11 February 2021
EMA/61734/2021
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: diphtheria / tetanus / pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed), diphtheria / tetanus / pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed) reduce antigens content

Procedure no.: PSUSA/00001126/202007

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TETRAVAC Injektionssuspension Diphtherie-, Tetanus-, Pertussis-(azellulär, Komponenten) und Poliomyelitis-(inaktiviert) Impfstoff, adsorbiert	SE/H/0154/001	2-00226	SANOFI PASTEUR EUROPE	AT
REPEVAX Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	2-00284	SANOFI PASTEUR EUROPE	AT
REPEVAX Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	2-00284	SANOFI PASTEUR EUROPE	AT
Boostrix®Polio - Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis azellulär- und Poliomyelitis (inaktiviert) - Impfstoff (adsorbiert, verminderter Antigengehalt)	DE/H/0466/003	2-00302	GLAXOSMITHKLINE PHARMA GMBH.	AT
DTaP-IPV Vakzine SSI Injektionssuspension in einer Fertigspritze. Diphtherie-, Tetanus-, Pertussis-(azelluläre Komponente) und Polio- (inaktiviert) Impfstoff (adsorbiert)	DK/H/0132/001	2-00232	AJ VACCINES A/S	AT
Tetravac, suspension injectable Vaccin diphtérique, tétanique, coquelucheux (acellulaire,	SE/H/0154/001	BE196122	SANOFI PASTEUR EUROPE	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
composant) et poliomyélitique (inactivé), adsorbé				
Tetravac, suspensie voor injectie Geadsorbeerd difterie-, tetanus-, kinkhoest- (acellulair, component) en poliovaccin (geïnactiveerd)	SE/H/0154/001	BE196122	SANOFI PASTEUR EUROPE	BE
Tetravac, Suspension zur Injektion Impfstoff (adsorbiert) gegen Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert)	SE/H/0154/001	BE196122	SANOFI PASTEUR EUROPE	BE
Infanrix-IPV, 0,5 ml/dosis, suspensie voor injectie	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Infanrix-IPV, 0,5 ml/Dosis Injizierbare Suspension	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Infanrix-IPV, 0,5 ml/dose, suspension injectable	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	BE
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	BE
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze Diphtherie-,	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	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
Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)				
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	BE
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	BE
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	BE241096	SANOFI PASTEUR EUROPE	BE
Boostrix Polio, suspension injectable en seringue pré-remplie Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélitique (inactivé)	DE/H/0466/003	BE270435	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Boostrix Polio, suspension	DE/H/0466/004	BE270426	GLAXOSMITHKLINE	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
injectable Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélitique (inactivé)			BIOLOGICALS S.A.	
Boostrix Polio, Injektionssuspension in einer Fertigspritze Impfstoff (adsorbiert, Inhalt antigenreduziert) gegen Diphtherie, Tetanus, Pertussis (azellulär) und Poliomyelitis (inaktiv)	DE/H/0466/003	BE270435	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Boostrix Polio, Injektionssuspension Impfstoff (adsorbiert, Inhalt antigenreduziert) gegen Diphtherie, Tetanus, Pertussis (azellulär) und Poliomyelitis (inaktiv)	DE/H/0466/004	BE270426	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Boostrix Polio, suspensie voor injectie in voorgevulde injectiespuit Vaccin (geadsorbeerd, antigengereduceerde inhoud) tegen difterie, tetanus, kinkhoest (acellulaire component) en poliomyelitis (geïnactiveerd)	DE/H/0466/003	BE270435	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Boostrix Polio, suspensie voor injectie Vaccin (geadsorbeerd, antigengereduceerde inhoud) tegen difterie, tetanus, kinkhoest (acellulaire component) en poliomyelitis (geïnactiveerd)	DE/H/0466/004	BE270426	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
АДАСЕЛ ПОЛИО, инжекционна суспензия в предварително напълнена спринцовка. Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана), с намалено антигенно съдържание	DE/H/0215/001	20200008	SANOFI PASTEUR	BG
АДАСЕЛ ПОЛИО, инжекционна суспензия в предварително напълнена спринцовка. Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана), с намалено антигенно съдържание	DE/H/0215/001	20200008	SANOFI PASTEUR	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI PASTEUR	BG
Бустрикс Полио инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/004	20120314	GLAXOSMITHKLINE EOOD	BG
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας	FR/H/0251/002	20232	GLAXOSMITHKLINE (CYPRUS) LIMITED	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(αδρανοποιημένης).				
