50 mikrogramů/100 mikrogramů dávkovaný prášek k inhalaci	SE/H/1590/001	14/278/18-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
PLUSVENT accuhaler 50 microgramos/100 microgramos/inhalación, Polvo para inhalación	not available	62.842	GLAXO S.A.	ES
PLUSVENT accuhaler 50 microgramos/250 microgramos/inhalación, Polvo para inhalación	not available	62.843	GLAXO S.A.	ES
PLUSVENT accuhaler 50 microgramos/500 microgramos/inhalación, Polvo para inhalación	not available	62.844	GLAXO S.A.	ES
Airflusal Forspiro 50 mikrogram/500 mikrogram/dos, inhalationspulver, avdelad dos	SE/H/1321/002	49082	SANDOZ A/S	SE
Airflusal Forspiro, 50 mikrogram/250 mikrogram/dos, inhalationspulver, avdelad dos	SE/H/1321/001	49081	SANDOZ A/S	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
Airflusal Forspiro 50 mikrogram/500 mikrogram/dose inhalasjonspulver, dosedispensert	SE/H/1321/002	13-9438	SANDOZ A/S	NO
Airflusal Forspiro 50 mikrogram/250 mikrogram/dose inhalasjonspulver, dosedispensert	SE/H/1321/001	13-9437	SANDOZ A/S	NO
AirFluSal Forspiro, inhalationspulver, afdelt	SE/H/1321/001	52138	SANDOZ A/S	DK
AirFluSal Forspiro, inhalationspulver, afdelt	SE/H/1321/002	52139	SANDOZ A/S	DK
AirFluSal Forspiro 50 Mikrogramm/250 Mikrogramm – einzeldosiertes Pulver zur Inhalation	SE/H/1408/001	135874	SANDOZ GMBH	AT
AirFluSal Forspiro 50 Mikrogramm/500 Mikrogramm – einzeldosiertes Pulver zur Inhalation	SE/H/1408/002	135875	SANDOZ GMBH	AT
Setirenol	SE/H/1408/001	50647	SANDOZ A/S	SE
SETIRENOL 50 MIKROGRAM/500 MIKROGRAM INHALATIONSPULVER, AVDELAD DOS	SE/H/1408/002	50648	SANDOZ A/S	SE
Airflusal Forspiro Fluticasona + Salmeterol 500 µg/dose + 50 µg/dose Pó para inalação	SE/H/1408/002	5622709	SANDOZ 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
em recipiente unidose				
AirFluSal Forspiro Fluticasona + Salmeterol 250 µg/dose + 50 µg/dose Pó para inalação em recipiente unidose	SE/H/1408/001	5622667	SANDOZ FARMACÊUTICA LDA.	PT
Seretide 25/50 Inhaler Suspenze k inhalaci v tlakovém obalu	not available	14/023/03-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
SERETIDE DISKUS 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	not available	H/00/01405/004	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE DISKUS 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	not available	H/00/01405/008	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE DISKUS 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	not available	H/00/01405/006	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE DISKUS 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	not available	H/00/01405/007	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE DISKUS 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	not available	H/00/01405/005	GLAXOSMITHKLINE D.O.O.	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
Samtoral Diskus standard 50 Mikrogramm/250 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/0170/002	1-22898	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	AT
Samtoral Diskus forte 50 Mikrogramm/500 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/0170/003	1-22897	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	AT
Viani 50 µg/250 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/0170/002	44918.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani forte 50 µg/500 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/0170/003	44918.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani Diskus 50 microgram/250 microgram/dose inhalation powder, pre-dispensed	SE/H/0170/002	14595	GLAXOSMITHKLINE AB	SE
Viani Diskus forte 50 microgram/500 microgram/dose inhalation powder, pre-dispensed	SE/H/0170/003	14596	GLAXOSMITHKLINE AB	SE
Veraspir Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/001	2875185	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/100	SE/H/0170/001	2875284	GLAXOSMITHKLINE - PRODUTOS	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
microgramas/dose pó para inalação, em recipiente unidose			FARMACEUTICOS, LDA	
Veraspir Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/001	4813382	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/001	2875383	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/001	4813481	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/002	2875482	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/002	2875581	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/002	4813580	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50	SE/H/0170/002	2875680	GLAXOSMITHKLINE -	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
microgramas/250 microgramas/dose pó para inalação, em recipiente unidose			PRODUTOS FARMACEUTICOS, LDA	
Veraspir Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/002	4813689	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/003	2875789	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/003	2875888	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/003	4813788	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/003	2875987	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Veraspir Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0170/003	4813887	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, 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
Samtoral Diskus levis 50 Mikrogramm/100 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/0170/001	1-22899	ALLEN PHARMAZEUTIKA GESELLSCHAFT M.B.H.	AT
Viani mite 50 µg/100 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/0170/001	44918.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Viani Diskus mite 50 microgram/100 microgram/dose inhalation powder, pre-dispensed	SE/H/0170/001	14594	GLAXOSMITHKLINE AB	SE
Thoreus Diskus 50/100 mikrogramm/adag adagolt inhalációs por	SE/H/0170/001	OGYI-T-9182/01	GLAXOSMITHKLINE KFT.	HU
Thoreus Diskus 50/500 mikrogramm/adag adagolt inhalációs por	SE/H/0170/003	OGYI-T-9182/03	GLAXOSMITHKLINE KFT.	HU
Thoreus Diskus 50/250 mikrogramm/adag adagolt inhalációs por	SE/H/0170/002	OGYI-T-9182/02	GLAXOSMITHKLINE KFT.	HU
Pavtide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0170/001	9573/2017/01	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0170/001	9573/2017/02	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0170/001	9573/2017/03	GLAXOSMITHKLINE (GSK) S.R.L.	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
Pavtide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0170/001	9573/2017/04	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0170/001	9573/2017/05	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0170/002	9574/2017/01	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0170/002	9574/2017/02	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0170/002	9574/2017/03	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0170/002	9574/2017/04	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0170/002	9574/2017/05	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0170/003	9575/2017/01	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/500	SE/H/0170/003	9575/2017/02	GLAXOSMITHKLINE (GSK) S.R.L.	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
microgram pulbere de inhalat				
Pavtide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0170/003	9575/2017/03	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0170/003	9575/2017/04	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Pavtide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0170/003	9575/2017/05	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Viani Diskus 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/002	H/17/02285/009	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/002	H/17/02285/010	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/002	H/17/02285/008	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/002	H/17/02285/006	GLAXOSMITHKLINE D.O.O.	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
Viani Diskus 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/002	H/17/02285/007	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/001	H/17/02285/005	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/001	H/17/02285/004	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/001	H/17/02285/003	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/001	H/17/02285/001	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/100 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/001	H/17/02285/002	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje,	SE/H/0170/003	H/17/02285/015	GLAXOSMITHKLINE D.O.O.	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
odmerjeni				
Viani Diskus 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/003	H/17/02285/014	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/003	H/17/02285/013	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/003	H/17/02285/011	GLAXOSMITHKLINE D.O.O.	SI
Viani Diskus 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/0170/003	H/17/02285/012	GLAXOSMITHKLINE D.O.O.	SI
PAVTIDE DISKUS 100 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/0170/001	NL46895	LABORATOIRE GLAXOSMITHKLINE	FR
PAVTIDE DISKUS 250 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/0170/002	NL46896	LABORATOIRE GLAXOSMITHKLINE	FR
PAVTIDE DISKUS 500	SE/H/0170/003	NL46897	LABORATOIRE	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
microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose			GLAXOSMITHKLINE	
Aldopulmin 50 microgramos/500 microgramos/inhalación, polvo para inhalación (unidosis)	not available	84562	LABORATORIO ALDO-UNIÓN, S.L.	ES
Seretide 50, (50 ?g + 25 ?g)/dawke inhalacyjna, aerozol inhalacyjny, zawiesina	not available	9069	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Seretide 125, (125 mg + 25 mg)/dawkę inhalacyjną, aerozol inhalacyjny, zawiesina	not available	9070	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Seretide 250, (250 mg + 25 mg)/dawkę inhalacyjną, aerozol inhalacyjny, zawiesina	not available	9071	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Airflusal® Forspiro® 50 microgram/250 microgram/dosis inhalatiepoeder, voorverdeeld	SE/H/1407/001	BE464800	SANDOZ N.V.	BE
Airflusal® Forspiro® 50 microgram/500 microgram/dosis inhalatiepoeder, voorverdeeld	SE/H/1407/002	BE464817	SANDOZ N.V.	BE
Flaxxone Forspiro 50 mikrogram/500	SE/H/1407/002	50637	SANDOZ A/S	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
mikrogram inhalationspulver, avdelad dos				
Flaxxone Forspiro, 50 mikrogram/250 mikrogram inhalationspulver, avdelad dos	SE/H/1407/001	50636	SANDOZ A/S	SE
Zoreeda 25 microgram/125 microgram per actuation pressurised inhalation, suspension.	IE/H/0654/001	PL 36390/0239	CIPLA (EU) LIMITED	UK
Zoreeda 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/0654/002	PL 36390/0240	CIPLA (EU) LIMITED	UK
Zoreeda 25 microgram/125 microgram per actuation pressurised inhalation, suspension	IE/H/0654/001	PA 1963/009/001	CIPLA EUROPE NV	IE
Zoreeda 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/0654/002	PA 1963/009/002	CIPLA EUROPE NV	IE
Sereflo 25 microgram/125 microgram per actuation pressurised inhalation, suspension.	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	IE
Sereflo 25	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	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
microgram/250 microgram per actuation pressurised inhalation, suspension.				
