



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

31 October 2019
EMA/600419/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

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

Procedure no.: PSUSA/00010298/201903

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Afluria Injeksjonsvæske, suspensjon, i ferdigfylt sprøyte Vaksine mot influensa (inaktivert, splittvirus)	DE/H/1938/001	04-3084	SEQIRUS GMBH	NO
Afluria injektionsvätska, suspension i förfylld spruta Vaccin mot influensa (spjälkat virus inaktiverat)	DE/H/1938/001	20546	SEQIRUS GMBH	SE
Afluria Injektionsvätska, suspension i förfylld spruta. Vaccin mot influensa (spjälkat virus, inaktiverat)	DE/H/1938/001	20234	SEQIRUS GMBH	FI
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1938/001	043216023	SEQIRUS GMBH	IT
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1938/001	043216035	SEQIRUS GMBH	IT
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1938/001	043216047	SEQIRUS GMBH	IT
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1938/001	043216050	SEQIRUS GMBH	IT
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino	DE/H/1938/001	043216062	SEQIRUS GMBH	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
influenzale (virus split (frammentato), inattivato)				
AFLURIA Sospensione iniettabile, in siringa pre-riempita. Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1938/001	043216011	SEQIRUS GMBH	IT
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	5643317	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 Suspensão injetável, em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/1938/001	5643309	SEQIRUS GMBH	PT
AFLURIA suspensie injectabilă, în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/01	SEQIRUS GMBH	RO

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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, 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	7123/2014/04	SEQIRUS GMBH	RO
AFLURIA suspensie injectabilă, în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/05	SEQIRUS GMBH	RO
AFLURIA suspensie injectabilă, în seringă preumplută Vaccin gripal (virion fragmentat, inactivat)	DE/H/1938/001	7123/2014/06	SEQIRUS GMBH	RO
AFLURIA Suspensie voor injectie, in een voorgevulde spuit. Influenzavaccin (split virus, geïnactiveerd)	DE/H/1938/001	BE285284	SEQIRUS GMBH	BE
AFLURIA Suspensie voor injectie, in een voorgevulde spuit. Influenzavaccin (split virus, geïnactiveerd)	DE/H/1938/001	RVG 31924	SEQIRUS GMBH	NL
AFLURIA Suspension injectable dans une	DE/H/1938/001	1716/06040039	SEQIRUS GMBH	LU

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seringue préremplie. Vaccin contre l'influenza (virion fragmenté, inactif)				
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
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	702573 – 3	SEQIRUS GMBH	ES
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	715322 - 1	SEQIRUS GMBH	ES
Afluria Suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	DE/H/1938/001	715321 - 4	SEQIRUS GMBH	ES
AFLURIA Suspension zur Injektion in Fertigspritzen. Influenza-Impfstoff	DE/H/1938/001	BE285284	SEQIRUS GMBH	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
(Spaltvirus, inaktiviert)				
AFLURIA Suspension zur Injektion in Fertigspritzen. Influenza-Impfstoff (Spaltvirus, inaktiviert)	DE/H/1938/001	1716/06040039	SEQIRUS GMBH	LU
AFLURIA Ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο γρίπης (τμήμα ιού, αδραντοποιημένο)	DE/H/1938/001	21056	SEQIRUS GMBH	GR
AFLURIA, injekční suspenze v predplněné injekční stříkacce Vakcína proti chřipce (štepový virion, inaktivovaná)	DE/H/1938/001	59/241/14-C	SEQIRUS GMBH	CZ
AFLURIA, injektioneste, suspensio, esitäytetty ruisku. Influenssarokote (pilkottu, inaktivoitu)	DE/H/1938/001	20234	SEQIRUS GMBH	FI
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	BE285284	SEQIRUS GMBH	BE
AFLURIA®, Suspension zur Injektion in Fertigspritzen Influenza-Impfstoff (Spaltvirus, inaktiviert)	DE/H/1938/001	PEI.H.03523.01.1	SEQIRUS GMBH	DE
a-RIX Injektionssuspension in einer Fertigspritze Grippeimpfung	DE/H/0124/001	BE147244	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE

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(fragmentiertes, inaktiviertes Virion)				
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
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
a-RIX-Tetra, suspension injectable en seringue préremplie Vaccin antigrippal (virion fragmenté, inactivé)	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
Fluarix 2016/2017,	DE/H/0124/001	RVG 22307	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
suspensie voor injectie in een voorgevulde spuit Influenzavaccin (gesplitst virion, geïnactiveerd)				
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 suspensión inyectable en jeringa precargada Vacuna antigripal (de virus fraccionados e inactivados)	DE/H/0124/001	60.772	GLAXOSMITHKLINE, S.A.	ES
Fluarix Tetra injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1939/001	LT/1/18/4236/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix Tetra injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1939/001	LT/1/18/4236/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix Tetra injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1939/001	LT/1/18/4236/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix Tetra injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1939/001	LT/1/18/4236/004	GLAXOSMITHKLINE LIETUVA UAB	LT
Fluarix Tetra injekcinė suspensija užpildytame švirkšte Vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1939/001	LT/1/18/4236/005	GLAXOSMITHKLINE LIETUVA UAB	LT

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inaktivuota)				
Fluarix Tetra injekčná suspenzia naplnená v injekčnej striekacke Ockovacia látka proti chrípke (štiepený virión, inaktivovaná)	DE/H/1939/001	59/0114/14-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Fluarix Tetra injekční suspenze v předplněné injekční stříkačce Vakcína proti chrípce (štěpený virion, inaktivovaný)	DE/H/1939/001	59/145/14-C	GLAXOSMITHKLINE BIOLOGICALS S.A.	CZ
Fluarix Tetra injektioneste, suspensio, esitäytetyssä ruiskussa Influenzavakcine (virusfragmentit, inaktivoitu)	DE/H/1939/001	35880	GLAXOSMITHKLINE OY	FI
Fluarix Tetra Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/1939/001	235552	GLAXOSMITHKLINE PHARMA GMBH.	AT
Fluarix Tetra sospensione iniettabile in siringa preimpiata Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1939/001	043132036	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preimpiata Vaccino influenzale (virus split frammentato), inattivato)	DE/H/1939/001	043132048	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
Fluarix Tetra sospensione iniettabile in siringa preimpiata Vaccino influenzale (virus split	DE/H/1939/001	043132051	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT

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(frammentato), inattivato)				
Fluarix Tetra sospensione iniettabile in siringa preriempita Vaccino influenzale (virus split (frammentato), inattivato)	DE/H/1939/001	043132063	GLAXOSMITHKLINE BIOLOGICALS S.A.	IT
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 suspensija injekcijam pilnšlirce Gripas vakcina (škelts, inaktivets virions) Vaccinum influenzae inactivatum ex virorum fragmentis praeparatum	DE/H/1939/001	18-0098	GLAXOSMITHKLINE LATVIA SIA	LV
Fluarix Tetra suspension for injection in pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1939/001	PA 1077/134/001	SMITHKLINE BEECHAM LTD	IE
Fluarix Tetra suspension for injection in pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1939/001	MA170/01101	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
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 suspenzija za injekciju u napunjenoj štrcaljki, cjepivo protiv influence (fragmentirani virion), inaktivirano	DE/H/1939/001	HR-H-048213854	GLAXOSMITHKLINE D.O.O.	HR

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Fluarix Tetra suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/1939/001	H/18/02481/002	GLAXOSMITHKLINE D.O.O.	SI
Fluarix Tetra suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/1939/001	H/18/02481/003	GLAXOSMITHKLINE D.O.O.	SI
Fluarix Tetra suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/1939/001	H/18/02481/004	GLAXOSMITHKLINE D.O.O.	SI
Fluarix Tetra suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/1939/001	H/18/02481/005	GLAXOSMITHKLINE D.O.O.	SI
Fluarix Tetra suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti gripi z delci virionov, inaktivirano	DE/H/1939/001	H/18/02481/001	GLAXOSMITHKLINE D.O.O.	SI
Fluarix Tetra süstesuspensioon süstlis Gripivaktsiin (inaktiveeritud purustatud virus)	DE/H/1939/001	966018	GLAXOSMITHKLINE EESTI OÜ	EE
Fluarix Tetra szuszpenziós injekció eloretöltött fecskendoben Influenza vakcina (split-vírus, inaktívált)	DE/H/1939/001	OGYI-T-23435/01	GLAXOSMITHKLINE KFT.	HU
Fluarix Tetra szuszpenziós injekció eloretöltött	DE/H/1939/001	OGYI-T-23435/02	GLAXOSMITHKLINE KFT.	HU