TETRAXIM, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα, Εμβόλιο (προσροφημένο), διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο)	HU/H/0406/001	022511	SANOFI PASTEUR	CY
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	MARKET AUTHORISATION APPLICATION - 310 - EU MRP -	59/265/19-C	SANOFI PASTEUR	CZ
Infanrix Polio injekční suspenze v předplněné injekční stříkačce Adsorbovaná vakcína proti difterii, tetanu, pertusi (acelulární komponenta) a poliomyelitidě (inaktivovaná vakcína).	FR/H/0251/002	59/268/05-C	GLAXOSMITHKLINE BIOLOGICALS S.A.	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	MARKET AUTHORISATION APPLICATION - 310 - EU MRP -	59/265/19-C	SANOFI PASTEUR	CZ
Boostrix Polio injekční suspenze v předplněné injekční stříkačce Adsorbovaná vakcína proti difterii, tetanu, pertusi (acelulární komponenta) a poliomyelitidě (inaktivovaná vakcína) se sníženým obsahem antigenů.	DE/H/0466/003	59/497/07-C	GLAXOSMITHKLINE S.R.O.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tetraxim, injekční suspenze v předplněné injekční stříkačce Adsorbovaná vakcína proti difterii, tetanu, pertusi (acelulární komponenta), poliomyelitidě (inaktivovaná)	HU/H/0406/001	59/285/16-C	SANOFI PASTEUR	CZ
TETRAVAC Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis (azellulär)-Poliomyelitis (inaktiviert)-Adsorbat-Impfstoff	not available	1A/97	SANOFI PASTEUR EUROPE	DE
REPEVAX Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	PEI.H.02354.01.1	SANOFI PASTEUR EUROPE	DE
REPEVAX Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	PEI.H.02354.01.1	SANOFI PASTEUR EUROPE	DE
Boostrix® Polio - Suspension zur Injektion in Glasflaschchen Diphtherie- Tetanus-Pertussis (azelluHire Bestandteile)-inaktivierter Poliomyelitis Kombinationsimpfstoff (adsorbiert) zur Auf:frischimpfung	DE/H/0466/004	PEI.H.02950.01.2	GLAXOSMITHKLINE GMBH & CO. KG	DE
Boostrix® Polio Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis (azellulär,	DE/H/0466/003	PEI.H.02950.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
aus Komponenten)- und Poliomyelitis (inaktiviert)-Adsorbatimpfstoff (mit reduziertem Antigengehalt)				
DiTeKiPol Booster, injektionsvæske, suspension, fyldt injektionssprøjte	not available	35304	AJ VACCINES A/S	DK
Tetravac Injektionsvæske, suspension Difteri-, tetanus-, pertussis-(acellulær, komponent) og poliovaccine (inaktiveret), adsorberet	SE/H/0154/001	30186	SANOFI PASTEUR EUROPE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er))	DE/H/0215/001	33442	SANOFI PASTEUR EUROPE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er))	DE/H/0215/001	33442	SANOFI PASTEUR EUROPE	DK
Boostrix Polio, injektionsvæske, suspension, hætteglas Difteri-, tetanus-, kighoste (acellulær)-, polio (inaktiveret)-vaccine (adsorberet)	DE/H/0466/004	49762	GLAXOSMITHKLINE PHARMA A/S	DK
Boostrix Polio, injektionsvæske, suspension, fyldt injektionssprøjte	DE/H/0466/003	49761	GLAXOSMITHKLINE PHARMA A/S	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Injektionsvæske				
DiTeKiPol, injektionsvæske, suspension i fyldt engangssprøjte	DK/H/0132/001	17843	AJ VACCINES A/S	DK
Infanrix Polio, süstesuspensioon süstlis. Difteeria, teetanuse, läkaköha (atsellulaarne komponentvaktsiin) ja poliomüeliidi (inaktiveeritud) vaktsiin (adsorbeeritud).	FR/H/0251/002	483205	GLAXOSMITHKLINE BIOLOGICALS S.A.	EE
TETRAXIM, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüeliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI PASTEUR	EE
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomielitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001	84.886	SANOFI PASTEUR	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomielitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001	84.886	SANOFI PASTEUR	ES
Boostrix Polio suspensión inyectable en jeringa precargada Vacuna antidiftérica, antitetánica, antitos ferina (componente acelular),	DE/H/0466/003	66.462	GLAXOSMITHKLINE, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
antipoliomielitis (inactivada) (adsorbida, contenido antigénico reducido)				
Tetravac, injektioneste, suspensio Adsorboitu kurkkumätä-, jäykkäkouristus-, hinkuyskä- (soluton, komponentti) ja polio (inaktivoitu) - rokote..	SE/H/0154/001	13676	SANOFI PASTEUR EUROPE	FI
Tetravac injektionsvätska, suspension Difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat) vaccin, adsorberat	SE/H/0154/001	13676	SANOFI PASTEUR EUROPE	FI
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä-, jäykkäkouristus-, hinkuyskä- (soluton komponentti) ja polio- (inaktivoitu) -rokote (adsorboitu, alhaisempi antigeenimäärä)	DE/H/0215/001	17190	SANOFI PASTEUR EUROPE	FI
REPEVAX Injektionsvätska, suspension i förfylld spruta Difteri-, tetanus-, pertussis- (acellulärt, komponent) och (inaktiverat) poliovaccin (adsorberat, lägre antigenhalt)	DE/H/0215/001	17190	SANOFI PASTEUR EUROPE	FI
Infanrix-Polio, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat	FR/H/0251/002	20846	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
Infanrix-Polio injektioneste, suspensio esitäytetyssä ruiskussa.Kurkkumätä-,	FR/H/0251/002	20846	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
jäykkäkouristus-, hinkuyskä (soluton, komponentti), poliomyeliitti (inaktivoitu) rokote (adsorboitu).				