Combisal 25 microgram /50 microgram per metered dose pressurized inhalation, suspension	UK/H/6076/001	PL 36532/0001	GENETIC SPA	UK
Combisal 25microgram /125 microgram per metered dose pressurized inhalation, suspensio	UK/H/6076/002	PL 36532/0002	GENETIC SPA	UK
Combisal 25 microgram /250 microgram per metered dose pressurised inhalation, suspension.	UK/H/6076/003	PL 36532/0003	GENETIC SPA	UK
Airsus 25 microgrammi / 50 microgrammi / per dose predosata sospensione pressurizzata per inalazione	UK/H/6076/001	044250013	GENETIC SPA	IT
Airsus 25 microgrammi / 125 microgrammi / per dose predosata sospensione pressurizzata per inalazione	UK/H/6076/002	044250025	GENETIC SPA	IT
Airsus 25 microgrammi / 250 microgrammi / per dose predosata	UK/H/6076/003	044250037	GENETIC SPA	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
sospensione pressurizzata per inalazione				
Azodal 25 microgram /50 microgram per metered dose pressurised inhalation, suspension	UK/H/6076/001	30277/15/14-05-2018	GENETIC SPA	GR
Azodal 25 microgram /125 microgram per metered dose pressurised inhalation, suspension	UK/H/6076/002	30278/15/14-05-2018	GENETIC SPA	GR
Azodal 25 microgram /250 microgram per metered dose pressurised inhalation, suspension	UK/H/6076/003	30279/15/14-05-2018	GENETIC SPA	GR
Fluticomb, (25 mikrogramów + 50 mikrogramów)/dawkę odmierzoną, aerozol inhalacyjny, zawiesina	UK/H/6076/001	25349	Zakłady Farmaceutyczne POLPHARMA S.A.	PL
Fluticomb, (25 mikrogramów + 125 mikrogramów)/dawkę odmierzoną, aerozol inhalacyjny, zawiesina	UK/H/6076/002	25350	Zakłady Farmaceutyczne POLPHARMA S.A.	PL
Fluticomb, (25 mikrogramów + 250 mikrogramów)/dawkę odmierzoną, aerozol inhalacyjny, zawiesina	UK/H/6076/003	25351	Zakłady Farmaceutyczne POLPHARMA S.A.	PL
Avenor 25 microgram /50 microgram per	UK/H/6816/003/DC	PL 36532/0004 - 0001	GENETIC SPA	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
metered dose pressurized inhalation, suspension				
Avenor 25 microgram /125 microgram per metered dose pressurized inhalation, suspension	UK/H/6816/003/DC	PL 36532/0005 - 0001	GENETIC SPA	UK
Avenor 25 microgram /250 microgram per metered dose pressurized inhalation, suspension	UK/H/6816/003/DC	PL 36532/0006 - 0001	GENETIC SPA	UK
Salmeterol/Fluticasone Genetic 25 μικρογραμμάρια /50 μικρογραμμάρια ανά καθορισμένη δόση εναιώρημα για εισπνοή υπό πίεση.	PT/H/2388/001	41405/ 08-04-2019	GENETIC SPA	GR
Salmeterol/Fluticasone Genetic 25 μικρογραμμάρια /125 μικρογραμμάρια ανά καθορισμένη δόση εναιώρημα για εισπνοή υπό πίεση.	PT/H/2388/002	41406/ 08-04-2019	GENETIC SPA	GR
Salmeterol/Fluticasone Genetic 25 μικρογραμμάρια /250 μικρογραμμάρια ανά καθορισμένη δόση εναιώρημα για εισπνοή υπό πίεση.	PT/H/2388/003	41407/ 08-04-2019	GENETIC SPA	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
PROPIONATE DE FLUTICASONE/SALMETEROL GENETIC 50 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	PT/H/2388/001	34009 301 757 9 8	GENETIC SPA	FR
PROPIONATE DE FLUTICASONE/SALMETEROL GENETIC 125 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	PT/H/2388/002	34009 301 758 0 4	GENETIC SPA	FR
PROPIONATE DE FLUTICASONE/SALMETEROL GENETIC 250 microgrammes/25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	PT/H/2388/003	34009 301 758 1 1	GENETIC SPA	FR
Salmeterol/Flutikazón Genetic 25 mikrogramov/50 mikrogramov/dávka inhalacná suspenzia v tlakovom obale	PT/H/2388/001-003/DC	14/0084/19-S	GENETIC SPA	SK
Salmeterol/Flutikazón Genetic 25 mikrogramov/125mikrogramov/dávka inhalacná suspenzia v tlakovom	PT/H/2388/001-003/DC	14/0085/19-S	GENETIC SPA	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
obale				
Salmeterol/Flutikazón Genetic 25 mikrogramov/250 mikrogramov/dávka inhalacná suspenzia v tlakovom obale	PT/H/2388/001-003/DC	14/0086/19-S	GENETIC SPA	SK
Salmeterol/Fluticasone Genetic 25 mikrogramů/50 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	PT/H/2388/001-003/DC	14/494/17-C	GENETIC SPA	CZ
Salmeterol/Fluticasone Genetic 25 mikrogramu/125 mikrogramu/dávka suspenze k inhalaci v tlakovém obalu	PT/H/2388/001-003/DC	14/495/17-C	GENETIC SPA	CZ
Salmeterol/Fluticasone Genetic 25 mikrogramu/250 mikrogramu/dávka suspenze k inhalaci v tlakovém obalu	PT/H/2388/001-003/DC	14/496/17-C	GENETIC SPA	CZ
Salmeterol/flutikazon Genetic 25 mikrogramov/50 mikrogramov/vpih inhalacijska suspenzija pod tlakom	PT/H/2388/001	H/19/02587/001	GENETIC SPA	SI
Salmeterol/flutikazon Genetic 25 mikrogramov/125	PT/H/2388/002	H/19/02587/002	GENETIC SPA	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
mikrogramov/vpih inhalacijska suspenzija pod tlakom				
Salmeterol/flutikazon Genetic 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	PT/H/2388/003	H/19/02587/003	GENETIC SPA	SI
Салметерол/Флутиказон Генетик 25 микрограма/50 микрограма в една отмерена доза суспензия под налягане за инхалация.	PT/H/2388/001	BG/MA/MP-46439/08-07-2019	GENETIC SPA	BG
Салметерол/Флутиказон Генетик 25 микрограма/125 микрограма в една отмерена доза суспензия под налягане за инхалация.	PT/H/2388/002	BG/MA/MP-46440/08-07-2019	GENETIC SPA	BG
Салметерол/Флутиказон Генетик 25 микрограма/250 микрограма в една отмерена доза суспензия под налягане за инхалация.	PT/H/2388/003	BG/MA/MP-46441/08-07-2019	GENETIC SPA	BG
Salmeterolo e Fluticasone Genetic 25 microgrammi/50 microgrammi/dose	PT/H/2388/001	046084012	GENETIC SPA	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
predosata, sospensione pressurizzata per inalazione				
Salmeterolo e Fluticasone Genetic 25 microgrammi/125 microgrammi/dose predosata, sospensione pressurizzata per inalazione	PT/H/2388/002	046084024	GENETIC SPA	IT
Salmeterolo e Fluticasone Genetic 25 microgrammi/250 microgrammi/dose predosata, sospensione pressurizzata per inalazione	PT/H/2388/003	046084036	GENETIC SPA	IT
Inaladuo 25 microgramos/50 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	64.064	SMITHKLINE BEECHAM FARMA, S.A.	ES
Inaladuo 25 microgramos/125 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	64.065	GLAXOSMITHKLINE, S.A.	ES
Inaladuo 25 microgramos/250 microgramos/inhalación, suspensión para inhalación en envase a	not available	64.066	SMITHKLINE BEECHAM FARMA, S.A.	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
presión				
Salflutin 50 Mikrogramm/100 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/001	97198.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Salflutin 50 Mikrogramm/250 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/002	97199.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Salflutin 50 Mikrogramm/500 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/003	97200.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Salmeterol/Fluticason Glenmark 50 Mikrogramm/500 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/003	138371	GLENMARK PHARMACEUTICALS NORDIC AB	AT
Salmeterol/Fluticason Glenmark 50 Mikrogramm/250 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/002	138370	GLENMARK PHARMACEUTICALS NORDIC AB	AT
Salmeterol/Fluticason Glenmark 50 Mikrogramm/100 Mikrogramm einzeldosiertes Pulver zur Inhalation	AT/H/0669/001	138369	GLENMARK ARZNEIMITTEL GMBH	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
Salmeterol/Fluticasone 1A Farma 50 mikrogram/500 mikrogram/dos, Inhalationspulver, avdelad dos	SE/H/1663/002	55540	1A FARMA A/S	SE
AirFluSal Forspiro, (50 mikrogramów + 250 mikrogramów)/dawkę inhalacyjną, proszek do inhalacji, podzielony	SE/H/1448/001	23169	SANDOZ GMBH	PL
AirFluSal Forspiro, (50 mikrogramów + 500 mikrogramów)/dawkę inhalacyjną, proszek do inhalacji, podzielony	SE/H/1448/002	23170	SANDOZ GMBH	PL
Roadasan Forspiro, 50 mikrogram/250mikrogram/dos inhalationspulver, avdelad dos	SE/H/1448/001	51642	SANDOZ A/S	SE
Roadasan Forspiro, 50 mikrogram/500mikrogram/dos inhalationspulver, avdelad dos	SE/H/1448/002	51643	SANDOZ A/S	SE
Sereflo 25 microgram/125 microgram per actuation pressurised inhalation,suspension	IE/H/0653/001	PL 36390/0237	CIPLA (EU) LIMITED	UK
Sereflo 25 microgram/250 microgram per actuation pressurised inhalation,suspension	IE/H/0653/002	PL 36390/0238	CIPLA (EU) LIMITED	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
Inhatec 50 microgram/500 microgram/ dose inhalation powder, pre-dispensed.	SE/H/1856/001	PL 49518/0003	WELLNEX GMBH	UK
Salmeterol/Fluticasone Wellnex 50 mikrog/500 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/1856/001	36309	WELLNEX GMBH	FI
Flusarion Easyhaler 50 mikrogram/250 mikrogram/dos inhalationspulver	SE/H/1694/001	56281	ORION CORPORATION	SE
Flusarion Easyhaler 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1694/002	56282	ORION CORPORATION	SE
Flusarion Easyhaler 50 Mikrogramm/500 Mikrogramm pro Inhalation, Pulver zur Inhalation	SE/H/1694/002	99809.00.00	ORION CORPORATION	DE
Flusarion Easyhaler 50 Mikrogramm/250 Mikrogramm pro Inhalation, Pulver zur Inhalation	SE/H/1694/001	99808.00.00	ORION CORPORATION	DE
PLUSVENT 25 microgramos/50 microgramos/ inhalación, Suspensión para inhalación en envase a presión	not available	63.869	GLAXO S.A.	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
PLUSVENT 25 microgramos/125 microgramos/ inhalación, Suspensión para inhalación en envase a presión	not available	63.870	GLAXO S.A.	ES
PLUSVENT 25 microgramos/250 microgramos/ inhalación, Suspensión para inhalación en envase a presión	not available	63.871	GLAXO S.A.	ES
Salfuler Easyhaler 50/500 microgram/dosis, inhalatiepoeder	SE/H/1700/02/DC	RVG 121013	ORION CORPORATION	NL
Salfuler Easyhaler, 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1700/02/DC	56294	ORION CORPORATION	SE
Brisomax Inalador 50 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703782	BIAL - PORTELA & Cª , SA	PT
Brisomax Inalador 125 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703881	BIAL - PORTELA & Cª , SA	PT
Brisomax Inalador 250 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703980	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
Duohal 25 micrograms/125 micrograms /dose pressurised inhalation, suspension	SE/H/1599/001	53791	CIPLA EUROPE NV	SE
Duohal 25 micrograms/250 micrograms /dose pressurised inhalation, suspension	SE/H/1599/002	53792	CIPLA EUROPE NV	SE
Салметерол и флутиказон Сипла 25 микрограма/ 125 микрограма на впръскване, суспензия под налягане за инхалация	SE/H/1599/001	20170119	CIPLA EUROPE NV	BG
Салметерол и флутиказон Сипла 25 микрограма/ 250 микрограма на впръскване, суспензия под налягане за инхалация	SE/H/1599/002	20170120	CIPLA EUROPE NV	BG
PROPIONATE DE FLUTICASONE/SALMETEROL CIPLA 125 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 301 564 3 8	CIPLA EUROPE NV	FR
PROPIONATE DE FLUTICASONE/SALMETE	SE/H/1599/001	34009 301 564 5 2	CIPLA EUROPE NV	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
ROL CIPLA 250 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé				
PROPIONATE DE FLUTICASONE/SALMETE ROL CIPLA 125 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 550 581 8 2	CIPLA EUROPE NV	FR
PROPIONATE DE FLUTICASONE/SALMETE ROL CIPLA 125 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 550 581 9 9	CIPLA EUROPE NV	FR
PROPIONATE DE FLUTICASONE/SALMETE ROL CIPLA 125 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 550 582 0 5	CIPLA EUROPE NV	FR