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fecskendoben Influenza vakcina (split-virus, inaktivált)				
Fluarix Tetra szuszpenziós injekció eloretöltött fecskendoben Influenza vakcina (split-virus, inaktivált)	DE/H/1939/001	OGYI-T-23435/04	GLAXOSMITHKLINE KFT.	HU
Fluarix Tetra szuszpenziós injekció eloretöltött fecskendoben Influenza vakcina (split-virus, inaktivált)	DE/H/1939/001	OGYI-T-23435/05	GLAXOSMITHKLINE KFT.	HU
Fluarix Tetra, injektionsvæske, suspension i fyldt injektionsprøjte	DE/H/1939/001	60754	GLAXOSMITHKLINE PHARMA A/S	DK
Fluarix Tetra, injektionsvätska, suspension i förfylld spruta Influenzavaccin (spjälkat virus, inaktiverat)	DE/H/1939/001	57495	GLAXOSMITHKLINE AB	SE
Fluarix Tetra, stungulyf, dreifa í áfylltri sprautu Inflúensubóluefni (veiruhlutar, deyddir)	DE/H/1939/001	IS/1/18/068/01	GLAXOSMITHKLINE PHARMA A/S	IS
Fluarix Tetra, suspensão injetável em seringa pré-cheia Vacina contra a gripe (vírião fragmentado, inativado)	DE/H/1939/001	5751078	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Fluarix Tetra, suspensie voor injectie in een voorgevulde spuit Influenzavaccin (gesplitst virion, geïnactiveerd)	DE/H/1939/001	RVG 122632	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
Fluarix Tetra, zawiesina do wstrzykiwan w ampulko-strzykawce Szczepionka przeciw grypie (rozszczepiony wirion, inaktywowana)	DE/H/1939/001	24888	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Fluarix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Αντιγριπικό εμβόλιο (τμήμα ιού, αδρανοποιημένο)	DE/H/1939/001	022907	GLAXOSMITHKLINE (CYPRUS) LIMITED	CY
Fluarix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Αντιγριπικό εμβόλιο (τμήμα ιού, αδρανοποιημένο)	DE/H/1939/001	3062501	GLAXOSMITHKLINE BIOLOGICALS S.A.	GR
Fluarix Tetraqsuspension for injection in pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1939/001	PL 10592/0302	SMITHKLINE BEECHAM LTD	UK
Fluarix, stungulyf, dreifa í áfylltum sprautum Inflúensubóluefni (veiruhlutur, deyddir)	DE/H/0124/001	930249	GLAXOSMITHKLINE PHARMA A/S	IS
Fluarix, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/0124/001	2454684	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	2704880	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix, suspensão injetável em seringa pré-	DE/H/0124/001	2943587	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS	PT

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cheia Vacina contra a gripe (virião fragmentado, inativado)			FARMACEUTICOS LDA	
Fluarix, suspensão injetável em seringa pré-cheia Vacina contra a gripe (virião fragmentado, inativado)	DE/H/0124/001	5304670	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Fluarix® suspension for injection in a pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/0124/001	MA 172/01001	SMITHKLINE BEECHAM LTD	MT
Fluarix® Tetraqsuspension for injection in pre-filled syringe Influenza vaccine (split virion, inactivated)	DE/H/1939/001	18-12139	GLAXOSMITHKLINE AS	NO
FLUARIXTETRA suspension injectable en seringue préremplie Vaccin grippal inactivé à virion fragmenté	DE/H/1939/001	NL42097	LABORATOIRE GLAXOSMITHKLINE	FR
Influsplit SSW® 2014/2015 Injektionssuspension in einer Fertigspritze Influenza-Spaltimpfstoff (inaktiviert)	DE/H/0124/001	PEI.H.11676.01.1	GLAXOSMITHKLINE BIOLOGICALS S.A.	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

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ISTIVAC, suspensão injetável em seringa pré-cheia. Vacina contra a gripe (virião fragmentado, inativado)	FR/H/0122/001	2638286	SANOFI PASTEUR EUROPE	PT
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 fragmentado, inativado)	FR/H/0122/001	2638484	SANOFI PASTEUR EUROPE	PT
MUTAGRIP, suspension injectable en seringue	FR/H/122/01	352 298-6	SANOFI PASTEUR EUROPE	FR