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä-, jäykkäkouristus-, hinkuyskä- (soluton komponentti) ja polio- (inaktivoitu) -rokote (adsorboitu, alhaisempi antigeenimäärä)	DE/H/0215/001	17190	SANOFI PASTEUR EUROPE	FI
REPEVAX Injektionsvätska, suspension i förfylld spruta Difteri-, tetanus-, pertussis- (acellulärt, komponent) och (inaktiverat) poliovaccin (adsorberat, lägre antigenhalt)	DE/H/0215/001	17190	SANOFI PASTEUR EUROPE	FI
Boostrix Polio injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0466/003	19479	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
Boostrix Polio - injektioneste, suspensio esitäytetyssä ruiskussa Kurkkumätä-, jäykkäkouristus-, hinkuyskä (soluton, komponentti), poliomyeliitti (inaktivoitu) rokote, (adsorboitu, matala antigeenipitoisuus)	DE/H/0466/003	19479	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
DiTeKiPol Injektionsvätska, suspension i förfylld spruta. Difteri, tetanus, pertussis (acellulär, komponent) och	DK/H/0132/001	13440	AJ VACCINES A/S	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
poliomyelit (inaktiverat) vaccin (adsorberat)				
DiTeKiPol, injektioneste suspensio, esitäytetyssä ruiskussa Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja (inaktivoituja) polio (-viruksia sisältävä) rokote (adsorboitu)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
DiTeKiPol, injektioneste suspensio, esitäytetyssä ruiskussa Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja (inaktivoituja) polio (-viruksia sisältävä) rokote (adsorboitu)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
DiTeKiPol Injektionsvätska, suspension i förfylld spruta. Difteri, tetanus, pertussis (acellulär, komponent) och poliomyelit (inaktiverat) vaccin (adsorberat)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
DiTeKiPol, injektioneste suspensio, esitäytetyssä ruiskussa Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja (inaktivoituja) polio (-viruksia sisältävä) rokote (adsorboitu)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
DiTeKiPol Injektionsvätska, suspension i förfylld spruta. Difteri, tetanus, pertussis (acellulär, komponent) och poliomyelit (inaktiverat) vaccin (adsorberat)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
DiTeKiPol, injektioneste suspensio, esitäytetyssä	DK/H/0132/001	13440	AJ VACCINES A/S	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
ruiskussa Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja (inaktivoituja) polio (-viruksia sisältävä) rokote (adsorboitu)				
DiTeKiPol Injektionsvätska, suspension i förfylld spruta. Difteri, tetanus, pertussis (acellulär, komponent) och poliomyelit (inaktiverat) vaccin (adsorberat)	DK/H/0132/001	13440	AJ VACCINES A/S	FI
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 360 647 6 8	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 348 221 2 4	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 360 644 7 8	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et	not available	34009 360 645 3 9	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
poliomyélitique (inactivé), adsorbé				
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 348 223 5 3	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 351 668 4 5	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 368 746 3 3	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 368 748 6 2	SANOFI PASTEUR EUROPE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 368 749 2 3	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
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 368 750 0 5	SANOFI PASTEUR EUROPE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 347 823 9 8	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 391 896 8 0	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 348 222 9 2	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 359 667 7 3	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin	not available	34009 359 668 3 4	SANOFI PASTEUR	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
diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé				
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 368 751 7 3	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 368 752 3 4	SANOFI PASTEUR	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), adsorbé	not available	34009 391 897 4 1	SANOFI PASTEUR	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 648-2	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique,	DE/H/0215/001	359 647-6	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
coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))				
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 645-3	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 740-5	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 739-7	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 738-0	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
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 737-4	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 644-7	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 643-0	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 642-4	SANOFI PASTEUR EUROPE	FR
INFANRIXTETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux acellulaire, poliomyélitique	FR/H/0251/002	NL21433	LABORATOIRE GLAXOSMITHKLINE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
inactivé, adsorbé				
INFANRIXTETRA, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire, poliomyélitique inactivé, adsorbé	FR/H/0251/002	34009 355 246 7 6	LABORATOIRE GLAXOSMITHKLINE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 648-2	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 647-6	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 645-3	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé),	DE/H/0215/001	368 740-5	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
(adsorbé, à teneur réduite en antigène(s))				
