26 October 2017
EMA/715509/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: influenza vaccine (split virion, inactivated) (non centrally authorised products)

Procedure no.: PSUSA/00010298/201703

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

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
Afluria Sospensione iniettabile, in siringa preriempita. Vaccino influenzale (virione split, inattivato)	DE/H/1938/001	043216011	SEQIRUS GMBH	IT
Afluria Sospensione iniettabile, in siringa preriempita. Vaccino influenzale (virione split, inattivato)	DE/H/1938/001	043216023	SEQIRUS GMBH	IT
Afluria Sospensione iniettabile, in siringa preriempita. Vaccino influenzale (virione split, inattivato)	DE/H/1938/001	043216035	SEQIRUS GMBH	IT
Afluria Sospensione iniettabile, in siringa preriempita. Vaccino influenzale (virione split, inattivato)	DE/H/1938/001	043216047	SEQIRUS GMBH	IT
Afluria Suspensão injetável, em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/1938/001	5643309	SEQIRUS GMBH	PT
Afluria Suspensão injetável, em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/1938/001	5643317	SEQIRUS GMBH	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
Afluria Suspensão injetável, em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/1938/001	5643325	SEQIRUS GMBH	PT
Afluria Suspensão injetável, em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/1938/001	5643333	SEQIRUS GMBH	PT
Afluria suspensie injectabila, în seringa preumpluta Vaccin gripal (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/03	SEQIRUS GMBH	RO
Afluria suspensie injectabilă, în seringă preumplută Vaccin gripal* (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/01	SEQIRUS GMBH	RO
Afluria suspensie injectabilă, în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/02	SEQIRUS GMBH	RO
Afluria suspensie injectabilă, în seringă preumplută Vaccin gripal (virion fragmentat,	DE/H/1938/001	7123/2014/04	SEQIRUS GMBH	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
inactivat)				
Afluria suspensie voor injectie, in een voorgevulde spuit. Influenzavaccin (split virus, geïnactiveerd)	DE/H/1938/001	1716/06040039	SEQIRUS GMBH	LU
Afluria Suspension injectable dans une seringue préremplie. accin contre l'influenza (virion fragmenté, inactivé)	DE/H/1938/001	1716/06040039	SEQIRUS GMBH	LU
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	705470 – 2	SEQIRUS GMBH	ES
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	705471 – 9	SEQIRUS GMBH	ES
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	702570 – 2	SEQIRUS GMBH	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
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	702573 – 3	SEQIRUS GMBH	ES
AFLURIA Ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο γρίπης (τμήμα ιού, αδρανοποιημένο)	DE/H/1938/001	21056	SEQIRUS GMBH	GR
Afluria, injekcní suspenze v predplnené injekcní stríkacce Vakcína proti chřipce (štěpený virion, inaktivovaná)	DE/H/1938/001	59/241/14-C	SEQIRUS GMBH	CZ
Afluria, injektionsvæske, suspension, fyldt injektionssprøjte	DE/H/1938/001	37439	SEQIRUS GMBH	DK
AFLURIA, Suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/1938/001	34009 279 598 9 9	SEQIRUS GMBH	FR
AFLURIA, Suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/1938/001	34009 279 602 6 0	SEQIRUS GMBH	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
AFLURIA, Suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/1938/001	34009 279 600 3 1	SEQIRUS GMBH	FR
AFLURIA, Suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/1938/001	34009 279 599 5 0	SEQIRUS GMBH	FR
a-RIX Injektionssuspension in einer Fertigspritze Grippeimpfung (fragmentiertes, inaktiviertes Virion)	DE/H/0124/001	BE147244	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
a-RIX, suspensie voor injectie in een voorgevulde spuit Griepvaccin (gefragmenteerd, geïnactiveerd virion)	DE/H/0124/001	BE147244	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
a-RIX, suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/0124/001	BE147244	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
a-RIX, suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/0124/001	2008089872	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
a-RIX, suspension injectable en seringue préremplie Vaccin grippal (virion fragmenté, inactivé)	DE/H/0124/001	2008089872	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
a-RIX-Tetra Injektionssuspension in Fertigspritze Grippeimpfung (Spalt-Virion, inaktiviert)	DE/H/1939/001	BE456924	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