PROPIONATE DE FLUTICASONE/SALMETE ROL CIPLA 250 microgrammes/25 microgrammes/ dose,	SE/H/1599/001	34009 550 582 1 2	CIPLA EUROPE NV	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
suspension pour inhalation en flacon pressurisé				
PROPIONATE DE FLUTICASONE/SALMETEROL CIPLA 250 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 550 582 3 6	CIPLA EUROPE NV	FR
PROPIONATE DE FLUTICASONE/SALMETEROL CIPLA 250 microgrammes/25 microgrammes/ dose, suspension pour inhalation en flacon pressurisé	SE/H/1599/001	34009 550 582 4 3	CIPLA EUROPE NV	FR
Fluterolo Easyhaler 50 mikrogram/250 mikrogram/dos inhalationspulver	SE/H/1693/01/DC	56279	ORION CORPORATION	SE
Fluterolo Easyhaler 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1693/02/DC	56280	ORION CORPORATION	SE
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961012	ORION CORPORATION	IT
Safubref 50 microgrammi/500 microgrammi/dose,	SE/H/1693/002	045961048	ORION CORPORATION	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
polvere per inalazione				
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961024	ORION CORPORATION	IT
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961036	ORION CORPORATION	IT
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961075	ORION CORPORATION	IT
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961087	ORION CORPORATION	IT
Safubref 50 microgrammi/250 microgrammi/dose, polvere per inalazione	SE/H/1693/001	045961099	ORION CORPORATION	IT
Safubref 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1693/002	045961051	ORION CORPORATION	IT
Safubref 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1693/002	045961063	ORION CORPORATION	IT
Safubref 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1693/002	045961101	ORION CORPORATION	IT
Safubref 50	SE/H/1693/002	045961113	ORION CORPORATION	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
microgrammi/500 microgrammi/dose, polvere per inalazione				
Safubref 50 microgrammi/500 microgrammi/dose, polvere per inalazione	SE/H/1693/002	045961125	ORION CORPORATION	IT
Salmex 50 microgrammes / 500 microgrammes / dose, poudre pour inhalation en recipient unidose	UK/H/6498/001/DC	2019070196	APC INSTYTUT SP. Z O.O.	LU
Seretide Inhaler CFC-Free 25 microgram/250 microgram suspensie de inhalat presurizată	not available	887/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Inhaler CFC-Free 25 microgram/125 microgram suspensie de inhalat presurizata	not available	886/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Inhaler CFC-Free 25 microgram/50 microgram suspensie de inhalat presurizata	not available	885/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Dysk 250, (250 mg + 50 mg)/dawkę inhalacyjną, proszek do inhalacji	not available	8325	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Seretide Dysk 100, (100 mg + 50 mg)/dawkę inhalacyjną, proszek do inhalacji	not available	8324	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Seretide Dysk 500, (500 mg + 50 mg)/dawkę	not available	8326	GLAXOSMITHKLINE (IRELAND) LIMITED	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
inhalacyjną, proszek do inhalacji				
Sirdupla 25 microgram/125 microgram per metered dose pressurised inhalation, suspension.	DE/H/5724/001	PL 04569/1449	GENERICS [UK] LIMITED	UK
Sirdupla 25 microgram/250 microgram per metered dose pressurised inhalation, suspension.	DE/H/5724/002	PL 04569/1450	GENERICS [UK] LIMITED	UK
SERKEP 25 mikrogram/125 mikrogram per dose inhalasjonsaerosol, suspensjon	DE/H/5724/001	16-11186	MYLAN AB	NO
SERKEP 25 mikrogram/250 mikrogram per dose inhalasjonsaerosol, suspensjon.	DE/H/5724/002	16-11187	MYLAN AB	NO
Serkep 25 mikrogram/125 mikrogram/dos inhalationsspray, suspension	DE/H/5724/001	54831	MYLAN AB	SE
Serkep 25 mikrogram/250 mikrogram/dos inhalationsspray, suspension	DE/H/5724/002	54832	MYLAN 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
Serkep 25 míkróg/125 míkróg í afmældum skammti af innúðalyfi, dreifu.	DE/H/5724/001	IS/1/16/106/01	MYLAN AB	IS
Serkep 25 míkróg/250 míkróg í afmældum skammti af innúðalyfi, dreifu.	DE/H/5724/002	IS/1/16/106/02	MYLAN AB	IS
Serkep, 25 mikrogrammi/125 mikrogrammi mõõdetud annuses, inhalatsiooniaerosool, suspensioon	DE/H/5724/001	925216	MYLAN S.A.S	EE
Serkep, 25 mikrogrammi/250 mikrogrammi mõõdetud annuses, inhalatsiooniaerosool, suspensioon	DE/H/5724/002	925316	MYLAN S.A.S	EE
Serkep 25/125 mikrogramai/dozėje suslėgtoji įkvepiamoji suspensija	DE/H/5724/001	LT/1/16/4018/001	MYLAN S.A.S	LT
Serkep 25/250 mikrogramų/dozėje suslėgtoji įkvepiamoji suspensija	DE/H/5724/002	LT/1/16/4018/002	MYLAN S.A.S	LT
Serkep 25/125 mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija	DE/H/5724/001	16-0255	MYLAN S.A.S	LV
Serkep 25/250	DE/H/5724/002	16-0256	MYLAN S.A.S	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
mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija				
Seritomyl 25 Mikrogramm/125 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5724/001	BE503831	MYLAN BVBA/SPRL	BE
Seritomyl 25 microgrammes / 125 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5724/001	BE503831	MYLAN BVBA/SPRL	BE
Seritomyl, 25 microgram/125 microgram per afgemeten dosis, aërosol, suspensie	DE/H/5724/001	BE503831	MYLAN BVBA/SPRL	BE
Seritomyl 25 Mikrogramm/250 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5724/002	BE503840	MYLAN BVBA/SPRL	BE
Seritomyl 25 microgrammes / 250 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5724/002	BE503840	MYLAN BVBA/SPRL	BE
Seritomyl, 25	DE/H/5724/002	BE503840	MYLAN BVBA/SPRL	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
microgram/250 microgram per afgemeten dosis, aërosol suspensie				
Sirdupla 25 mikrograma/125 mikrograma po potisku, stlačeni inhalat, suspenzija	DE/H/5724/001	HR-H-254713416	MYLAN HRVATSKA D.O.O.	HR
Sirdupla 25 mikrograma/250 mikrograma po potisku, stlačeni inhalat, suspenzija	DE/H/5724/002	HR-H-709386868	MYLAN HRVATSKA D.O.O.	HR
Serkep 25 microgramas + 125 microgramas por dose calibrada, suspensão pressurizada para inalação	DE/H/5724/001	5699822	MYLAN, LDA	PT
Serkep 25 microgramas + 250 microgramas por dose calibrada, suspensão pressurizada para inalação	DE/H/5724/002	5699830	MYLAN, LDA	PT
Serkep 25 Mikrogramm/125 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	DE/H/5724/001	137380	ARCANA ARZNEIMITTEL GMBH	AT
Serkep 25 Mikrogramm/250 Mikrogramm pro Sprühstoß	DE/H/5724/002	137381	ARCANA ARZNEIMITTEL GMBH	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
Druckgasinhalation, Suspension				
Sirdupla 25 mikrogramů/125 mikrogramů/dávka, suspenze k inhalaci v tlakovém obalu	DE/H/5724/001	14/679/16-C	GENERICS [UK] LIMITED	CZ
Sirdupla 25 mikrogramů/250 mikrogramů/dávka, suspenze k inhalaci v tlakovém obalu	DE/H/5724/002	14/680/16-C	GENERICS [UK] LIMITED	CZ
Serzyl 25 microgrammi/125 microgrammi per dose predosata, sospensione pressurizzata per inalazione	DE/H/5724/001	045124017	MYLAN S.P.A.	IT
Serzyl 25 microgrammi/250 microgrammi per dose predosata, sospensione pressurizzata per inalazione	DE/H/5724/002	045124029	MYLAN S.P.A.	IT
Seritomyl 25 microgrammes / 250 microgrammes par dose mesurée, suspension pour inhalation en flacon pressurisé	DE/H/5724/002	2017070248	MYLAN BVBA/SPRL	LU
Seritomyl 25 microgrammes / 125 microgrammes par dose mesurée, suspension	DE/H/5724/001	2017070247	MYLAN BVBA/SPRL	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
pour inhalation en flacon pressurisé				
Serkep 25 μικρογραμμάρια /125 μικρογραμμάρια ανά δοσιμετρημένη δόση εναιώρημα για εισπνοή υπό πίεση	DE/H/5724/001	59075/17/29-01-2018	MYLAN S.A.S	GR
Serkep 25 μικρογραμμάρια /250 μικρογραμμάρια ανά δοσιμετρημένη δόση εναιώρημα για εισπνοή υπό πίεση	DE/H/5724/002	49517/17/29-01-2018	MYLAN S.A.S	GR
Serkep, inhalationsspray, suspension	DE/H/5724/001	57839	MYLAN AB	DK
Serkep, inhalationsspray, suspension	DE/H/5724/001	57840	MYLAN AB	DK
Serkep 25 mikrog/125 mikrog/annos inhalaatiosumute, suspensio	DE/H/5724/001	34226	MYLAN AB	FI
Serkep 25 mikrogram/125 mikrogram/dos inhalationsspray, suspension	DE/H/5724/001	34226	MYLAN AB	FI
Serkep 25 mikrog/250 mikrog/annos inhalaatiosumute, suspensio	DE/H/5724/002	34227	MYLAN AB	FI
Serkep 25 mikrogram/250 mikrogram/dos	DE/H/5724/002	34227	MYLAN AB	FI

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
inhalationsspray, suspension				
Serkep 25 mikrogramov/125 mikrogramov inhalačná suspenzia v tlakovom obale	DE/H/5724/001	14/0492/16-S	MYLAN IRELAND LIMITED	SK
Serkep 25 mikrogramov/250 mikrogramov inhalačná suspenzia v tlakovom obale	DE/H/5724/002	14/0493/16-S	MYLAN IRELAND LIMITED	SK
Serhalyx 25/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	DE/H/5724/001	OGYI-T-23117/01	MYLAN IRELAND LIMITED	HU
Serhalyx 25/250 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	DE/H/5724/002	OGYI-T-23117/02	MYLAN IRELAND LIMITED	HU
Serkep 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	DE/H/5724/001	H/17/02354/001	MYLAN IRELAND LIMITED	SI
Serkep 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	DE/H/5724/002	H/17/02354/002	MYLAN IRELAND LIMITED	SI
Serkep 25 mikrogramů/125 mikrogramů/dávka suspenze k inhalaci v	DE/H/5724/001	14/679/16-C	MYLAN IRELAND LIMITED	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
tlakovém obalu				
Serkep 25 mikrogramů/250 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	DE/H/5724/002	14/680/16-C	MYLAN IRELAND LIMITED	CZ
Sirdupla 25 microgram/250 microgram per metered dose pressurised inhalation, suspension	DE/H/5724/002	PA 577/201/002	MCDERMOTT LABORATORIES LTD	IE
Sirdupla 25 microgram/125 microgram per metered dose pressurised inhalation, suspension	DE/H/5724/001	PA 0577/201/001	MCDERMOTT LABORATORIES LTD	IE
Sirdupla 25 microgram/125 microgram per metered dose pressurised inhalation, suspension	DE/H/5724/001	MA1270/02201	MYLAN IRELAND LIMITED	MT
Sirdupla 25 microgram/250 microgram per metered dose pressurised inhalation, suspension.	DE/H/5724/002	MA1270/02202	MYLAN IRELAND LIMITED	MT
Seritomyl 25 Mikrogramm/125 Mikrogramm pro abgemessene Dosis Druckgasinhalation,	DE/H/5724/001	2017070247	MYLAN BVBA/SPRL	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
Suspension				
Seritomyl 25 Mikrogramm/250 Mikrogramm pro abgemessene Dosis Druckgasinhalation, Suspension	DE/H/5724/002	2017070248	MYLAN BVBA/SPRL	LU
Serkep 25 Mikrogramm/125 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	DE/H/5724/001	91143.00.00	MYLAN GERMANY GMBH	DE
Serkep 25 Mikrogramm/250 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	DE/H/5724/002	91144.00.00	MYLAN GERMANY GMBH	DE
ROLENIUM 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0972/001	041500012/M	ELPEN PHARMACEUTICAL CO. INC.	IT
ROLENIUM 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0972/002	041500024/M	ELPEN PHARMACEUTICAL CO. INC.	IT
Relanio 50 míkróg/250 míkróg/skammt innöndunarduft, afmældir skammtar.	SE/H/0972/001	IS/1/13/094/01	ELPEN PHARMACEUTICAL CO. INC.	IS
Relanio 50 míkróg/500	SE/H/0972/002	IS/1/13/094/02	ELPEN PHARMACEUTICAL	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
míkróg/skammt innöndunarduft, afmældir skammtar.			CO. INC.	