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 739-7	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 738-0	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	368 737-4	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 644-7	SANOFI PASTEUR EUROPE	FR
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique,	DE/H/0215/001	359 643-0	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
coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))				
REPEVAX, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigène(s))	DE/H/0215/001	359 642-4	SANOFI PASTEUR EUROPE	FR
BOOSTRIXTETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	34009 367 738 7 5	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXTETRA, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigens)	DE/H/0466/003	34009 367 740 1 8	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXTETRA, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et	DE/H/0466/003	34009 367 741 8 6	LABORATOIRE GLAXOSMITHKLINE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigens)				
BOOSTRIXTETRA, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigens)	DE/H/0466/003	34009 367 739 3 6	LABORATOIRE GLAXOSMITHKLINE	FR
Tetravac, ενέσιμο εναιώρημα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο)	SE/H/0154/001	14662/02-03-2018 & 106295/17/17-04-2018	SANOFI PASTEUR EUROPE	GR
REPEVAX, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων)).	DE/H/0215/001/E1	88313/10-07-2019	SANOFI PASTEUR EUROPE	GR
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένης)	FR/H/0251/002	2682901	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
REPEVAX, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη	DE/H/0215/001/E1	88313/10-07-2019	SANOFI PASTEUR EUROPE	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων)).				
Boostrix Polio ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσροφημένο, μειωμένο(α) περιεχόμενο(α) σε αντιγόνο(α) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένο)	DE/H/0466/003	2650601	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
ADACEL POLIO, suspenzija za injekciju u napunjenoj štrcaljki Cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano) sa smanjenim sadržajem antigena, adsorbirano	DE/H/0215/001	HR-H-773372732	SANOFI PASTEUR	HR
ADACEL POLIO, suspenzija za injekciju u napunjenoj štrcaljki Cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano) sa smanjenim sadržajem antigena, adsorbirano	DE/H/0215/001	HR-H-773372732	SANOFI PASTEUR	HR
TETRAXIM, suspenzija za injekciju u napunjenoj štrcaljki, cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano), adsorbirano	HU/H/0406/001	HR-H-185201369	SANOFI PASTEUR	HR
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz,	DE/H/0215/001	OGYI-T-23632/01	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
pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)				
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/02	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/03	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/04	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/05	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz	DE/H/0215/001	OGYI-T-23632/06	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) vakcina (adszorbeált, csökkentett antigéntartalmú)				
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/07	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/08	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/09	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/10	SANOFI PASTEUR	HU
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina	FR/H/0251/002	OGYI-T-20352/02	GLAXOSMITHKLINE KFT.	HU
Infanrix IPV szuszpenziós	FR/H/0251/002	OGYI-T-20352/05	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
injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina				
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina	FR/H/0251/002	OGYI-T-20352/06	GLAXOSMITHKLINE KFT.	HU
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina	FR/H/0251/002	OGYI-T-20352/01	GLAXOSMITHKLINE KFT.	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/01	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/02	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált,	DE/H/0215/001	OGYI-T-23632/03	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
csökkentett antigéntartalmú)				
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/04	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/05	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/06	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/07	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/08	SANOFI PASTEUR	HU
ADACEL POLIO szuszpenziós	DE/H/0215/001	OGYI-T-23632/09	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
injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)				
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001	OGYI-T-23632/10	SANOFI PASTEUR	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/02	GLAXOSMITHKLINE KFT.	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/03	GLAXOSMITHKLINE KFT.	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/04	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
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/07	GLAXOSMITHKLINE KFT.	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/08	GLAXOSMITHKLINE KFT.	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén tartalmú)	DE/H/0466/003	OGYI-T-20497/01	GLAXOSMITHKLINE KFT.	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H60406/001	OGYI-T-9452/01	SANOFI PASTEUR	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/02	SANOFI PASTEUR	HU
Tetraxim szuszpenziós injekció	HU/H/0406/001	OGYI-T-9452/03	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
előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)				
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/04	SANOFI PASTEUR	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/05	SANOFI PASTEUR	HU