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préremplie Vaccin grippal (inactivé, à virion fragmenté)				
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	320 153-2	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	352 299-2	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	352 300-0	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	352 301-7	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	352 302-3	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	352 304-6	SANOFI PASTEUR EUROPE	FR
MUTAGRIP, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/122/01	320 152-6	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
fragmenté)				
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
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
Trivalent Influenza Vaccine (Split Virion, Inactivated) High Dose, suspension for injection in pre-filled syringe	not available	PL 04425/0756	AVENTIS PHARMA LTD	UK
Trivalent Influenza Vaccine (Split Virion, Inactivated) High Dose, suspension for injection in pre-filled syringe	not available	PL 04425/0756	AVENTIS PHARMA LTD	UK
VAXIGRIP ENFANTS, suspension injectable en seringue préremplie Vaccin grippal (inactivé, à virion fragmenté)	FR/H/0139/001	NL 22 390	SANOFI PASTEUR	FR
Vaxigrip injekční suspenze v předplněné injekční stříkačce Vakcína proti chřipce (inaktivovaná, štěpený virion)	FR/H/0121/001	59/1035/94-C	SANOFI PASTEUR	CZ
Vaxigrip injektionsvätska, suspension, förfylld spruta	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
influenzavaccin (spjalkat virus, inaktiverat)				
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/02	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/03	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/04	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/05	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/06	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/07	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/08	SANOFI PASTEUR	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-8606/09	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/10	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/11	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/12	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/13	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/14	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/15	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött	FR/H/0121/001	OGYI-T-8606/16	SANOFI PASTEUR	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
fecskendőben Influenza vakcina (hasított vírus, inaktivált)				
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/17	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/18	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/19	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/20	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/21	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus, inaktivált)	FR/H/0121/001	OGYI-T-8606/22	SANOFI PASTEUR	HU
Vaxigrip szuszpenziós injekció előretöltött fecskendőben Influenza vakcina (hasított vírus,	FR/H/0121/001	OGYI-T-8606/23	SANOFI PASTEUR	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-8606/24	SANOFI PASTEUR	HU
Vaxigrip Tetra injekčná suspenzia naplnená v injekčnej striekačke tetravalentná očkovacia látka proti chrípke (štiepený virión, inaktivovaná)	DE/H/1949/001	59/0415/16-S	SANOFI PASTEUR	SK
Vaxigrip Tetra injekční suspenze v předplněné injekční stříkačce Tetravalentní vakcína proti chřipce (štěpený virion, inaktivovaný)	DE/H/1949/001	59/370/16-C	SANOFI PASTEUR	CZ
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, inaktivált)	DE/H/1949/001	OGYI-T-23068/01	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/02	SANOFI PASTEUR	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-23068/03	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/04	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/05	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/06	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/08	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/09	SANOFI PASTEUR	HU
Vaxigrip Tetra	DE/H/1949/001	OGYI-T-23068/10	SANOFI PASTEUR	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
szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)				
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/11	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/12	SANOFI PASTEUR	HU
Vaxigrip Tetra szuszpenziós injekció előretöltött fecskendőben kvadrivalens influenza vakcina (hasított vírus, inaktivált)	DE/H/1949/001	OGYI-T-23068/07	SANOFI PASTEUR	HU
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	DE/H/1949/001	044898017	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
siringa preriempita 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	044898029	SANOFI PASTEUR EUROPE	IT
Vaxigrip Tetra, sospensione iniettabile in siringa preriempita Vaccino influenzale quadrivalente preparato con virus frammentati "split", inattivati	DE/H/1949/001	044898031	SANOFI PASTEUR EUROPE	IT