Rolenium 50 Mikrogramm/250 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/0972/001	81933.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Rolenium 50 Mikrogramm /500 Mikrogramm /Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/0972/002	81934.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Relanio 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/0972/001/DC	44235	ELPEN PHARMACEUTICAL CO. INC.	SE
Relanio 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/0972/002	44236	ELPEN PHARMACEUTICAL CO. INC.	SE
DIMENIUM 50 mikrogramu/250 mikrogramu dávkovaný prášek k inhalaci	SE/H/0972/001	14/581/11-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
DIMENIUM 50mikrogramů/500 mikrogramů, dávkovaný prášek k inhalaci	SE/H/0972/002	14/582/11-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
DIMENIUM 50 mikrogramov/250 mikrogramov/dávka	SE/H/0972/001	14/0405/11-S	ELPEN PHARMACEUTICAL CO. INC.	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
dávkovaný inhalačný prášok				
DIMENIUM 50 mikrogramov/500 mikrogramov/dávka dávkovaný inhalačný prášok	SE/H/0972/002	14/0406/11-S	ELPEN PHARMACEUTICAL CO. INC.	SK
DIMENIUM 250 µg/dose + 50 µg/dose, pó para inalação em recipiente unidose	SE/H/0972/001	5400676	ELPEN PHARMACEUTICAL CO. INC.	PT
DIMENIUM 500 µg/dose + 50 µg/dose, pó para inalação em recipiente unidose	SE/H/0972/002	5400700	ELPEN PHARMACEUTICAL CO. INC.	PT
Salmeterol/Fluticasone Hexal, 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1322/002	49084	HEXAL AG	SE
Lastenol Forspiro, 50 mikrogram/500 mikrogram inhalationspulver, avdelad dos	SE/H/1404/002	50635	SANDOZ A/S	SE
AIRFLUSAL FORSPIRO	SE/H/1404/002	043262029	SANDOZ S.P.A.	IT
Salmeson 50 mikrog/250 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/1191/002	30089	ELPEN PHARMACEUTICAL CO. INC.	FI
Salmeson 50 mikrog/500 mikrog/annos inhalaatiojauhe,	SE/H/1191/003	30090	ELPEN PHARMACEUTICAL CO. INC.	FI

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
annosteltu				
SALMESON 250 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/002	34009 277 074 2 1	ELPEN PHARMACEUTICAL CO. INC.	FR
SALMESON 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/003	34009 277 077 1 1	ELPEN PHARMACEUTICAL CO. INC.	FR
Symflusal, (50 mikrogramów + 250 mikrogramów)/dawke, proszek do inhalacji, podzielony	SE/H/1191/002	21380	SYMPHAR SP. Z O.O.	PL
Symflusal, (50 mikrogramów + 500 mikrogramów)/dawke, proszek do inhalacji, podzielony	SE/H/1191/003	21381	SYMPHAR SP. Z O.O.	PL
SALMESON 250 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/002	34009 300 095 8 1	ELPEN PHARMACEUTICAL CO. INC.	FR
SALMESON 250 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/002	34009 277 075 9 9	ELPEN PHARMACEUTICAL CO. INC.	FR
SALMESON 250 microgrammes/50 microgrammes/dose,	SE/H/1191/002	34009 550 064 4 2	ELPEN PHARMACEUTICAL CO. INC.	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
poudre pour inhalation en récipient unidose				
SALMESON 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/003	34009 300 095 9 8	ELPEN PHARMACEUTICAL CO. INC.	FR
SALMESON 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/003	34009 277 078 8 9	ELPEN PHARMACEUTICAL CO. INC.	FR
SALMESON 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/1191/003	34009 550 064 5 9	ELPEN PHARMACEUTICAL CO. INC.	FR
Salmeson 50 mikrog/dose + 250 mikrog/dose inhalasjonspulver, dosedispensert	SE/H/1191/002/DC	11-8552	ELPEN PHARMACEUTICAL CO. INC.	NO
Salmeson 50 mikrog/dose + 500 mikrog/dose inhalasjonspulver, dosedispensert	SE/H/1191/003/DC	11-8553	ELPEN PHARMACEUTICAL CO. INC.	NO
Salmeson 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1191/002	46640	ELPEN PHARMACEUTICAL CO. INC.	SE
Salmeson 50 mikrogram/500	SE/H/1191/003	46641	ELPEN PHARMACEUTICAL CO. INC.	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
mikrogram/dos inhalationspulver, avdelad dos				
SERETIDE 25 mikrogramov/125 mikrogramov/vpih inhalacijska suspenzija pod tlakom	not available	H/00/01405/002	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE 25 mikrogramov/250 mikrogramov/vpih inhalacijska suspenzija pod tlakom	not available	H/00/01405/003	GLAXOSMITHKLINE D.O.O.	SI
SERETIDE 25 mikrogramov/50 mikrogramov/vpih inhalacijska suspenzija pod tlakom	not available	H/00/01405/001	GLAXOSMITHKLINE D.O.O.	SI
Seretide Inhaler 25 mikrograma + 50 mikrograma u jednoj dozi stlaceni inhalat, suspenzija	not available	HR-H-513256189	GLAXOSMITHKLINE D.O.O.	HR
Seretide Inhaler 25 mikrograma + 250 mikrograma u jednoj dozi stlačeni inhalat, suspenzija	not available	HR-H-572113579	GLAXOSMITHKLINE D.O.O.	HR
Серетид Дискус 50 микрограма/100 микрограма/доза прах за инхалация, предварително дозиран	not available	20000234	GLAXOSMITHKLINE EOOD	BG
Серетид Дискус 50	not available	20000232	GLAXOSMITHKLINE EOOD	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
микрограма/250 микрограма/доза прах за инхалация, предварително дозиран				
Серетид Дискус 50 микрограма/500 микрограма/доза прах за инхалация, предварително дозиран	not available	20000233	GLAXOSMITHKLINE EOOD	BG
Fulsalmide Easyhaler 50 microgram/250 microgram/dose, inhalation powder	SE/H/1697/001	56287	ORION CORPORATION	SE
Fulsalmide Easyhaler 50 microgram/500 microgram/dose, inhalation powder	SE/H/1697/02/DC	56288	ORION CORPORATION	SE
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 250 microgrammes/50 microgrammes/ dose, poudre pour inhalation	SE/H/1697/001	34009 301 757 0 5	ORION CORPORATION	FR
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 250 microgrammes/50 microgrammes/ dose, poudre pour inhalation	SE/H/1697/001	34009 550 635 3 7	ORION CORPORATION	FR
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 250	SE/H/1697/001	34009 550 635 4 4	ORION CORPORATION	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
microgrammes/50 microgrammes/ dose, poudre pour inhalation				
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 500 microgrammes/50 microgrammes/ dose, poudre pour inhalation	SE/H/1697/002	34009 301 757 1 2	ORION CORPORATION	FR
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 500 microgrammes/50 microgrammes/ dose, poudre pour inhalation	SE/H/1697/002	34009 550 635 5 1	ORION CORPORATION	FR
PROPIONATE DE FLUTICASONE/SALMETEROL ORION 500 microgrammes/50 microgrammes/ dose, poudre pour inhalation	SE/H/1697/002	34009 550 635 6 8	ORION CORPORATION	FR
Saflutin 50/100 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/15/3820/001	UAB INTELI GENERICS NORD	LT
Saflutin 50/250 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/15/3820/002	UAB INTELI GENERICS NORD	LT
Saflutin 50/500	not available	LT/1/15/3820/003	UAB INTELI GENERICS	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
mikrogramų/dozėje dozuoti įkvepiamieji milteliai			NORD	
Seretide Inhaler, 25/50 mikrogrammi annuses inhalatsiooniaerosool, suspensioon	not available	406403	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Seretide Inhaler, 25/250 mikrogrammi annuses inhalatsiooniaerosool, suspensioon	not available	406303	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Seretide Inhaler, 25/125 mikrogrammi annuses inhalatsiooniaerosool, suspensioon	not available	406203	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Salmeterol/Flutikason Devatis 25 microgram/250 microgram/actuation, pressurised inhalation, suspension	SE/H/1683/003	57114	DEVATIS GMBH	SE
Salmeterol/Flutikason Devatis 25 microgram/250 microgram/actuation, pressurised inhalation, suspension	SE/H/1683/003	57114	DEVATIS GMBH	SE
Salmeterol/Flutikason Devatis 25 microgram/250 microgram/actuation, pressurised inhalation, suspension	SE/H/1683/003	57114	DEVATIS GMBH	SE
Salmeterol/Flutikason	SE/H/1683/003	57114	DEVATIS GMBH	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
Devatis 25 microgram/250 microgram/actuation, pressurised inhalation, suspension				
Salmeterol/Fluticasonpropionaat 25 microgram/125 microgram/dosis Vincion, aërosol, suspensie	not available	RVG 115995	VINCION BV	NL
Salmeterol/Fluticasonpropionaat 25 microgram/250 microgram/dosis Vincion, aërosol, suspensie	not available	RVG 115996	VINCION BV	NL
Sereflo 25 microgram/125 microgram per actuation pressurised inhalation, suspension.	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	IE
Sereflo 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	IE
Airflusal® Forspiro® 50 microgramos/250 microgramos/inhalación, polvo para inhalación (unidosis)	SE/H/1446/001	80248	SANDOZ FARMACÉUTICA, S.A.	ES
Airflusal® Forspiro® 50 microgramos/500 microgramos/inhalación, polvo para inhalación	SE/H/1446/002	80249	SANDOZ FARMACÉUTICA, S.A.	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
(unidosis)				
Mesaflu Forspiro	SE/H/1446/002	51635	SANDOZ A/S	SE
Mesaflu Forspiro	SE/H/1446/001	51634	SANDOZ A/S	SE
Aloflute 25 microgram/125 microgram per metered dose pressurised inhalation, suspension	UK/H/5068/001	PL 00068/0405	3M HEALTH CARE LIMITED	UK
Aloflute 25 microgram/250 microgram per metered dose pressurised inhalation, suspension	UK/H/5068/002	PL 00068/0406	3M HEALTH CARE LIMITED	UK
Aloflute 25 Mikrogramm/250 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	UK/H/5068/002	91263.00.00	JENSONR+ (IRELAND) LTD	DE
Aloflute 25 Mikrogramm/125 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	UK/H/5068/001	91262.00.00	JENSONR+ (IRELAND) LTD	DE
Maizar Diskus 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	not available	3600285	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Maizar Diskus 50 microgramas/250 microgramas/dose pó para inalação, em	not available	3600582	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, 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
recipiente unidose				
Maizar Diskus 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	not available	3600889	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