a-RIX-Tetra, suspensie voor injectie in een voorgevulde spuit Griepvaccin (gefragmenteerd, geïnactiveerd virion)	DE/H/1939/001	BE456924	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Enzira Suspension for injection, pre-filled syringe Influenza vaccine (Split Virion, inactivated)	DE/H/1938/001	PA1373/001/001	SEQIRUS GMBH	IE
Enzira® Suspension for injection, pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1938/001	PL 22236/0001	SEQIRUS GMBH	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
Fluarix 2013/2014, suspensie voor injectie in een voorgevulde spuit Influenzavaccin (gesplitst virion, geïnactiveerd)	DE/H/0124/001	RVG 22307	GLAXOSMITHKLINE B.V.	NL
Fluarix injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/0124/001	LT/1/05/0274/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/0124/001	LT/1/05/0274/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/0124/001	LT/1/5/0274/004	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/0124/001	LT/1/05/0274/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu	DE/H/0124/001	LT/1/05/0274/005	GLAXOSMITHKLINE LIETUVA UAB	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
fragmentu, inaktyvuota)				
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)	DE/H/0124/001	LT/1/5/0274/006	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)	DE/H/0124/001	LT/1/5/0274/007	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)	DE/H/0124/001	LT/1/05/0274/008	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)	DE/H/0124/001	LT/1/5/0274/009	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix injekcine suspensija užpildytame švirkšte|Vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)	DE/H/0124/001	LT/1/05/0274/010	GLAXOSMITHKLINE LIETUVA UAB	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
Fluarix injekcná suspenzia v naplnenej injekcnej striekacke Ockovacia látka proti chrípke (štiepený virión, inaktivovaná)	DE/H/0124/001	59/0270/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Fluarix injektioneste, suspensio esitäytetyssä ruiskussa Influenssarokote (virusfragmentit, inaktivoitu)	DE/H/0124/001	13153	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245230	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	29245242	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	29245255	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
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	29245267	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245242	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245255	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245178	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245180	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus	DE/H/0124/001	029245192	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
frammentati, inattivati)				
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245204	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245216	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati)	DE/H/0124/001	029245228	GLAXOSMITHKLINE S.P.A.	IT
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	3088/2010/04	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/07	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/01	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/07	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/08	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/09	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/10	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat,	DE/H/0124/001	8244/2015/04	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
inactivat)				
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/05	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/06	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/06	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/02	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Fluarix suspensie injectabilă în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/0124/001	8244/2015/03	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Fluarix suspensija injekcijām pilnšļircē Pretgripas vakcīna (šķelts, inaktivēts virions) Vaccinum influenzae inactivatum ex virorum fragmentis praeparatum	DE/H/0124/001	05-0197	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Fluarix suspension for injection in a pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/0124/001	PA 1077/025/001	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Fluarix suspension for injection in a pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/0124/001	MA 172/01001	SMITHKLINE BEECHAM PLC	MT
FLUARIX suspension injectable en seringue préremplie. Vaccin grippal inactivé à virion fragmenté	DE/H/124/01	NL10811	LABORATOIRE GLAXOSMITHKLINE	FR