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	DE/H/1949/001	044898068	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
preparato con virus frammentati "split", inattivati				
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente contra a gripe (virião fragmentado, inativado)	DE/H/1949/001	5686860	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente contra a gripe (virião fragmentado, inativado).	DE/H/1949/001	5686852	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente contra a gripe (virião fragmentado, inativado).	DE/H/1949/001	5686878	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente contra a gripe (virião fragmentado, inativado).	DE/H/1949/001	5686928	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente contra a gripe (virião fragmentado, inativado).	DE/H/1949/001	5686910	SANOFI PASTEUR EUROPE	PT
Vaxigrip Tetra, suspensão injetável em seringa pré-cheia. Vacina quadrivalente	DE/H/1949/001	5686902	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
contra a gripe (virião fragmentado, inativado).				
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, suspensie voor injectie in een voorgevulde spuit Quadrivalent griepvaccin (gesplitst virion, geïnactiveerd)	DE/H/1949/001	RVG 117963	SANOFI PASTEUR EUROPE	NL
Vaxigrip Tetra, suspension for injection in pre-filled syringe Quadrivalent influenza vaccine (split virion, inactivated)	DE/H/1949/001	MA573/00103	SANOFI PASTEUR	MT
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 fraccionados, inactivados)	DE/H/1949/001	81098	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
VAXIGRIP, injekčná suspenzia naplnená v injekčnej striekačke Očkovacia látka proti chrípke (štiepený virión, inaktivovaná)	FR/H/0121/001	59/0228/07-S	SANOFI PASTEUR	SK
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, injektionsvätska, suspension i förfylld spruta. Influensavaccin (spjälkat virus, inaktiverat)	FR/H/0121/001	14029	SANOFI PASTEUR EUROPE	SE
VAXIGRIP, sospensione iniettabile in siringa priempita. Vaccino influenzale preparato con virus frammentati "split", inattivati	FR/H/0121/001	026032312	SANOFI PASTEUR EUROPE	IT
VAXIGRIP, sospensione iniettabile in siringa priempita. Vaccino influenzale preparato con virus frammentati "split", inattivati	FR/H/0121/001	026032209	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	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
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	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
iniettabile in siringa preriempita. Vaccino influenzale preparato con virus frammentati "split", 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	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
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 (vírião fragmentado, inativado)	FR/H/0121/001	4317889	SANOFI PASTEUR EUROPE	PT
VAXIGRIP, suspensão injetável em seringa pré-cheia Vacina contra a gripe (vírião fragmentado,	FR/H/0121/001	2637882	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
inativado)				
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
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, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/02	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal	not available	1289/2008/03	SANOFI PASTEUR	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
(virus fragmentat, inactivat)				
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/04	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/05	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/06	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/07	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/08	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/09	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabilă Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/10	SANOFI PASTEUR	RO
VAXIGRIP, suspensie injectabila, Vaccin gripal (virus fragmentat, inactivat)	not available	1289/2008/01	SANOFI PASTEUR	RO
VAXIGRIP, suspensie voor injectie in een voorgevulde spuit.	FR/H/0121/001	RVG 22306	SANOFI PASTEUR EUROPE	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
Griepvaccin (gesplitst virion, geïnactiveerd)				
VAXIGRIP, suspension for injection in prefilled syringe. Influenza vaccine (split virion, inactivated).	not available	MA573/00101	SANOFI PASTEUR	MT
VAXIGRIP, suspension i fyldt injektionssprøjte, Influenzavaccine, split virion, (inaktiveret)	FR/H/0121/001	11708	SANOFI PASTEUR EUROPE	DK
VAXIGRIP, suspension injectable en flacon multidose Vaccin grippal (inactivé, à virion fragmenté)	FR/H/0121/002	NL 11 155-2	SANOFI PASTEUR	FR
VAXIGRIP, suspension injectable en seringue préremplie. Vaccin grippal (inactivé, à virion fragmenté).	FR/H/0121/001	NL 11 155-1	SANOFI PASTEUR	FR
VAXIGRIP, suspensión inyectable en jeringa precargada Vacuna antigripal (virus fraccionados, inactivados)	FR/H/0121/001	61.108	SANOFI PASTEUR EUROPE	ES