AirFluSal MDI 25 microgram/125 microgram per actuation pressurised inhalation, suspension	NL/H/3708/001	PL 04416/1475	SANDOZ LTD	UK
AirFluSal® MDI 25 microgram/250 microgram per actuation pressurised inhalation, suspension	NL/H/3708/002	PL 04416/1476	SANDOZ LTD	UK
Salmeterol/Fluticasonpropionaat Sandoz 25/125 microgram, aerosol, suspensie	NL/H/3708/001	RVG 118834	SANDOZ B.V.	NL
Salmeterol/Fluticasonpropionaat Sandoz 25/250 microgram, aerosol, suspensie	NL/H/3708/002	RVG 118835	SANDOZ B.V.	NL
Flusamer 50 mikrogrammi/250 mikrogrammi/annuses annustatud inhalatsioonipulber	EE/H/0224/001	779312	ELPEN PHARMACEUTICAL CO. INC.	EE
Flusamer 50 mikrogrammi/500 mikrogrammi/annuses annustatud inhalatsioonipulber	EE/H/0224/002	779212	ELPEN PHARMACEUTICAL CO. INC.	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
Флусамер 50 микрограма /250 микрограма/доза прах за инхалация, предварително дозиран.	EE/H/0224/001	20120291	ELPEN PHARMACEUTICAL CO. INC.	BG
Флусамер 50 микрограма /500 микрограма/доза прах за инхалация, предварително дозиран.	EE/H/0224/002	20120292	ELPEN PHARMACEUTICAL CO. INC.	BG
Flamerio 50 mikrogramų /250 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	EE/H/0224/001	LT/1/12/2888/001	ELPEN PHARMACEUTICAL CO. INC.	LT
Flamerio 50 mikrogramų /500 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	EE/H/0224/002	LT/1/12/2888/002	ELPEN PHARMACEUTICAL CO. INC.	LT
Flamerio 50 mikrogrami/250 mikrogrami/devā inhalācijas pulveris, dalīts	EE/H/0224/001	12-0049	ELPEN PHARMACEUTICAL CO. INC.	LV
Flamerio 50 mikrogrami/500 mikrogrami/devā inhalācijas pulveris, dalīts	EE/H/0224/002	12-0050	ELPEN PHARMACEUTICAL CO. INC.	LV
Flamerio 50 mikrogramov/250 mikrogramov na odmerek, odmerjeni	EE/H/0224/001/DC	H/12/00614/001	ELPEN PHARMACEUTICAL CO. INC.	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
prašek za inhaliranje				
Flamerio 50 mikrogramov/500 mikrogramov na odmerek, odmerjeni prasek za inhaliranje	EE/H/0224/002/DC	H/12/00614/002	ELPEN PHARMACEUTICAL CO. INC.	SI
Flusamer 50 Mikrogramm/250 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	EE/H/0224/001/DC	2012090037	ELPEN PHARMACEUTICAL CO. INC.	LU
Flusamer 50 Mikrogramm/500 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	EE/H/0224/002/DC	2012090038	ELPEN PHARMACEUTICAL CO. INC.	LU
Flusamer 50 microgramе/250 microgramе/doză pulbere de inhalat cu doze prestabilite	EE/H/0224/001	4501/2012/01	ELPEN PHARMACEUTICAL CO. INC.	RO
Flusamer 50 microgramе/500 microgramе/doză pulbere de inhalat cu doze prestabilite	EE/H/0224/002	4502/2012/01	ELPEN PHARMACEUTICAL CO. INC.	RO
Seretide Evohaler 25/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	not available	OGYI-T-8652/02	GLAXOSMITHKLINE KFT.	HU
Seretide Evohaler 25/250 mikrogramm/adag	not available	OGYI-T-8652/03	GLAXOSMITHKLINE 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
túlnyomásos inhalációs szuszpenzió				
Seretide Evohaler 25/50 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	not available	OGYI-T-8652/01	GLAXOSMITHKLINE KFT.	HU
Inaladuo Accuhaler 50 microgramos/250 microgramos/inhalación, polvo para inhalación.	not available	62.868	SMITHKLINE BEECHAM FARMA, S.A.	ES
Inaladuo Accuhaler 50 microgramos/500 microgramos/inhalación, polvo para inhalación.	not available	62.869	SMITHKLINE BEECHAM FARMA, S.A.	ES
Inaladuo Accuhaler 50 microgramos/100 microgramos/inhalación, polvo para inhalación.	not available	62.867	SMITHKLINE BEECHAM FARMA, S.A.	ES
Seretide Diskus 50/250 mikrogrami/devā pulveris inhalācijām	not available	00-0934	GLAXOSMITHKLINE LATVIA SIA	LV
Seretide Diskus 50/500 mikrogrami/devā pulveris inhalācijām	not available	00-0935	GLAXOSMITHKLINE LATVIA SIA	LV
Seretide Diskus 50/100 mikrogrami/devā pulveris inhalācijām	not available	00-0933	GLAXOSMITHKLINE LATVIA SIA	LV
Seretide 25/50 inhalasjonsaerosol, suspensjon	not available	00-1842	GLAXOSMITHKLINE AS	NO
Seretide 25/250 inhalasjonsaerosol, suspensjon	not available	00-1844	GLAXOSMITHKLINE AS	NO
Seretide 25/125	not available	00-1843	GLAXOSMITHKLINE AS	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
inhalasjonsaerosol, suspensjon				
Seretide Diskus 50/250 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/99/0481/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Seretide Diskus 50/100 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/99/0481/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Seretide Diskus 50/250 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/99/0481/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Seretide Diskus 50/500 mikrogramų/dozėje dozuoti įkvepiamieji milteliai	not available	LT/1/99/0481/004	GLAXOSMITHKLINE LIETUVA UAB	LT
ЕРФЛУЗАЛ ФОРСПИРО 50 МИКРОГРАМА + 250 МИКРОГРАМА /ДОЗА ПРАХ ЗА ИНХАЛАЦИЯ, ПРЕДВАРИТЕЛНО ДОЗИРАН	SE/H/1323/003	20150137	SANDOZ PHARMACEUTICALS D.D.	BG
AirFluSol Forspiro 50 mikrogramm/250 mikrogramm/adag adagolt inhalációs por	SE/H/1323/003	OGYI-T-22596/05	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/250 mikrogramm/adag adagolt inhalációs por	SE/H/1323/003	OGYI-T-22596/04	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/250	SE/H/1323/003	OGYI-T-22596/06	SANDOZ HUNGÁRIA 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
mikrogramm/adag adagolt inhalációs por				
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/01	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/02	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/03	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/04	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250 microgram/ doza, pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/05	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgram/250 microgram/doză pulbere unidoză de inhalat	SE/H/1323/003	7673/2015/06	S.C. SANDOZ S.R.L.	RO
AirFluSal® Forspiro® 50 microgram/250	SE/H/1323/003	7673/2015/07	S.C. SANDOZ S.R.L.	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
microgramе/ doza, pulbere unidoză de inhalat				
Salmeterol/Fluticasone Sandoz 50 mikrogram/250 mikrogram/dos Inhalationspulver, avdelad dos	SE/H/1323/003	50816	SANDOZ A/S	SE
Salmeterol/Fluticasonpropionaat Sandoz 25/125 microgram, aerosol, suspensie	NL/H/3709/001	RVG 118844	SANDOZ B.V.	NL
Salmeterol/Fluticasonpropionaat Sandoz 25/250 microgram, aerosol, suspensie	NL/H/3709/002	RVG 118845	SANDOZ B.V.	NL
PROPIONATE DE FLUTICASONE/SALMETEROL SANDOZ 125 microgrammes/ 25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	NL/H/3709/001	34009 301 720 4 9	SANDOZ	FR
PROPIONATE DE FLUTICASONE/SALMETEROL SANDOZ 250 microgrammes/ 25 microgrammes/dose, suspension pour inhalation en flacon pressurisé	NL/H/3709/002	34009 301 720 8 7	SANDOZ	FR
Sereflo 25	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	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
microgram/125 microgram per actuation pressurised inhalation, suspension.				
Sereflo 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	IE
AirFluSal Forspiro 50/500 microgram/dosis, inhalatiepoeder, voorverdeeld	SE/H/1447/002	RVG 116003	SANDOZ B.V.	NL
Casorol Forspiro, 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1447/002	51641	SANDOZ A/S	SE
Airflusal® Dosieraerosol 25 Mikrogramm/125 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	SE/H/1537/001	94827.00.00	CIPLA EUROPE NV	DE
Airflusal® Dosieraerosol 25 Mikrogramm/250 Mikrogramm pro Sprühstoß Druckgasinhalation, Suspension	SE/H/1537/002	94828.00.00	CIPLA EUROPE NV	DE
Zoreeda 25 mikrogram/125 mikrogram/dos inhalationsspray,	SE/H/1537/001	52740	CIPLA EUROPE NV	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
suspension				
Zoreeda 25 mikrogram/250 mikrogram/dos inhalationsspray, suspension	SE/H/1537/002	52741	CIPLA EUROPE NV	SE
Salmeterol/Fluticason Cipla 25 mikrogramu / 125 mikrogramu / dávka suspenze k inhalaci v tlakovém obalu	SE/H/1537/001	14/628/15-C	CIPLA EUROPE NV	CZ
Salmeterol/Fluticason Cipla 25 mikrogramu / 250 mikrogramu / dávka suspenze k inhalaci v tlakovém obalu	SE/H/1537/002	14/629/15-C	CIPLA EUROPE NV	CZ
Anasma Accuhaler 50 microgramos/500 microgramos/inhalación, polvo para inhalación.	not available	62.841	ALLEN FARMACÉUTICA, S.A	ES
Anasma Accuhaler 50 microgramos/250 microgramos/inhalación, polvo para inhalación	not available	62.840	ALLEN FARMACÉUTICA, S.A	ES
Anasma Accuhaler 50 microgramos/100 microgramos/inhalación, polvo para inhalación	not available	62.839	ALLEN FARMACÉUTICA, S.A	ES
Seretide 25/50 mikrogrami/deva aerosols inhalacijam, zem spiediena, suspensija	not available	01-0451	GLAXOSMITHKLINE LATVIA SIA	LV
Seretide 25/250	not available	01-0453	GLAXOSMITHKLINE LATVIA	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
mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija			SIA	
Seretide 25/125 mikrogrami/devā aerosols inhalācijām, zem spiediena, suspensija	not available	01-0452	GLAXOSMITHKLINE LATVIA SIA	LV
Seretide 50 Inhaler N Inhalacná suspenzia v tlakovom obale	not available	14/0113/01-S	GLAXOSMITHKLINE (IRELAND) LIMITED	SK
Seretide 125 Inhaler N inhalacná suspenzia v tlakovom obale	not available	14/0114/01-S	GLAXOSMITHKLINE (IRELAND) LIMITED	SK
Seretide 250 Inhaler N Inhalacna suspenzia v tlakovom obale	not available	14/0115/01-S	GLAXOSMITHKLINE (IRELAND) LIMITED	SK
Dimenium 50 mikrograma+100 mikrograma u jednoj dozi, prašak inhalata, dozirani	not available	HR-H-536223257	ELPEN PHARMACEUTICAL CO. INC.	HR
Dimenium 50 mikrograma+250 mikrograma u jednoj dozi, prašak inhalata, dozirani	not available	HR-H-505836307	ELPEN PHARMACEUTICAL CO. INC.	HR
Dimenium 50 mikrograma+500 mikrograma u jednoj dozi, prašak inhalata, dozirani	not available	HR-H-590491088	ELPEN PHARMACEUTICAL CO. INC.	HR
Sereflo 25	IE/H/0653/001	PA 1457/022/001	FANNIN LTD	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
microgram/125 microgram per actuation pressurised inhalation, suspension.				