Fluarix suspensión inyectable en jeringa precargada Vacuna antigripal (de virus fraccionados e inactivados)	DE/H/0124/001	60.772	GLAXOSMITHKLINE, S.A.	ES
FLUARIX suspenzija za injekciju u napunjenoj štrcaljki Cjepivo protiv	DE/H/0124/001	HR-H-996462312	GLAXOSMITHKLINE D.O.O.	HR

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influence (fragmentirani virioni), inaktivirano				
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi. cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/009	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/003	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/004	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/005	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/006	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
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/007	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/001	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/010	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/008	GLAXOSMITHKLINE D.O.O.	SI
Fluarix suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/0124/001	H/04/00626/002	GLAXOSMITHKLINE D.O.O.	SI
Fluarix szuszpenziós injekció eloretöltött fecskendoben Influenza vakcina (split-vírus,	DE/H/0124/001	OGYI-T-8421/01	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
inaktivált)				
Fluarix szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (split-vírus, inaktivált)	DE/H/0124/001	OGYI-T-8421/02	GLAXOSMITHKLINE KFT.	HU
Fluarix Tetra injekcná suspenzia naplnená v injekcnej striekacke Ockovacia látka proti chrípke (štiepený virión, inaktivovaná)	DE/H/1939/001	59/0114/14-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Fluarix Tetra Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/1939/001/DC	235552	GLAXOSMITHKLINE PHARMA GMBH.	AT
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132012	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split	DE/H/1939/001	043132024	GLAXOSMITHKLINE BIOLOGICALS 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
(frammentato), inattivato)				
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132036	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132048	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132051	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132063	GLAXOSMITHKLINE BIOLOGICALS 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
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132075	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra suspensión inyectable en jeringa precargada Vacuna antigripal (de virus fraccionados e inactivados)	DE/H/1939/001	78.568	GLAXOSMITHKLINE BIOLOGICALS S.A.	ES
Fluarix Tetra, injekcní suspenze v predplnené injekcní stríkacce Vakcína proti chripce (štepený virion, inaktivovaný)	DE/H/1939/001	59/145/14-C	GLAXOSMITHKLINE BIOLOGICALS S.A.	CZ
Fluarix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Αντιγριπικό εμβόλιο (τμήμα ιού, αδρανοποιημένο)	DE/H/1939/001	38342/1-9-2015	GLAXOSMITHKLINE AEBE	GR
Fluarix, injeksjonsvæske, suspensjon, i ferdigfylt sprøyte. Vaksine mot influensa (inaktivert, splittvirus).	DE/H/0124/001	02-1030	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
Fluarix, injektionsvæske, suspension, fyldt injektionssprøjte	DE/H/0124/001	15746	GLAXOSMITHKLINE PHARMA A/S	DK
Fluarix, injektionsvätska, suspension i förfylld spruta Influensavaccin (spjälkat virus, inaktiverat)	DE/H/124/01	14056	GLAXOSMITHKLINE AB	SE
Fluarix, injektionsvätska, suspension i förfylld spruta Vaccin mot influensa (spjälkat virus, inaktiverat)	DE/H/0124/001	13153	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
FLUARIX, sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati).	DE/H/124/01	29245230	GLAXOSMITHKLINE S.P.A.	IT
FLUARIX, sospensione iniettabile in siringa preriempita. Vaccino influenzale (virus frammentati, inattivati).	DE/H/124/01	029245267	GLAXOSMITHKLINE S.P.A.	IT
Fluarix, stungulyf, dreifa í áfylltum sprautum Inflúensubóluefni (veiruhlutar, deyddir)	DE/H/0124/001	930249 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Fluarix, suspensão injectável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inactivado)	DE/H/0124/001	2454684	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix, suspensão injectável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inactivado)	DE/H/0124/001	2704880	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix, suspensão injectável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inactivado)	DE/H/0124/001	5304670	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/0124/001	2943587	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix, süstesuspensioon süstlis. Gripivaktsiin (inaktiveeritud purustatud viirus)	DE/H/0124/001	160597	GLAXOSMITHKLINE BIOLOGICALS S.A.	EE