Tetravac, suspension for injection Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, adsorbed	SE/H/0154/001	PA 2131/009/001	SANOFI PASTEUR EUROPE	IE
REPEVAX, suspension for injection, in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	PA 2131/006/001	SANOFI PASTEUR EUROPE	IE
IPV Infanrix, suspension for injection in pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed)	FR/H/0251/002	PA 1077/108/001	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REPEVAX, suspension for injection, in pre-filled syringe	DE/H/0215/001	PA 2131/006/001	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
Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)				
IPV-Boostrix suspension for injection in pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed, reduced antigen(s) content	DE/H/0466/003	PA 1077/101/001	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Tetravac, stungulyf, dreifa Barnaveiki-, stífkrampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt)	SE/H/0154/001	970045	SANOFI PASTEUR EUROPE	IS
REPEVAX, stungulyf, dreifa, í áfylltri sprautu Barnaveiki, stífkrampa, kíghósta (frumulaust, samsett) og mænusóttarveiru (dautt) bóluefni, (aðsogað, minnkað mótefnisvaka innihald)	DE/H/0215/001	IS/1/02/020/01	SANOFI PASTEUR EUROPE	IS
REPEVAX, stungulyf, dreifa, í áfylltri sprautu Barnaveiki, stífkrampa, kíghósta (frumulaust, samsett) og mænusóttarveiru (dautt) bóluefni, (aðsogað, minnkað mótefnisvaka innihald)	DE/H/0215/001	IS/1/02/020/01	SANOFI PASTEUR EUROPE	IS
Boostrix Polio stungulyf, dreifa í áfylltri sprautu Bóluefni gegn barnaveiki, stífkrampa, kíghósta (frumulaust, hlutar) og mænusótt (deyddar veirur), (aðsogað, skert	DE/H/0466/003	IS/1/07/027/01	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
mótefnavakainnihald)				
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127011	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127023	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127035	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127047	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127062	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127050	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127074	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile	SE/H/0154/001	034127086	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 adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)				
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127098	SANOFI PASTEUR EUROPE	IT
Tetravac sospensione iniettabile Vaccino adsorbito antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato)	SE/H/0154/001	034127100	SANOFI PASTEUR EUROPE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290011	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290023	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico	DE/H/0215/001	048290035	SANOFI PASTEUR	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
(componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).				
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290047	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290050	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290062	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e	DE/H/0215/001	048290074	SANOFI PASTEUR	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
antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).				
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290086	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290098	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290100	SANOFI PASTEUR	IT
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato).	FR/H/0251/002	037157094	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
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato)	FR/H/0251/002	037157043	GLAXOSMITHKLINE S.P.A.	IT
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato).	FR/H/0251/002	037157068	GLAXOSMITHKLINE S.P.A.	IT
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato)	FR/H/0251/002	037157070	GLAXOSMITHKLINE S.P.A.	IT
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato).	FR/H/0251/002	037157031	GLAXOSMITHKLINE S.P.A.	IT
PolioInfanrix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-poliomielite (inattivato)	FR/H/0251/002	037157017	GLAXOSMITHKLINE S.P.A.	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e	DE/H/0215/001	048290011	SANOFI PASTEUR	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
antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).				
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290023	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290035	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290047	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato)	DE/H/0215/001	048290050	SANOFI PASTEUR	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
(adsorbito, contenuto antigenico ridotto).				
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290062	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290074	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290086	SANOFI PASTEUR	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico	DE/H/0215/001	048290098	SANOFI PASTEUR	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
ridotto).				
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001	048290100	SANOFI PASTEUR	IT
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e poliomielitico (inattivato)	DE/H/0466/003	036752020	GLAXOSMITHKLINE S.P.A.	IT
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e poliomielitico (inattivato)	DE/H/0466/003	036752069	GLAXOSMITHKLINE S.P.A.	IT
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e poliomielitico (inattivato)	DE/H/0466/003	036752018	GLAXOSMITHKLINE S.P.A.	IT
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e	DE/H/0466/003	036752057	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