Sereflo 25 microgram/250 microgram per actuation pressurised inhalation, suspension.	IE/H/653/002/DC	PA 1457/022/002	FANNIN LTD	IE
Saldisk 50 mikrograma+250 mikrograma u jednoj dozi, prasak inhalata, dozirani	not available	UP/I-530-09/12-01/696	SALVUS D.O.O.	HR
Saldisk 50 mikrograma+100 mikrograma u jednoj dozi, prasak inhalata, dozirani	not available	UP/I-530-09/12-01/695	SALVUS D.O.O.	HR
Saldisk 50 mikrograma+500 mikrograma u jednoj dozi, prasak inhalata, dozirani	not available	UP/I-530-09/12-01/697	SALVUS D.O.O.	HR
Fluticasona + Salmeterol Finao Biotech (100 µg + 50 µg)/ dose pó para inalação, em recipiente unidose	not available	5729728	FINAO BIOTECH, LDA	PT
Fluticasona + Salmeterol Finao Biotech (250 µg + 50 µg)/ dose pó para inalação, em recipiente unidose	not available	5729751	FINAO BIOTECH, 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
Fluticasona + Salmeterol Finao Biotech (500 µg + 50 µg)/ dose pó para inalação, em recipiente unidose	not available	5729769	FINAO BIOTECH, LDA	PT
Salmeterol/Fluticasone Cipla 25 mikrogram/125 mikrogram/dos inhalationsspray, suspension	SE/H/1208/001	49095	CIPLA EUROPE NV	SE
Salmeterol/Fluticasone Cipla 25 mikrogram/250 mikrogram/dos inhalationsspray, suspension	SE/H/1208/002	49096	CIPLA EUROPE NV	SE
FULLHALE 25 mikrogramov/125 mikrogramov/dávka inhalačná suspenzia v tlakovom obale	SE/H/1208/001	14/0135/14-S	CIPLA EUROPE NV	SK
SERROFLO 25 Mikrogramm / 125 Mikrogramm / Dosis Druckgasinhalation, Suspension	SE/H/1208/001	89763.00.00	CIPLA EUROPE NV	DE
FULLHALE 25 mikrogramov/250 mikrogramov/dávka inhalačná suspenzia v tlakovom obale	SE/H/1208/002	14/0136/14-S	CIPLA EUROPE NV	SK
SERROFLO 25 Mikrogramm / 250 Mikrogramm / Dosis Druckgasinhalation,	SE/H/1208/002	89764.00.00	CIPLA EUROPE NV	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
Suspension				
FULLHALE 25 mikrogramů/125 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	SE/H/1208/001	14/219/14-C	CIPLA EUROPE NV	CZ
FULLHALE 25 mikrogramů/250 mikrogramů/dávka suspenze k inhalaci v tlakovém obalu	SE/H/1208/002	14/220/14-C	CIPLA EUROPE NV	CZ
Salmeterol/Fluticasone Cipla 25 mikrogram/125 mikrogram/ dose inhalasjonsaerosol, suspensjon	SE/H/1208/001	13-9659	CIPLA EUROPE NV	NO
Salmeterol/Fluticasone Cipla 25 mikrogram/250 mikrogram/ dose inhalasjonsaerosol, suspensjon	SE/H/1208/002	13-9660	CIPLA EUROPE NV	NO
Seretide Evohaler 25/50 microgram per metered dose pressurised inhalation, suspension.	not available	19553	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Inhalok Forspiro 50 microgramos/250 microgramos/inhalación, polvo para inhalación (unidosis)	SE/H/1663/001	81.790	KERN PHARMA, S.L.	ES
Inhalok Forspiro 50 microgramos/500 microgramos/inhalación, polvo para inhalación	SE/H/1663/002	81.791	KERN PHARMA, S.L.	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
(unidosis)				
Flusamix Easyhaler 50 microgramos/500 microgramos/inhalación, polvo para inhalación	SE/H/1698/002	83644	ORION CORPORATION	ES
Flusamix Easyhaler, 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1698/002	56290	ORION CORPORATION	SE
AirFluSal Forspiro, (50 mikrogramów + 250 mikrogramów)/dawkę inhalacyjną, proszek do inhalacji, podzielony	SE/H/1448/001	23169	SANDOZ GMBH	PL
AirFluSal Forspiro, (50 mikrogramów + 500 mikrogramów)/dawkę inhalacyjną, proszek do inhalacji, podzielony	SE/H/1448/002	23170	SANDOZ GMBH	PL
Roadasan Forspiro, 50 mikrogram/250mikrogram/dos inhalationspulver, avdelad dos	SE/H/1448/001	51642	SANDOZ A/S	SE
Roadasan Forspiro, 50 mikrogram/500mikrogram/dos inhalationspulver, avdelad dos	SE/H/1448/002	51643	SANDOZ A/S	SE
Fusacomb Easyhaler 50 microgram/250 microgram/dose, inhalation powder	SE/H/1701/001	PL 27925/0093	ORION CORPORATION	UK
Fusacomb Easyhaler 50 microgram/500 microgram/dose,	SE/H/1701/002	PL 27925/0094	ORION CORPORATION	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
inhalation powder				
Fusacomb Easyhaler 50 microgram/500 microgram/dose, inhalation powder	SE/H/1701/002	56296	ORION CORPORATION	SE
Fusacomb Easyhaler 50 microgram/250 microgram/dose, inhalation powder	SE/H/1701/001	56295	ORION CORPORATION	SE
Seretide Evohaler 25 microgram/50 microgram per metered dose pressurised inhalation, suspension.	not available	MA 192/00905	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Seretide Evohaler 25 microgram/125 microgram per metered dose pressurised inhalation, suspension	not available	MA 192/00906	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Seretide Evohaler 25 microgram/250 microgram per metered dose pressurised inhalation, suspension	not available	MA 192/00907	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Brisomax Inalador 50 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703782	BIAL - PORTELA & Cª , SA	PT
Brisomax Inalador 125 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703881	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
Brisomax Inalador 250 microgramas/dose + 25 microgramas/dose suspensão pressurizada para inalação	not available	3703980	BIAL - PORTELA & Cª , SA	PT
AirFluSol Forspiro 50 mikrogramm/500 mikrogramm/adag adagolt inhalációs por	SE/H/1323/002	OGYI-T-22596/01	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/500 mikrogramm/adag adagolt inhalációs por	SE/H/1323/002	OGYI-T-22596/02	SANDOZ HUNGÁRIA KFT	HU
AirFluSol Forspiro 50 mikrogramm/500 mikrogramm/adag adagolt inhalációs por	SE/H/1323/002	OGYI-T-22596/03	SANDOZ HUNGÁRIA KFT	HU
AirFluSal Forspiro 50 microgram/500 microgram/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/01	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgram/500 microgram/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/02	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgram/500 microgram/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/03	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgram/500	SE/H/1323/002	6124/2014/04	S.C. SANDOZ S.R.L.	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
microgramе/doză pulbere unidoză de inhalat				
AirFluSal Forspiro 50 microgramе/500 microgramе/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/05	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgramе/500 microgramе/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/06	S.C. SANDOZ S.R.L.	RO
AirFluSal Forspiro 50 microgramе/500 microgramе/doză pulbere unidoză de inhalat	SE/H/1323/002	6124/2014/07	S.C. SANDOZ S.R.L.	RO
Airflusal® Forspiro® 50 Mikrogramm/500 Mikrogramm/Dosis einzeldosiertes Pulver zur Inhalation	SE/H/1323/002	89724.00.00	HEXAL AG	DE
Salmeterol/Fluticasone Sandoz 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1323/002	49086	SANDOZ A/S	SE
ЕРФЛУЗАЛ ФОРСПИРО 50 МИКРОГРАМА+ 500 МИКРОГРАМА/ДОЗА ПРАХ ЗА ИНХАЛАЦИЯ, ПРЕДВАРИТЕЛНО	SE/H/1323/002	20140018	SANDOZ PHARMACEUTICALS D.D.	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
ДОЗИРАН				
AirFluSal Forspiro 50/500 microgram/dosis, inhalatiepoeder, voorverdeeld	SE/H/1447/002	RVG 116003	SANDOZ B.V.	NL
Casorol Forspiro, 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1447/002	51641	SANDOZ A/S	SE
Airflusal Forspiro 50 mikrogramov/250 mikrogramov, dávkovaný inhalačný prášok	SE/H/1405/001	14/0327/14-S	SANDOZ PHARMACEUTICALS D.D.	SK
Airflusal Forspiro 50 mikrogramov/500 mikrogramov, dávkovaný inhalačný prášok	SE/H/1405/002	14/0328/14-S	SANDOZ PHARMACEUTICALS D.D.	SK
AirFluSal Forspiro, 50/250 mikrogrammi annuses, annustatud inhalatsioonipulber	SE/H/1405/001	858314	SANDOZ PHARMACEUTICALS D.D.	EE
AirFluSal Forspiro 50/250 mikrogrami/devā, pulveris inhalācijām, dozēts	SE/H/1405/001	14-0224	SANDOZ PHARMACEUTICALS D.D.	LV
AirFluSal Forspiro 50/500 mikrogrami/devā, pulveris inhalācijām, dozēts	SE/H/1405/002	14-0225	SANDOZ PHARMACEUTICALS D.D.	LV
AirFluSal Forspiro, 50/500 mikrogrammi annuses, annustatud inhalatsioonipulber	SE/H/1405/002	858214	SANDOZ PHARMACEUTICALS D.D.	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
AIRFLUSAN FORSPIRO 50 MIKROGRAMŮ/250 MIKROGRAMŮ dávkovaný prášek k inhalaci	SE/H/1405/001	14/411/14-C	SANDOZ S.R.O.	CZ
AIRFLUSAN FORSPIRO 50 MIKROGRAMŮ/500 MIKROGRAMŮ dávkovaný prášek k inhalaci	SE/H/1405/002	14/412/14-C	SANDOZ S.R.O.	CZ
AirFluSal Forspiro 50 mikrograma + 500 mikrograma u jednoj dozi, prašak inhalata, dozirani	SE/H/1405/002	HR-H-007513419	SANDOZ D.O.O.	HR
AirFluSal Forspiro 50 mikrograma + 250 mikrograma u jednoj dozi, prašak inhalata, dozirani	SE/H/1405/001	HR-H-597688265	SANDOZ D.O.O.	HR
Airflusan Forspiro 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/001	H/14/01918/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/001	H/14/01918/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje,	SE/H/1405/001	H/14/01918/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
odmerjeni				
Airflusan Forspiro 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/002	H/14/01918/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/002	H/14/01918/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/002	H/14/01918/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/002	H/14/01918/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/500 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/002	H/14/01918/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Molduo Forspiro, 50 mikrogram/250 mikrogram inhalationspulver	SE/H/1405/001	50645	SANDOZ A/S	SE
Molduo Forspiro, 50 mikrogram/500 mikrogram inhalationspulver	SE/H/1405/002	50646	SANDOZ A/S	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
Airflusan Forspiro 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/001	H/14/01918/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Airflusan Forspiro 50 mikrogramov/250 mikrogramov/odmerek prašek za inhaliranje, odmerjeni	SE/H/1405/001	H/14/01918/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
AirFluSal Forspiro, 50 μικρογραμμάρια/250 μικρογραμμάρια/δόση, κόνις για εισπνοή σε δόσεις	SE/H/1405/001	66491/06-10-2015	SANDOZ PHARMACEUTICALS D.D.	GR
AirFluSal Forspiro, 50 μικρογραμμάρια/500 μικρογραμμάρια/δόση, κόνις για εισπνοή σε δόσεις	SE/H/1405/002	66492/06-10-2015	SANDOZ PHARMACEUTICALS D.D.	GR
AirFluSal® Forspiro® 50 microgram/500 microgram/dose, inhalation powder, predispensed	SE/H/1405/002	PA0711/237/002	ROWEX LTD	IE
AirFluSal® Forspiro® 50 microgram/250 microgram/dose, inhalation powder, predispensed	SE/H/1405/001	PA0711/237/001	ROWEX LTD	IE
AirFluSal Forspiro 50 microgram/500 microgram per actuation inhalation powder, pre-	SE/H/1469/002	PL 04416/1431	SANDOZ 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
dispensed				
Salfluae Forspiro 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1469/002	51645	SANDOZ A/S	SE
Safumide Easyhaler 50 microgram/250 microgram/dose, inhalation powder	SE/H/1696/001	56285	ORION CORPORATION	SE
Safumide Easyhaler 50 microgram/500 microgram/dose, inhalation powder	SE/H/1696/02/DC	56286	ORION CORPORATION	SE
SALFLUAIR EASYHALER 250 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/001	34009 301 756 8 2	ORION CORPORATION	FR
SALFLUAIR EASYHALER 250 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/001	34009 550 634 9 0	ORION CORPORATION	FR
SALFLUAIR EASYHALER 250 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/001	34009 550 635 0 6	ORION CORPORATION	FR