Fluarix, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw grypie (rozszczepiony wirion,	DE/H/0124/001	12831	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
inaktywowana)				
FLUARIX, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο γρίπης (τμήμα ιού, αδρανοποιημένο)	DE/H/124/01	20302	GLAXOSMITHKLINE (CYPRUS) LIMITED	CY
Fluarix, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο γρίπης (τμήμα ιού, αδρανοποιημένο)	DE/H/0124/001	38341/1-9-2015	GLAXOSMITHKLINE AEBE	GR
Fluarix® Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/0124/001	2-00382	GLAXOSMITHKLINE PHARMA GMBH.	AT
Fluarix® suspension for injection in a pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/0124/001	PL 10592/0118	SMITHKLINE BEECHAM LTD	UK
Fluarix® Tetraqsuspension for injection in pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1939/001/DC	PL 10592/0302	SMITHKLINE BEECHAM 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
FLUARIXTETRA™ suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	DE/H/1939/001	NL42097	LABORATOIRE GLAXOSMITHKLINE	FR
IMMUGRIP, suspension injectable en seringue préremplie	not available	333 855-0	PIERRE FABRE MEDICAMENT	FR
INACTIVATED INFLUENZA VACCINE (SPLIT VIRION) BP, suspension for injection in prefilled syringe. Influenza vaccine (split virion, inactivated)	FR/H/0121/001	PA 2131/012/001	SANOFI PASTEUR EUROPE	IE
INACTIVATED INFLUENZA VACCINE (SPLIT VIRION) BP, suspension for injection in prefilled syringe. Influenza vaccine (split virion, inactivated)	FR/H/0121/001	PL 46602/0002	SANOFI PASTEUR EUROPE	UK
Influsplit SSW® 2014/2015 Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/124/01	PEI.H.11676.01.1	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Influsplit SSW® 2014/2015 Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/124/01	PEI.H.00084.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE
Influsplit SSW® 2014/2015|Injektionssuspension in einer Fertigspritze|Influenza-Spaltimpfstoff (inaktiviert)	DE/H/124/01	PEI.H.00084.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE
Influsplit Tetra® 2014/2015 Injektionssuspension in Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/1939/001	PEI.H.11629.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	8650309	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	2638286	SANOFI PASTEUR EUROPE	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
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	2638385	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	4317186	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	4316980	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	4317285	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	4317087	SANOFI PASTEUR EUROPE	PT
ISTIVAC, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião	FR/H/0122/001	2638484	SANOFI PASTEUR EUROPE	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
fragmentado, inativado)				
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 298-6	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	320 153-2	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 299-2	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 300-0	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 301-7	SANOFI PASTEUR EUROPE	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
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 302-3	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	352 304-6	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	FR/H/0122/001	320 152-6	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspensión inyectable en jeringa precargada. Vacuna antigripal (virus fraccionados, inactivados)	FR/H/0122/001	51.577	SANOFI PASTEUR EUROPE	ES
Quadrivalent Influenza Vaccine (split virion, inactivated), suspension for injection in pre-filled syringe Quadrivalent influenza vaccine (split virion, inactivated)	DE/H/1949/001	PA 2131/013/001	SANOFI PASTEUR EUROPE	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
Quadrivalent Influenza Vaccine (split virion, inactivated), suspension for injection in pre-filled syringe Quadrivalent influenza vaccine (split virion, inactivated)	DE/H/1949/001	PL 46602/0017	SANOFI PASTEUR EUROPE	UK
VAXIGRIP Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	FR/H/0121/001	PEI.H.00189.01.1	SANOFI PASTEUR EUROPE	DE
VAXIGRIP ENFANTS, suspension injectable en seringue préremplie|Vaccin grippal (inactivé, à virion fragmenté)	FR/H/0139/001	NL 22 390	SANOFI PASTEUR SA	FR
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/002	SANOFI PASTEUR SA	LT
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/003	SANOFI PASTEUR SA	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