poliomielitico (inattivato)				
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e poliomielitico (inattivato)	DE/H/0466/003	036752032	GLAXOSMITHKLINE S.P.A.	IT
PolioBoostrix sospensione iniettabile in siringa preriempita. Vaccino (adsorbito, a ridotto contenuto di antigeni) difterico, tetanico, pertossico (componente acellulare) e poliomielitico (inattivato)	DE/H/0466/003	036752044	GLAXOSMITHKLINE S.P.A.	IT
TETRAXIM injekcine suspensija užpildytame švirkšte, Vakcina nuo difterijos, stabliges, kokliušo (nelasteline, komponentine) ir nuo poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/001	SANOFI PASTEUR	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte. Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/002	SANOFI PASTEUR	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte. Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/003	SANOFI PASTEUR	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte. Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir	not available	LT/1/99/2298/004	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
nuo poliomielito (inaktyvuota), (adsorbuota)				
TETRAXIM injekcinė suspensija užpildytame švirkšte. Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/005	SANOFI PASTEUR	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte. Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/006	SANOFI PASTEUR	LT
Infanrix Polio injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos (D), stabligės (T), kokliušo (neląstelinė, komponentinė) (Pa) ir nuo poliomielito (inaktyvuota) (IPV), (adsorbuota)	FR/H/0251/002	LT/1/05/0301/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Infanrix Polio injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos (D), stabligės (T), kokliušo (neląstelinė, komponentinė) (Pa) ir nuo poliomielito (inaktyvuota) (IPV), (adsorbuota)	FR/H/0251/002	LT/1/05/0301/001	GLAXOSMITHKLINE LIETUVA UAB	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/002	GLAXOSMITHKLINE LIETUVA UAB	LT
BOOSTRIX POLIO injekcinė	DE/H/0466/003	LT/1/08/0971/005	GLAXOSMITHKLINE LIETUVA	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 Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)			UAB	
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/006	GLAXOSMITHKLINE LIETUVA UAB	LT
BOOSTRIX POLIO injekcinė suspensija Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/004	LT/1/08/0971/003	GLAXOSMITHKLINE LIETUVA UAB	LT
BOOSTRIX POLIO injekcine suspensija Vakcina nuo difterijos, stabliges, kokliušo (nelasteline, komponentine) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenu kiekio)	DE/H/0466/004	LT/1/08/0971/004	GLAXOSMITHKLINE LIETUVA UAB	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/001	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
Tetravac, Suspension zur Injektion Impfstoff (adsorbiert) gegen Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert)	SE/H/0154/001	2008089883	SANOFI PASTEUR EUROPE	LU
Tetravac, suspension injectable Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé), adsorbé	SE/H/0154/001	2008089883	SANOFI PASTEUR EUROPE	LU
Infanrix-IPV, 0,5 ml/dose, suspension injectable	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
Infanrix-IPV, 0,5 ml/Dosis Injizierbare Suspension Impfstoff gegen Diphtherie, Tetanus, Keuchhusten und Poliomyelitis	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))	MARKET AUTHORISATION APPLICATION - 310 - EU MRP -	2002090039	SANOFI PASTEUR EUROPE	LU
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	2002090039	SANOFI PASTEUR EUROPE	LU
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique,	MARKET AUTHORISATION APPLICATION - 310 - EU	2002090039	SANOFI PASTEUR EUROPE	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tétanique, coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))	MRP -			
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001	2002090039	SANOFI PASTEUR EUROPE	LU
Boostrix Polio, suspension injectable en seringue pré-remplie Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélitique (inactivé)	DE/H/0466/003	2009020265	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
Boostrix Polio, Injektionssuspension Impfstoff (adsorbiert, Inhalt antigenreduziert) gegen Diphtherie, Tetanus, Pertussis (azellulär) und Poliomyelitis (inaktiv)	DE/H/0466/003	2009020265	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
TETRAXIM suspensija injekcijai pilnšļircē Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivētā poliomielīta vakcīna. Diphtheria, tetanus, pertussis (acellular component) and poliomyelitis (inactivated)	not available	02-0343	SANOFI PASTEUR	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
vaccine (adsorbed)				
Infanrix Polio suspensija injekcijām pilnšļircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomielīta (inaktivēta) vakcīna (adsorbēta)	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Boostrix Polio suspensija injekcijai pilnšļircēs Difterijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna (adsorbēta, ar samazinātu antigēnu saturu)	DE/H/0466/003	07-0238	GLAXOSMITHKLINE LATVIA SIA	LV
Boostrix Polio suspensija injekcijai Difterijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna (adsorbēta, ar samazinātu antigēnu saturu)	DE/H/0466/003	07-0239	GLAXOSMITHKLINE LATVIA SIA	LV
IPV-Boostrix suspension for injection Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed, reduced antigen(s) content)	DE/H/0466/003	MA 170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
IPV-Boostrix suspension for injection Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed, reduced antigen(s) content)	DE/H/0466/004	MA 170/00103	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
Infanrix-IPV, suspensie voor injectie Difterie-(D), tetanus-(T),	not available	RVG 34568	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
pertussis-(acellulair, component) (Pa) en poliomyelitis-(geïnactiveerd) (IPV) vaccin.				