SALFLUAIR EASYHALER 500 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/002	34009 301 756 9 9	ORION CORPORATION	FR
SALFLUAIR EASYHALER 500 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/002	34009 550 635 1 3	ORION CORPORATION	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
SALFLUAIR EASYHALER 500 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/1696/002	34009 550 635 2 0	ORION CORPORATION	FR
Seretide Diskus 50 microgrammes/250 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/002	2009 02 0198	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide Diskus 50 mikrogram / 100 mikrogram/dos inhalationspulver, avdelad dos	SE/H/169/001	13822	GLAXOSMITHKLINE OY	FI
Seretide Diskus 50 mikrogram / 250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/0169/002	13823	GLAXOSMITHKLINE OY	FI
Seretide Diskus 50 mikrogram / 500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/0169/003	13824	GLAXOSMITHKLINE OY	FI
Seretide Diskus 50 microgram/100 microgram/dosis, inhalatiepoeder, voorverdeeld	SE/H/169/001	BE200855	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide Diskus 50 Mikrogramm/100 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/0169/001	BE200855	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Seretide Diskus 50 Mikrogramm/250 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/169/002	BE200873	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SeretideDiskus 50 microgram/250 microgram/dosis, inhalatiepoeder, voorverdeeld.	SE/H/0169/002	BE200873	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide Diskus 50 microgram/500 microgram/dosis, inhalatiepoeder, voorverdeeld.	SE/H/169/003	BE200882	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide Diskus 50 Mikrogramm/500 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/169/003	BE200882	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide Diskus 50 Mikrogramm/100 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/169/001	2009 02 0197	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide Diskus 50 Mikrogramm/250 Mikrogramm/Dosis, einzeldosiertes Pulver zur Inhalation	SE/H/0169/002	2009 02 0198	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide Diskus 50 Mikrogramm/500 Mikrogramm/Dosis, einzeldosiertes Pulver	SE/H/0169/003	2009 02 0199	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
zur Inhalation				
Seretide Diskus 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/003	034371068	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 mikrogramov/100 mikrogramov dávkovaný inhalačný prášok	SE/H/0169/001	14/0010/17-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
atmadisc mite 50 µg/100 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/169/001	44915.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
atmadisc 50 µg/250 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/0169/002	44915.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
atmadisc forte 50 µg/500 µg Diskus Einzeldosiertes Pulver zur Inhalation	SE/H/169/003	44915.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Seretaide Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/169/001	4812889	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide Diskus® levis 50 Mikrogramm/100 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/169/001	1-22902	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seretide Diskus 50 microgrammes/100 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/001	BE200855	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Seretide	SE/H/169/001	30386	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Diskus 50 mikrog/100 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/169/001	13822	GLAXOSMITHKLINE OY	FI
SERETIDE DISKUS 100 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/0169/001	NL25846	LABORATOIRE GLAXOSMITHKLINE	FR
Seretide Diskus 50/100 μικρογραμμάρια/δόση κόνεως για εισπνοή σε δόσεις	SE/H/169/001	2439201	GLAXOSMITHKLINE AEBE	GR
Seretide 100 Diskus 50 microgram/100 microgram/dose inhalation powder, pre-dispensed.	SE/H/169/001	PA 1077/46/1	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seretide Diskus 50 microgrammi/100 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/001	034371017	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 microgrammi/100 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/001	034371043	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 microgrammi/100 microgrammi/dose di	SE/H/0169/001	034371070	GLAXOSMITHKLINE 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
polvere per inalazione in contenitore monodose				
Seretide Diskus 50 microgrammes/100 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/001	2009 02 0197	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide Diskus 50 microgram/100 microgram/dosis - inhalatiepoeder, voorverdeeld.	SE/H/169/001	RVG 23529	GLAXOSMITHKLINE B.V.	NL
Seretaide Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/169/001	2874287	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/169/001	2874386	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/001	2874485	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/100 microgramas/dose pó para inalação, em recipiente unidose	SE/H/169/001	4812780	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide Accuhaler 50 microgramos/100	SE/H/0169/001	62.667	GLAXOSMITHKLINE, S.A.	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
microgramos/inhalación, polvo para inhalación				
Seretide Diskus mite 50 mikrogram/100 mikrogram/dos inhalationspulver, avdelad dos	SE/H/169/001	14591	GLAXOSMITHKLINE AB	SE
Seretide Accuhaler 50 microgram /100 microgram /dose inhalation powder, pre-dispensed	SE/H/169/001	PL10949/0314	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Seretide Diskus® standard 50 Mikrogramm/250 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/0169/002	1-22901	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seretide Diskus 50 microgrammes/250 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/002	BE200873	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide	SE/H/169/002	30387	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Diskus 50 mikrog/250 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/169/002	13823	GLAXOSMITHKLINE OY	FI
SERETIDE DISKUS 250 microgrammes/50 microgrammes/dose, poudre pour inhalation	SE/H/0169/002	NL25847	LABORATOIRE GLAXOSMITHKLINE	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
en récipient unidose				
Seretide Diskus 50/250 μικρογραμμάρια/δόση κόνεως για εισπνοή σε δόσεις	SE/H/0169/002	2439202	GLAXOSMITHKLINE AEBE	GR
Seretide 250 Diskus 50 microgram/250 microgram/dose inhalation powder, pre-dispensed.	SE/H/169/002	PA 1077/46/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seretide Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/002	034371029	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/002	034371056	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/002	034371082	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/169/002	14592	GLAXOSMITHKLINE AB	SE
Seretide Diskus 50 microgram/250 microgram/dosis - inhalatiepoeder,	SE/H/0169/002	RVG 23530	GLAXOSMITHKLINE 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
voorverdeeld				
Seretaide Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/002	4812988	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/002	2874782	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/002	2874683	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/002	2874584	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/250 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/002	4813085	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide Accuhaler 50 microgramos/250 microgramos/inhalación, polvo para inhalación	SE/H/0169/002	62.668	GLAXOSMITHKLINE S.A.	ES
Seretide Accuhaler 50 microgram /250 microgram /dose inhalation powder, pre-	SE/H/0169/002	PL10949/0315	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	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
dispensed				
Seretide Diskus® forte 50 Mikrogramm/500 Mikrogramm - einzeldosiertes Pulver zur Inhalation	SE/H/0169/003	1-22900	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seretide Diskus 50 microgrammes/500 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/003	BE200882	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seretide	SE/H/169/003	30388	GLAXOSMITHKLINE PHARMA A/S	DK
Seretide Diskus 50 mikrog/500 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/169/003	13824	GLAXOSMITHKLINE OY	FI
SERETIDE DISKUS 500 microgrammes/50 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/0169/003	NL25848	LABORATOIRE GLAXOSMITHKLINE	FR
Seretide Diskus 50/500 μικρογραμμάρια/δόση κόνεως για εισπνοή σε δόσεις	SE/H/0169/003	2439203	GLAXOSMITHKLINE AEBE	GR
Seretide 500 Diskus 50 microgram/500 microgram/dose inhalation powder, pre-dispensed.	SE/H/169/003	PA 1077/46/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seretide Diskus 50 microgrammi/500	SE/H/0169/003	034371031	GLAXOSMITHKLINE 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
microgrammi/dose di polvere per inalazione in contenitore monodose				
Seretide Diskus 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0169/003	034371094	GLAXOSMITHKLINE S.P.A.	IT
Seretide Diskus 50 microgrammes/500 microgrammes/dose, poudre pour inhalation en récipient unidose	SE/H/169/003	2009 02 0199	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seretide Diskus 50 microgram/500 microgram/dosis - inhalatiepoeder, voorverdeeld	SE/H/169/003	RVG 23531	GLAXOSMITHKLINE B.V.	NL
Seretaide Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/003	4813283	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/003	4813184	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/003	2875086	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretaide Diskus, 50	SE/H/0169/003	2874980	GLAXO WELLCOME	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
microgramas/500 microgramas/dose pó para inalação, em recipiente unidose			FARMACÊUTICA, LDA	
Seretaide Diskus, 50 microgramas/500 microgramas/dose pó para inalação, em recipiente unidose	SE/H/0169/003	2874881	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Seretide Accuhaler 50 microgramos/500 microgramos/inhalación, polvo para inhalación	SE/H/0169/003	62.669	GLAXOSMITHKLINE S.A.	ES
Seretide Diskus forte 50 mikrogram/500 mikrogram/dos inhalationspulver, avdelad dos	SE/H/169/003	14593	GLAXOSMITHKLINE AB	SE
Seretide Accuhaler 50 microgram /500 microgram /dose inhalation powder, predispensed	SE/H/169/003	PL10949/0316	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Seretide Diskus 50 microgram/500 microgram/ dose inhalation powder, pre-dispensed.	SE/H/169/003	MA 192/00903	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Seretide Diskus 50 microgram/250 microgram/ dose inhalation powder, pre-dispensed	SE/H/169/002	MA 192/00902	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Seretide Diskus 100,	SE/H/0169/001	272599	GLAXOSMITHKLINE	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
50/100 mikrogrammi annustatud inhalatsioonipulber			(IRELAND) LIMITED	