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/004	SANOFI PASTEUR SA	LT
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/005	SANOFI PASTEUR SA	LT
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/006	SANOFI PASTEUR SA	LT
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/007	SANOFI PASTEUR SA	LT
VAXIGRIP injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	FR/H/0121/001	LT/1/07/0789/008	SANOFI PASTEUR SA	LT
VAXIGRIP injekcine suspensija užpildytame švirkšte, Vakcina nuo gripo (iš virionu	FR/H/121/01	N1_LT/1/07/0789/001	SANOFI PASTEUR SA	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
fragmentu, inaktyvuota)				
VAXIGRIP Injekcná suspenzia naplnenej v injekcnej striekacke Ockovacia látka proti chrípke (štiepený virión, inaktivovaná)	FR/H/0121/001	59/0228/07-S	SANOFI PASTEUR SA	SK
VAXIGRIP Injektionssuspension in einer Fertigspritze Influenza Impfstoff (Spaltimpfstoff, inaktiviert)	FR/H/0121/001	2-00215	SANOFI PASTEUR EUROPE	AT
VAXIGRIP Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	FR/H/0121/001	BE108184	SANOFI PASTEUR EUROPE	BE
VAXIGRIP Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	FR/H/0121/001	2009010131	SANOFI PASTEUR EUROPE	LU
VAXIGRIP suspensija injekcijam pilnšlirce, Gripas vakcina (škelts virions, inaktiveta),	FR/H/121/01	07-0162	SANOFI PASTEUR SA	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
Vaccinum influenzae inactivatum ex virorum fragmentis praeparatum				
VAXIGRIP suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano Sevi 2017/2018	not available	H/97/01621/001	SANOFI PASTEUR SA	SI
Vaxigrip szuszpenziós injekció eloretöltött fecskendoben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/12	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/02	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/03	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/04	SANOFI PASTEUR SA	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
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/05	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/06	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/07	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/08	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/09	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus,	FR/H/0121/001	OGYI-T-8606/10	SANOFI PASTEUR SA	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
inaktivált)				
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/11	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/13	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/14	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/15	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/16	SANOFI PASTEUR SA	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
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/17	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/18	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/19	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/20	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/21	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus,	FR/H/0121/001	OGYI-T-8606/22	SANOFI PASTEUR SA	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
inaktivált)				
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/23	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/24	SANOFI PASTEUR SA	HU
Vaxigrip szuszpenziós injekció eloretöltött fecskendoben, Influenza vakcina (split virion, inaktivált)	FR/H/121/01	OGYI-T-8606/01	SANOFI PASTEUR SA	HU
Vaxigrip Tetra Injektionssuspension in einer Fertigspritze Tetravalenter Influenza-Spaltimpfstoff (inaktiviert)	DE/H/1949/001	PEI.H.11808.01.1	SANOFI PASTEUR EUROPE	DE
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus,	DE/H/1949/001	OGYI-T-23068/01	SANOFI PASTEUR SA	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
inaktivált)				
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/02	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/03	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/04	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/05	SANOFI PASTEUR SA	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
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/06	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/07	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/08	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/09	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus,	DE/H/1949/001	OGYI-T-23068/10	SANOFI PASTEUR SA	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
inaktivált)				
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/11	SANOFI PASTEUR SA	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/12	SANOFI PASTEUR SA	HU