TRIAXIS® POLIO, suspensie voor injectie in voorgevulde spuit Difterie, Tetanus, Kinkhoest (acellulair, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001	RVG 27569	SANOFI PASTEUR EUROPE	NL
TRIAXIS® POLIO, suspensie voor injectie in voorgevulde spuit Difterie, Tetanus, Kinkhoest (acellulair, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001	RVG 27569	SANOFI PASTEUR EUROPE	NL
Boostrix Polio suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, verminderd aantal antigenen)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Boostrix Polio injectieflacon suspensie voor injectie Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, verminderd aantal antigenen)	DE/H/0466/004	RVG 35124	GLAXOSMITHKLINE B.V.	NL
TETRAVAC, injeksjonsvæske, suspensjon Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt	not available	97-704	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
(inaktivert), (adsorbert)				
REPEVAX suspensjon til injeksjon i en ferdigfylt sprøyte Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert) (adsorbert, redusert antigeninnhold)	DE/H/0215/001	02-1069	SANOFI PASTEUR EUROPE	NO
Infanrix Polio injeksjonsvæske, suspensjon i ferdigfylt sprøyte. Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	FR/H/0251/002	05-3312	GLAXOSMITHKLINE AS	NO
REPEVAX suspensjon til injeksjon i en ferdigfylt sprøyte Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert) (adsorbert, redusert antigeninnhold)	DE/H/0215/001	02-1069	SANOFI PASTEUR EUROPE	NO
Boostrix Polio injeksjonsvæske, suspensjon, i ferdigfylt sprøyte Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert, redusert innhold av antigen(er))	DE/H/0466/003	04-2685	GLAXOSMITHKLINE AS	NO
Boostrix Polio injeksjonsvæske, suspensjon, i ferdigfylt sprøyte Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert, redusert innhold av antigen(er)).	DE/H/0466/003	04-2685	GLAXOSMITHKLINE AS	NO
Adacel Polio, zawiesina do wstrzykiwan w ampulko-	DE/H/0215/001	25842	SANOFI PASTEUR	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
strzykawce. Szczepionka przeciw blonicy, tezcowi, krztuscowi (bezkomorkowa, zlozona) i poliomyelitis (inaktywowana), absorbowana,o zmniejszonej zawartosci antydenow				
Infanrix-IPV, zawiesina do wstrzykiwan w ampulkostrzykawce Szczepionka przeciw blonicy, tezcowi, krztuscowi (bezkomórkowa, zlozona) i poliomyelitis (inaktywowana), adsorbowana	FR/H/0251/002	11884	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Adacel Polio, zawiesina do wstrzykiwan w ampulko-strzykawce. Szczepionka przeciw blonicy, tezcowi, krztuscowi (bezkomorkowa, zlozona) i poliomyelitis (inaktywowana), absorbowana,o zmniejszonej zawartosci antydenow	DE/H/0215/001	25842	SANOFI PASTEUR	PL
Boostrix Polio, zawiesina do wstrzykiwań Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0466/003	14240	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Boostrix Polio, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i	DE/H/0466/003	14241	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
poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów				
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce, Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana	HU/H/0406/001	23545	SANOFI PASTEUR SA	PL
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5376389	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	2782480	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5458583	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5458781	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5376488	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável	SE/H/0154/001	2782589	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
Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)				
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5458682	SANOFI PASTEUR EUROPE	PT
Tetravac, suspensão injetável Vacina adsorvida contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada)	SE/H/0154/001	5458484	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083283	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459680	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083796	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina	DE/H/0215/001	5459581	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
(adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)				
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459789	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459482	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083382	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083887	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria,	DE/H/0215/001	4083184	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
tétano, tosse convulsa (componente acelular) e poliomielite (inativada)				
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083986	SANOFI PASTEUR EUROPE	PT
Infanrix Tetra, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativado)	FR/H/0251/002	5621081	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Infanrix Tetra, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativado)	FR/H/0251/002	5267356	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083283	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459680	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável	DE/H/0215/001	4083796	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
em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)				
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459581	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459789	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	5459482	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083382	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de	DE/H/0215/001	4083887	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
antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)				
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083184	SANOFI PASTEUR EUROPE	PT
REPEVAX suspensão injectável em seringa pré-cheia Vacina (adsorvida, teor reduzido de antigénio(s)) contra a difteria, tétano, tosse convulsa (componente acelular) e poliomielite (inativada)	DE/H/0215/001	4083986	SANOFI PASTEUR EUROPE	PT
Boostrix Polio Suspensão injetável Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/004	5212881	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/004	5212980	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável em seringa pré-cheia Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite	DE/H/0466/003	5212683	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(inativado) (adsorvida, com conteúdo reduzido de antigénio(s))				
Boostrix Polio Suspensão injetável em seringa pré-cheia Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/003	5212782	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável em seringa pré-cheia Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/003	5270889	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável em seringa pré-cheia Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/003	5270988	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável em seringa pré-cheia Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))	DE/H/0466/003	5271085	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Boostrix Polio Suspensão injetável em seringa pré-cheia	DE/H/0466/003	5271184	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vacina contra a difteria, tétano e tosse convulsa (componente acelular) e poliomielite (inativado) (adsorvida, com conteúdo reduzido de antigénio(s))			FARMACEUTICOS LDA	
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/01	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/02	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/03	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/04	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis	DE/H/0215/001	12959/2020/05	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
(acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))				
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/06	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/07	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/08	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/09	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat),	DE/H/0215/001	12959/2020/10	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
(adsorbit, cu conţinut redus de antigen(e))				
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/01	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/02	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/03	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/04	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/05	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
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/06	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/07	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/08	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/09	SANOFI PASTEUR	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomielitic (inactivat), (adsorbit, cu conţinut redus de antigen(e))	DE/H/0215/001	12959/2020/10	SANOFI PASTEUR	RO
TETRAXIM suspensie injectabilă în seringă preumplută	not available	4490/2012/01	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
Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit				
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit	not available	4490/2012/02	SANOFI PASTEUR	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit	not available	4490/2012/03	SANOFI PASTEUR	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit	not available	4490/2012/04	SANOFI PASTEUR	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit	not available	4490/2012/05	SANOFI PASTEUR	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/03	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/02	GLAXOSMITHKLINE (GSK) S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/01	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/04	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/05	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) şi poliomielitic (inactivat) (adsorbit, cu conţinut redus de antigen(e))	DE/H/0466/003	8900/2016/06	GLAXOSMITHKLINE (GSK) S.R.L.	RO
Tetravac, injektionsvätska, suspension. Difteri-, tetanus-, pertussis- (acellulärt, komponent) och poliomyelitisvaccin (inaktiverat) adsorberat	SE/H/0154/001	13464	SANOFI PASTEUR EUROPE	SE
Repevax, injektionsvätska, suspension i förfylld spruta	MARKET AUTHORISATION	59538	SANOFI PASTEUR	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
Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	APPLICATION - 310 - EU MRP -			
Infanrix Polio, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	FR/H/0251/002	22287	GLAXOSMITHKLINE AB	SE
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	MARKET AUTHORISATION APPLICATION - 310 - EU MRP -	59538	SANOFI PASTEUR	SE
Boostrix Polio, injektionsvätska, suspension i förfylld spruta. Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0466/003	21068	GLAXOSMITHKLINE AB	SE
DiTeKiPol, injektionsvätska, suspension, förfylld spruta. Difteri, Tetanus, Pertussis (acellulär, komponent) och Poliomyelit (inaktiverat) Vaccin (adsorberat)	DK/H/0132/001	14398	AJ VACCINES A/S	SE
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis	DE/H/0215/001	H/20/02707/001	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
(acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)				
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/002	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/003	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/004	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/006	SANOFI PASTEUR	SI
Adacel Polio suspenzija za	DE/H/0215/001	H/20/02707/005	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
injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)				
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/007	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/008	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/009	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated)	DE/H/0215/001	H/20/02707/010	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
Vaccine (adsorbed, reduced antigen(s) content)				
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/001	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/002	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/003	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/004	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi	DE/H/0215/001	H/20/02707/006	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
Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)				
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/005	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/007	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/008	SANOFI PASTEUR	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/009	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
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	H/20/02707/010	SANOFI PASTEUR	SI
Boostrix Polio suspenzija za injiciranje cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/005	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/003	H/10/00651/001	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/002	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano	DE/H/0466/004	H/10/00651/003	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
(adsorbirano, z zmanjšano vsebnostjo antigenov)				
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/004	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/006	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/007	GLAXOSMITHKLINE D.O.O.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskemu kašlju – acelularno in otroški paralizi - inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/004	H/10/00651/008	GLAXOSMITHKLINE D.O.O.	SI
Adacel Polio injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná s	DE/H/0215/001	59/0035/20-S	SANOFI PASTEUR	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
redukovaným obsahom antigénov) proti záškrtu, tetanu, čiernemu kašľu (nebunkové zložky) a detskej obrne (inaktivovaná)				
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)inaktivovaná	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Adacel Polio injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná s redukovaným obsahom antigénov) proti záškrtu, tetanu, čiernemu kašľu (nebunkové zložky) a detskej obrne (inaktivovaná)	DE/H/0215/001	59/0035/20-S	SANOFI PASTEUR	SK
Boostrix Polio Injekčná suspenzia naplnená v injekčnej striekačke Očkovacia látka (adsorbovaná, so zníženým obsahom antigénu (antigénov)) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka pertussis) a detskej obrne (inaktivovaná)	DE/H/0466/003	59/0373/07-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
TETRAXIM Injekčná suspenzia naplnená v injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka),	HU/H/0406/001	59/0291/16-S	SANOFI PASTEUR	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
detskej obrne (inaktivovaná)				
Tetravac, suspension for injection Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, adsorbed	SE/H/0154/001	PL 46602/0018	SANOFI PASTEUR EUROPE	XI
REPEVAX, suspension for injection, in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	PL 46602/0005	SANOFI PASTEUR EUROPE	XI
Infanrix-IPV, suspension for injection in pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed)	FR/H/0251/002	PL 10592/0209	SMITHKLINE BEECHAM LTD	XI
REPEVAX, suspension for injection, in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001	PL 46602/0005	SANOFI PASTEUR EUROPE	XI
Boostrix-IPV suspension for injection in pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed, reduced antigen(s) content)	DE/H/0466/003	PL 10592/0214	SMITHKLINE BEECHAM LTD	XI