Seretide Diskus 50 mikrogramů/100 mikrogramů Dávkovaný prášek k inhalaci	SE/H/0169/001	14/101/00-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Seretide Diskus 50 mikrogramů/250 mikrogramů Dávkovaný prášek k inhalaci	SE/H/0169/002	14/102/00-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Seretide Diskus 50 mikrogramů/500 mikrogramů Dávkovaný prášek k inhalaci	SE/H/0169/003	14/103/00-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Seretide Diskus 250, 50/250 mikrogrammi annustatud inhalatsioonipulber	SE/H/0169/002	272499	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Seretide Diskus 500, 50/500 mikrogrammi annustatud inhalatsioonipulber	SE/H/0169/003	458604	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Seretide Diskus 50/100 mikrogramm/adag adagolt inhalációs por	SE/H/169/001	OGYI-T-7626/01	GLAXOSMITHKLINE KFT.	HU
Seretide Diskus 50/500 mikrogramm/adag adagolt inhalációs por	SE/H/0169/003	OGYI-T-7626/03	GLAXOSMITHKLINE KFT.	HU
Seretide Diskus 50/250 mikrogramm/adag adagolt inhalációs por	SE/H/0169/002	OGYI-T-7626/02	GLAXOSMITHKLINE KFT.	HU
Seretide Diskus 50 mikrogramov/250	SE/H/0169/002	14/0011/17-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	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
mikrogramov dávkovaný inhalačný prášok				
Seretide Diskus 50 mikrogramov/500 mikrogramov dávkovaný inhalačný prášok	SE/H/0169/003	14/0012/17-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Seretide 50 míkróg/100 míkróg/skammt innöndunarduft, afmældir skammtar í Diskos	SE/H/169/003	980374 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide 50 míkróg/250 míkróg/skammt innöndunarduft, afmældir skammtar í Diskos	SE/H/169/003	980375 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide Diskus 50 mikrograma+500 mikrograma u jednoj dozi, prašak inhalata, dozirani	SE/H/169/003	HR-H-251056275	GLAXOSMITHKLINE D.O.O.	HR
Seretide Diskus 50 mikrograma+250 mikrograma u jednoj dozi, prašak inhalata, dozirani	SE/H/169/002	HR-H-562619185	GLAXOSMITHKLINE D.O.O.	HR
Seretide Diskus 50 mikrograma+100 mikrograma u jednoj dozi, prašak inhalata, dozirani	SE/H/169/001	HR-H-491933173	GLAXOSMITHKLINE D.O.O.	HR
Seretide Diskus 50 micrograme/100 micrograme pulbere de	SE/H/0169/001	9570/2017/01	GLAXOSMITHKLINE (IRELAND) LIMITED	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
inhalat				
Seretide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0169/001	9570/2017/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0169/001	9570/2017/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/100 microgram pulbere de inhalat	SE/H/0169/001	9570/2017/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0169/002	9571/2017/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0169/002	9571/2017/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0169/002	9571/2017/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0169/002	9571/2017/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/250 microgram pulbere de inhalat	SE/H/0169/002	9571/2017/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50	SE/H/0169/003	9572/2017/01	GLAXOSMITHKLINE	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
microgram/500 microgram pulbere de inhalat			(IRELAND) LIMITED	
Seretide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0169/003	9572/2017/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0169/003	9572/2017/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0169/003	9572/2017/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide Diskus 50 microgram/500 microgram pulbere de inhalat	SE/H/0169/003	9572/2017/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Seretide 50 míkróg/500 míkróg/skammt innöndunarduft, afmældir skammtar í Diskos	SE/H/169/003	980376 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
Seretide Diskus 50/100 μικρογραμμάρια/δόση κόνεως για εισπνοή σε δόσεις	SE/H/169/001	022630	GLAXOSMITHKLINE (CYPRUS) LIMITED	CY
Seretide Diskus 50/250 μικρογραμμάρια/δόση κόνεως για εισπνοή σε δόσεις	SE/H/169/002	022631	GLAXOSMITHKLINE (CYPRUS) LIMITED	CY
Seretide Diskus 50/500 μικρογραμμάρια/δόση	SE/H/169/003	022632	GLAXOSMITHKLINE (CYPRUS) LIMITED	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
κόνεως για εισπνοή σε δόσεις				
Salmeterol/Fluticasonpropionat Easyhaler 50 Mikrogramm/250 Mikrogramm pro Inhalation, Pulver zur Inhalation	SE/H/1695/01/DC	99810.00.00	ORION CORPORATION	DE
Salmeterol/Fluticasonpropionat Easyhaler 50 Mikrogramm/500 Mikrogramm pro Inhalation, Pulver zur Inhalation	SE/H/1695/02/DC	99811.00.00	ORION CORPORATION	DE
Aerion Easyhaler 50 mikrogram/250 mikrogram/dos inhalationspulver	SE/H/1695/001	56283	ORION CORPORATION	SE
Aerion Easyhaler 50 mikrogram/500 mikrogram/dos inhalationspulver	SE/H/1695/002	56284	ORION CORPORATION	SE
AirFluSal Aerosol 25/125 microgram, aerosol, suspensie	NL/H/3710/001	RVG 118836	SANDOZ B.V.	NL
AirFluSal Aerosol 25/250 microgram, aerosol, suspensie	NL/H/3710/002	RVG 118837	SANDOZ B.V.	NL
Zoreeda 25 Mikrogramm/125 Mikrogramm/Dosis Druckgasinhalation, Suspension	AT/H/0517/001	135937	CIPLA EUROPE NV	AT
Zoreeda 25	AT/H/0517/002	135938	CIPLA EUROPE NV	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
Mikrogramm/250 Mikrogramm/Dosis Druckgasinhalation, Suspension				
Salmeterol/flutikazon Cipla 25 mikrograma + 125 mikrograma u jednoj dozi, stlačeni inhalat, suspenzija	AT/H/0517/001	HR-H-607845986	CIPLA EUROPE NV	HR
Salmeterol/flutikazon Cipla 25 mikrograma + 250 mikrograma u jednoj dozi, stlačeni inhalat, suspenzija	AT/H/0517/002	HR-H-705296771	CIPLA EUROPE NV	HR
SALMETEROLO e FLUTICASONE DOC 25 microgrammi / 125 microgrammi/erogazione sospensione pressurizzata per inalazione	AT/H/0517/001	043019013	CIPLA EUROPE NV	IT
SALMETEROLO e FLUTICASONE DOC 25 microgrammi / 250 microgrammi/erogazione sospensione pressurizzata per inalazione	AT/H/0517/002	043019025	CIPLA EUROPE NV	IT
Fullhale 25 mikrogramm/125 mikrogramm/adag túlnyomásos inhalációs szuszpenzió	AT/H/0517/001	OGYI-T-22783/01	CIPLA EUROPE NV	HU
Fullhale 25	AT/H/0517/002	OGYI-T-22783/02	CIPLA EUROPE NV	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
mikrogramm/250 mikrogramm/adag túlnyomásos inhalációs szuszpenzió				
Salmeterol/Fluticasona Kern Pharma 25 microgramos/125 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	81.181	KERN PHARMA, S.L.	ES
Salmeterol/Fluticasona Kern Pharma 25 microgramos/250 microgramos/inhalación, suspensión para inhalación en envase a presión	not available	81.182	KERN PHARMA, S.L.	ES
Stalpex 50 microgram/500 microgram/dose inhalation powder, pre-dispensed	IE/H/0928/001	PL 25258/0296	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Salflutin 50 microgram/500 microgram/dosis inhalatiepoeder, voorverdeeld	IE/H/0928/001	RVG 120133	GLENMARK PHARMACEUTICALS NORDIC AB	NL
Salmex 50 microgram/500 microgram/dose inhalation powder, pre-dispensed	IE/H/0928/001	MA1357/00101	GLENMARK PHARMACEUTICALS NORDIC AB	MT
Stalpex 50	IE/H/0928/001	PA22815/001/001	GLENMARK	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
microgram/500 microgram/dose inhalation powder, pre-dispensed			PHARMACEUTICALS NORDIC AB	
Aliflus Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/002	034463051	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/100 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/001	034463075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/003	034463099	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/100 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/001	034463048	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/100 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/001	034463012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/002	034463087	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
Aliflus Diskus 50 microgrammi/250 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/002	034463024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/003	034463036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus Diskus 50 microgrammi/500 microgrammi/dose di polvere per inalazione in contenitore monodose	SE/H/0170/003	034463063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus 25 microgrammi/125 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5771/002	034463113	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus 25 microgrammi/250 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5771/003	034463125	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Aliflus 25 microgrammi/50 microgrammi/dose sospensione pressurizzata per inalazione	DE/H/5771/001	034463101	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Salmeterol/Fluticason	SE/H/1797/001	IS/1/19/047/01	WELLNEX GMBH	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
Wellnex 50 míkrógrömm/100 míkrógrömm/skammt innöndunarduft, afmældir skammtar				
Salmeterol/Fluticason Wellnex 50 míkrógrömm/250 míkrógrömm/skammt innöndunarduft, afmældir skammtar.	SE/H/1797/002	IS/1/19/047/02	WELLNEX GMBH	IS
Inhatec 50 microgram/ 100 microgram/ dose inhalation powder, pre-dispensed	SE/H/1797/001	PL 49518/0001	WELLNEX GMBH	UK
Inhatec 50 microgram/ 250 microgram/ dose inhalation powder, pre-dispensed	SE/H/1797/002	PL 49518/0002	WELLNEX GMBH	UK
Salmeterol/Fluticasonpropionaat Wellnex 50 microgram/100 microgram inhalatiepoeder, voorverdeeld.	SE/H/1797/001	RVG 122998	WELLNEX GMBH	NL
Salmeterol/Fluticasonpropionaat Wellnex 50 microgram/250 microgram inhalatiepoeder, voorverdeeld.	SE/H/1797/002	RVG 123002	WELLNEX GMBH	NL
Salmeterol/Fluticasone Wellnex 50 mikrogram/100	SE/H/1797/001	57731	WELLNEX GMBH	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
mikrogram/dos inhalationspulver, avdelad dos				
Salmeterol/Fluticasone Wellnex 50 mikrogram/250 mikrogram/dos inhalationspulver, avdelad dos	SE/H/1797/002	57732	WELLNEX GMBH	SE
Salmeterol/Fluticasone Wellnex 50 mikrogram/100 mikrogram inhalasjonspulver, dosedispensert	SE/H/1797/001	18-12213	WELLNEX GMBH	NO
Salmeterol/Fluticasone Wellnex 50 mikrogram/250 mikrogram inhalasjonspulver, dosedispensert	SE/H/1797/002	18-12214	WELLNEX GMBH	NO
Salmeterol/Fluticasone Wellnex 50 mikrog/100 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/1797/001	36053	WELLNEX GMBH	FI
Salmeterol/Fluticasone Wellnex 50 mikrog/250 mikrog/annos inhalaatiojauhe, annosteltu	SE/H/1797/002	36054	WELLNEX GMBH	FI
Salmeterol/Fluticason Wellnex 50 mikrogram/100	SE/H/1797/001	61027	WELLNEX GMBH	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
mikrogram/dosis inhalationspulver, afdelt				
Salmeterol/Fluticason Wellnex 50 mikrogram/250 mikrogram/dosis inhalationspulver, afdelt	SE/H/1797/002	61028	WELLNEX GMBH	DK
Seretide Diskus 50/250 inhalasjonspulver, dosedispenert	not available	98/4518	GLAXOSMITHKLINE AS	NO
Seretide Diskus 50/100 inhalasjonspulver, dosedispenert	not available	98/4517	GLAXOSMITHKLINE AS	NO
Seretide Diskus 50/500 inhalasjonspulver, dosedispenert	not available	98/4519	GLAXOSMITHKLINE AS	NO
Seretide Inhaler 25/125 mikrogramai/dozėje suslėgtoji įkvepiamoji suspensija	not available	LT/1/01/1576/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Seretide Inhaler 25/50 mikrogramų/dozėje suslėgtoji įkvepiamoji suspensija	not available	LT/1/01/1576/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Seretide Inhaler 25/250 mikrogramų/dozėje suslėgtoji įkvepiamoji suspensija	not available	LT/1/01/1576/003	GLAXOSMITHKLINE LIETUVA UAB	LT