Vaxigrip Tetra, injekcnà suspenzia naplnenà v injekcnej, striekacke tetravalentnà ockovacia, làtka, proti, chripke (stiepeny virion inaktivovanà)	DE/H/1949/001	59/0415/16-S	SANOFI PASTEUR SA	SK
Vaxigrip Tetra, injekcní suspenze v predplnené injekcní stríkacce, Tetravalentní vakcína proti chripce (štepený virion, inaktivovaný)	DE/H/1949/001	59/370/16-C	SANOFI PASTEUR SA	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
Vaxigrip Tetra, Injektionssuspension in einer Fertigspritze Quadrivalenter Influenza-Impfstoff (inaktiviert, gespalten)	DE/H/1949/001	BE501511	SANOFI PASTEUR EUROPE	BE
Vaxigrip Tetra, Injektionssuspension in einer Fertigspritze Quadrivalenter Influenza-Impfstoff (inaktiviert, gespalten)	DE/H/1949/001	2017040149	SANOFI PASTEUR EUROPE	LU
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita, Vaccino influenzale quadrivalente preparato con virus frammentati "split", inattivati	DE/H/1949/001	044898017	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita, Vaccino influenzale quadrivalente preparato con virus frammentati "split", inattivati	DE/H/1949/001	044898029	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita,	DE/H/1949/001	044898031	SANOFI PASTEUR EUROPE	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
Vaccino influenzale quadrivalente preparato con virus frammentati “split”, inattivati				
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita, Vaccino influenzale quadrivalente preparato con virus frammentati “split”, inattivati	DE/H/1949/001	044898043	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita, Vaccino influenzale quadrivalente preparato con virus frammentati “split”, inattivati	DE/H/1949/001	044898056	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita, Vaccino influenzale quadrivalente preparato con virus frammentati “split”, inattivati	DE/H/1949/001	044898068	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a	DE/H/1949/001	5686852	SANOFI PASTEUR EUROPE	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
gripe (viriao fragmentado, inativado)				
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a gripe (viriao fragmentado, inativado)	DE/H/1949/001	5686928	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a gripe (viriao fragmentado, inativado)	DE/H/1949/001	5686878	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a gripe (viriao fragmentado, inativado)	DE/H/1949/001	5686860	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a gripe (viriao fragmentado, inativado)	DE/H/1949/001	5686910	SANOFI PASTEUR EUROPE	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
Vaxigrip Tetra, suspensao injetavel em seringa pre-cheia, Vacina quadrivalente contra a gripe (viriao fragmentado, inativado)	DE/H/1949/001	5686902	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensie voor injectie in een voorgevulde spuit Quadrivalent griepvaccin (gesplitst virion, geïnactiveerd)	DE/H/1949/001	BE501511	SANOFI PASTEUR EUROPE	BE
Vaxigrip Tetra, suspension injectable en seringue préremplie Vaccin grippal quadrivalent (virion fragmenté, inactivé)	DE/H/1949/001	BE501511	SANOFI PASTEUR EUROPE	BE
Vaxigrip Tetra, suspension injectable en seringue préremplie Vaccin grippal quadrivalent (virion fragmenté, inactivé)	DE/H/1949/001	2017040149	SANOFI PASTEUR EUROPE	LU
Vaxigrip Tetra, suspension inyectable en jeringa precargada Vacuna antigripal tetravalente (virus	DE/H/1949/001	81.098	SANOFI PASTEUR EUROPE	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
fraccionados, inactivados)				
VAXIGRIP, injekcní suspenze v predplnené injekcní stríkacce, Vakcína proti chripce (inaktivovaná, štepený virion)	FR/H/121/01	59/1035/94-C	SANOFI PASTEUR SA	CZ
VAXIGRIP, injeksjonsvæske, suspensjon i ferdigfylt sprøyte. Vaksine mot influensa (inaktivert, splittvirus).	FR/H/0121/001	07-4931	SANOFI PASTEUR EUROPE	NO
Vaxigrip, injektioneste, suspensio, esitäytetyssä ruiskussa. Influenssarokote (virusfragmentit, inaktivoitu)	FR/H/0121/001	13130	SANOFI PASTEUR EUROPE	FI
Vaxigrip, injektionsvæske, suspension, fyldt injektionssprøjte	FR/H/0121/001	11708	SANOFI PASTEUR EUROPE	DK
Vaxigrip, injektionsvätska, suspension i förfylld spruta. Influensavaccin	FR/H/0121/001	13130	SANOFI PASTEUR EUROPE	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
(spjälkat virus, inaktiverat)				
VAXIGRIP, injektionsvätska, suspension i förfylld spruta. Influensavaccin (spjälkat virus, inaktiverat)	FR/H/0121/001	14029	SANOFI PASTEUR EUROPE	SE
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032209	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032375	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032274	SANOFI PASTEUR EUROPE	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