VAXIGRIP, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο)	FR/H/0121/001	42844/05-09-2017	SANOFI PASTEUR EUROPE	GR
VAXIGRIP, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα. Αντιγριπικό εμβόλιο (split virion, αδρανοποιημένο).	FR/H/0121/001	20507	SANOFI PASTEUR	CY
VaxigripTetra injekcine	DE/H/1949/001	LT/1/16/3975/001	SANOFI PASTEUR	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
suspensija užpildytame švirkšte Keturvalente vakcina nuo gripo (iš virionu fragmentu, inaktyvuota)				
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/002	SANOFI PASTEUR	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	LT
VaxigripTetra injekcinė suspensija užpildytame švirkšte Keturvalentė vakcina nuo gripo (iš virionų fragmentų, inaktyvuota)	DE/H/1949/001	LT/1/16/3975/004	SANOFI PASTEUR	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	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	LT
VaxigripTetra Injektionssuspension in	DE/H/1949/001	137203	SANOFI PASTEUR EUROPE	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
einer Fertigspritze Tetravalenter Influenza- Impfstoff (Spaltvirus, inaktiviert)				
VaxigripTetra injektionsvätska, suspension i förfylld spruta Quadrivalent influensavaccin (spjälkat virus, inaktiverat)	DE/H/1949/001	33654	SANOFI PASTEUR EUROPE	FI
VaxigripTetra suspensie injectabila în seringă preumpluta	DE/H/1949/001	9198/2016/01	SANOFI PASTEUR	RO
VaxigripTetra suspensie injectabila în seringă preumpluta	DE/H/1949/001	9198/2016/02	SANOFI PASTEUR	RO
VaxigripTetra suspensie injectabila în seringă preumpluta	DE/H/1949/001	9198/2016/04	SANOFI PASTEUR	RO
VaxigripTetra suspensie injectabila în seringă preumpluta	DE/H/1949/001	9198/2016/05	SANOFI PASTEUR	RO
VaxigripTetra suspensie injectabila în seringă preumpluta	DE/H/1949/001	9198/2016/06	SANOFI PASTEUR	RO
VaxigripTetra suspensie injectabila în seringă preumpluta V	DE/H/1949/001	9198/2016/03	SANOFI PASTEUR	RO
VaxigripTetra suspensija injekcijām pilnšjircē Četrvērtīga gripas vakcīna (šķelts virions, inaktivēta) Quadrivalent influenza vaccine (split virion, inactivated)	DE/H/1949/001	16-0141	SANOFI PASTEUR	LV
VaxigripTetra suspensija za injiciranje v napoljnjeni	DE/H/1949/001	H/16/02251/001	SANOFI PASTEUR	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
injekcijski brizgi 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/16/02251/002	SANOFI PASTEUR	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/003	SANOFI PASTEUR	SI
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	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	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	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi	DE/H/1949/001	H/16/02251/007	SANOFI PASTEUR	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/16/02251/008	SANOFI PASTEUR	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	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	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	SI
VaxigripTetra suspenzija za injiciranje v napolnjeni injekcijski brizgi Štirivalentno cepivo proti gripi (razcepljeni virioni, inaktivirano)	DE/H/1949/001	H/16/02251/012	SANOFI PASTEUR	SI
Vaxigriptetra, injekcijske, suspenzije i ferdigfylt sprøyte.	DE/H/1949/001	15-10871	SANOFI PASTEUR EUROPE	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
Kvadrivalent vaksine mot influensa (inaktivert, splittvirus)				
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 Fyrvalent influensavaccin (spjälkat virus, inaktiverat)	DE/H/1949/001	53400	SANOFI PASTEUR EUROPE	SE
VaxigripTetra, stungulyf, dreifa í áfylltri sprautu. Fjörgilt influensubóluefni (klofin veiruögn, deydd)	DE/H/1949/001	IS/1/16/068/01	SANOFI PASTEUR EUROPE	IS
VaxigripTetra, suspension injectable en seringue préremplie Vaccin grippal quadrivalent (inactivé, à virion fragmenté)	DE/H/1949/001	NL46320	SANOFI PASTEUR	FR
VaxigripTetra, suspenzija za injekciju u napunjenoj štrcaljki Cetverovalentno cjepivo protiv influence (fragmentirani virion, inaktivirano)	DE/H/1949/001	HR-H-888872987	SANOFI PASTEUR	HR
VaxigripTetra, süstesuspensioon süstlis	DE/H/1949/001	919416	SANOFI PASTEUR	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	PL
VaxigripTetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Τετραδύναμο αντιγριπικό εμβόλιο (split virion, αδραντοποιημένο)	DE/H/1949/001	022512	SANOFI PASTEUR	CY
VaxigripTetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Τετραδύναμο αντιγριπικό εμβόλιο (split virion, αδραντοποιημένο)	DE/H/1949/001	51755/05-09-2017	SANOFI PASTEUR EUROPE	GR
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
ВаксигрипТетра инжекционна суспензия в предварително напълнена спринцовка Четирилентна ваксина	DE/H/1949/001	20160310	SANOFI PASTEUR	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
срещу грип (фрагментиран инактивиран вирион)				
Флуарикс Тетра инжекционна суспензия в предварително напълнена спринцовка Ваксина срещу грип (фрагментиран вирион, инактивирана)	DE/H/1939/001	20180166	GLAXOSMITHKLINE EOOD	BG