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032387	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032286	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032399	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032298	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”,	FR/H/0121/001	026032401	SANOFI PASTEUR EUROPE	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
inattivati				
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032300	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032312	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032324	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati “split”, inattivati	FR/H/0121/001	026032336	SANOFI PASTEUR EUROPE	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
VAXIGRIP, stungulyf, dreifa í áfylltri sprautu. Inflúensubóluefni (klofin veiruögn, deydd).	FR/H/0121/001	IS/1/07/042/01	SANOFI PASTEUR EUROPE	IS
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	2637882	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	2637981	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	2638088	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	4317780	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	4317681	SANOFI PASTEUR EUROPE	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
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	2638187	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	4317988	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0121/001	4317889	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensie voor injectie in een voorgevulde spuit. Griepvaccin (gesplitst virion, geïnactiveerd).	FR/H/0121/001	BE108184	SANOFI PASTEUR EUROPE	BE
VAXIGRIP, suspensie voor injectie in een voorgevulde spuit. Griepvaccin (gesplitst virion, geïnactiveerd)	FR/H/0121/001	RVG 22306	SANOFI PASTEUR EUROPE	NL
VAXIGRIP, suspension injectable en flacon multidose, Vaccin grippal inactivé à virion	FR/H/0121/002	NL 11 155-2	SANOFI PASTEUR SA	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
fragmenté				
VAXIGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé à virion fragmenté)	FR/H/0121/001	2009010131	SANOFI PASTEUR EUROPE	LU
VAXIGRIP, suspension injectable en seringue préremplie, VACCIN GRIPPAL INACTIVE A VIRION FRAGMENTE	FR/H/121/01	NL 11 155-1	SANOFI PASTEUR SA	FR
VAXIGRIP, suspension injectable en seringue préremplie. Vaccin grippal (inactivé à virion fragmenté).	FR/H/0121/001	BE108184	SANOFI PASTEUR EUROPE	BE
VAXIGRIP, suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	FR/H/0121/001	61.108	SANOFI PASTEUR EUROPE	ES
VAXIGRIP, süstesuspensioon süstlis (inaktiveeritud gripi vaktsiin, purustatud viirus)	FR/H/0121/001	186597	SANOFI PASTEUR SA	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
VAXIGRIP, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο)	FR/H/0121/001	68726/10-11-2016	SANOFI PASTEUR EUROPE	GR
VAXIGRIP, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα, Αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο).	FR/H/0121/001	20507	SANOFI PASTEUR SA	CY
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/001	SANOFI PASTEUR SA	LT
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/002	SANOFI PASTEUR SA	LT
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/003	SANOFI PASTEUR SA	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
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/004	SANOFI PASTEUR SA	LT
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/005	SANOFI PASTEUR SA	LT
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/006	SANOFI PASTEUR SA	LT
VaxigripTetra Injektionssuspension in einer Fertigspritze Tetravalenter Influenza-Impfstoff (Spaltvirus, inaktiviert)	DE/H/1949/001	137203	SANOFI PASTEUR EUROPE	AT
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat,	DE/H/1949/001	9198/2016/01	SANOFI PASTEUR SA	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
inactivat)				
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat, inactivat)	DE/H/1949/001	9198/2016/02	SANOFI PASTEUR SA	RO
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat, inactivat)	DE/H/1949/001	9198/2016/03	SANOFI PASTEUR SA	RO
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat, inactivat)	DE/H/1949/001	9198/2016/04	SANOFI PASTEUR SA	RO
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat, inactivat)	DE/H/1949/001	9198/2016/05	SANOFI PASTEUR SA	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
VaxigripTetra suspensie inectabilà în seringà preumpluta Vaccin antigripal tetravalent (virion fragmentat, inactivat)	DE/H/1949/001	9198/2016/06	SANOFI PASTEUR SA	RO
VaxigripTetra suspensija injekcijam pilnšlirce Cetrvertiga gripas vakcina (škelts virions, inaktiveta) Vaccinum influenzae inactivatum ex virorum fragmentis praeparatum	DE/H/1949/001	16-0141	SANOFI PASTEUR SA	LV
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/001	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/002	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi	DE/H/1949/001	H/16/02251/003	SANOFI PASTEUR SA	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
Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)				
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/004	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/005	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/006	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/007	SANOFI PASTEUR SA	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
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/008	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/009	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/010	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/011	SANOFI PASTEUR SA	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni,	DE/H/1949/001	H/16/02251/012	SANOFI PASTEUR SA	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
inaktivirano)				
Vaxigriptetra, injeksjonsvæske, suspensjon i ferdigfylt sprøyte. Vaksine mot influensa (inaktivert, splittvirus).	DE/H/1949/001	15-10871	SANOFI PASTEUR EUROPE	NO
VaxigripTetra, injektioneste, suspensio, esitäytetty ruisku. Nelivalenttinen influenssarokote (virusfragmentit, inaktivoitu)	DE/H/1949/001	33654	SANOFI PASTEUR EUROPE	FI
Vaxigriptetra, injektionsvæske, suspension i fyldt injektionssprøjte	DE/H/1949/001	56583	SANOFI PASTEUR EUROPE	DK
VaxigripTetra, injektionsvätska, suspension i förfylld spruta Quadrivalent influensavaccin (spjälkat virus, inaktiverat)	DE/H/1949/001	33654	SANOFI PASTEUR EUROPE	FI
VaxigripTetra, injektionsvätska, suspension i förfylld	DE/H/1949/001	53400	SANOFI PASTEUR EUROPE	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
spruta Quadrivalent influensavaccin (spjälkat virus, inaktiverat)				
VaxigripTetra, stungulyf, dreifa í áfylltri sprautu. Fjórgilt inflúensubóluefni (klofin veiruögn, deydd)	DE/H/1949/001	IS/1/16/068/01	SANOFI PASTEUR EUROPE	IS
VaxigripTetra, suspensie voor injectie Quadrivalent griepvaccin (gesplitst virion, geïnactiveerd)	DE/H/1949/001	RVG 117963	SANOFI PASTEUR EUROPE	NL
VaxigripTetra, suspension for injection in pre-filled syringe	DE/H/1949/001	MA573/00103	SANOFI PASTEUR SA	MT
VaxigripTetra, suspension injectable en seringue préremplie, Vaccin grippal quadrivalent (inactivé, à virion fragmenté)	DE/H/1949/001	NL 46 320	SANOFI PASTEUR SA	FR
VaxigripTetra, suspenzija za injekciju u napunjenoj štrcaljki Cetverovalentno cjepivo protiv influence (fragmentirani virion, inaktivirano)	DE/H/1949/001	HR-H-888872987	SANOFI PASTEUR SA	HR
VaxigripTetra, süstesuspensioon süstlis,	DE/H/1949/001	919416	SANOFI PASTEUR SA	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
Neljavalentne gripivaktsiin (purustatud viirus, inaktiveeritud)				
VaxigripTetra, zawiesina do wstrzykiwan w ampulko-strzykawce Czterowalentna szczepionka przeciw grypie (rozszczepiony wirion), inaktywowana	DE/H/1949/001	23540	SANOFI PASTEUR SA	PL
VaxigripTetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Τετραδύναμο αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο)	DE/H/1949/001	022512	SANOFI PASTEUR SA	CY
VaxigripTetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Τετραδύναμο αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο)	DE/H/1949/001	57068/15/16-12-2016	SANOFI PASTEUR EUROPE	GR
α-RIX-Tetra, suspension injectable en seringue préremplie Vaccin antigrippal (virion fragmenté, inactivé)	DE/H/1939/001/DC	BE456924	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
α-RIX-Tetra, suspension injectable en seringue préremplie Vaccin antigrippal (virion fragmenté, inactivé)	DE/H/1939/001	2014090216	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
α-RIX-Tetra, suspension injectable en seringue préremplie Vaccin antigrippal (virion fragmenté, inactivé)	DE/H/1939/001	2014090216	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
ВАКСИГРИП 0,5 ml инжекционна суспензия, Ваксина срещу грип (фрагментиран инактивиран вирион), Инжекционна суспензия в предварително напълнена спринцовка	FR/H/121/01	20000248	SANOFI PASTEUR SA	BG
ВаксигрипТетра инжекционна суспензия в предварително напълнена спринцовка Четирилентна ваксина срещу грип (фрагментиран инактивиран вирион)	DE/H/1949/001	20160310	SANOFI PASTEUR SA	BG
Флуарикс, инжекционна суспензия в предварително	DE/H/0124/001	9700046	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
напълнена спринцовка Ваксина срещу грип (фрагментиран инактивиран вирион)				