



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

12 February 2026
EMADOC-1700519818-2982954
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): diphtheria / tetanus / pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed), diphtheria / tetanus / pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed, reduced antigen(s) content)

EURD list No. PSUSA/00001126/202507

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
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
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
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT
Repevax Injektionssuspension in einer Fertigspritze	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)- Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)				
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)- Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)- Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)- Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)- Poliomyelitis(inaktiviert)-Impfstoff	DE/H/0215/001-002	2-00284	SANOFI WINTHROP INDUSTRIE	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(adsorbiert, mit reduziertem Antigengehalt)				
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest- (acellulair, component) en poliomyelitis- (geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest- (acellulair, component) en poliomyelitis- (geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest- (acellulair, component) en poliomyelitis- (geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest-	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	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
(acellulair, component) en poliomyelitis-(geïnactiveerd) vaccin (geadsorbeerd)				
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest-(acellulair, component) en poliomyelitis-(geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest-(acellulair, component) en poliomyelitis-(geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactif) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest-(acellulair, component) en poliomyelitis-(geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	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
Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)				
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	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
Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)			INDUSTRIE	
Tetravac, Injektionssuspension in einer Fertigspritze Diphtherie, Tetanus, Pertussis (mit azellulärer Komponente) und Poliomyelitis (inaktiviert) Impfstoff (adsorbiert)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactivé) (adsorbé)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	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
Tetravac, suspensie voor injectie in een voorgevulde spuit Difterie-, tetanus-, kinkhoest- (acellulair, component) en poliomyelitis- (geïnactiveerd) vaccin (geadsorbeerd)	SE/H/0154/001	BE196122	SANOFI WINTHROP INDUSTRIE	BE
Boostrix Polio, suspension injectable en seringue pré-remplie Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélique (inactivé)	DE/H/0466/003	BE270435	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Boostrix Polio, Injektionssuspension in einer Fertigspritze Impfstoff (adsorbiert, Inhalt antigenreduziert) gegen Diphtherie, Tetanus, Pertussis (azellulär) und Poliomyelitis (inaktiviert)	DE/H/0466/003	BE270435	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
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélique inactivé (contenu	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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
réduit en antigène(s) adsorbé(s))				
<p>TRIAXIS POLIO</p> <p>Injektionssuspension in einer Fertigspritze</p> <p>Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)</p>	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
<p>TRIAXIS POLIO</p> <p>Injektionssuspension in einer Fertigspritze</p> <p>Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)</p>	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
<p>TRIAXIS POLIO</p> <p>Injektionssuspension in einer Fertigspritze</p> <p>Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)</p>	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
<p>TRIAXIS POLIO</p> <p>Injektionssuspension in einer Fertigspritze</p> <p>Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)</p>	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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
Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)			INDUSTRIE	
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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
(geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)				
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/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique,	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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
coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))				
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/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO, suspensie voor injectie in voorgevulde spuit. Difterie, Tetanus, Pertussis (acellulaire, component) en Poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerd antigeengehalte)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
TRIAXIS POLIO Injektionssuspension in einer Fertigspritze	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	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
Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)				
TRIAxis POLIO Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	BE241096	SANOFI WINTHROP INDUSTRIE	BE
Infanrix-IPV, 0,5 ml/Dosis Injizierbare Suspension	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Infanrix-IPV, 0,5 ml/dosis, suspensie voor injectie	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	BE
Infanrix-IPV, 0,5 ml/dose, suspension injectable	not available	BE212152	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG
Бустрикс Полио инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	DE/H/0466/003	20120314	GLAXOSMITHKLINE BIOLOGICALS S.A.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
полиомиелит (инактивирана)				
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	BG
ТЕТРАКСИМ 0,5 ml инжекционна суспензия Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и полиомиелит (инактивирана)	not available	20060041	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	BG
АДАСЕЛ ПОЛИО инжекционна суспензия в предварително напълнена спринцовка Адсорбирана ваксина срещу дифтерия, тетанус, коклюш (безклетъчна, компонентна) и	DE/H/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
полиомиелит (инактивирана), с намалено антигенно съдържание/				
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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	20200008	SANOFI WINTHROP INDUSTRIE	BG
TetraVac, ενέσιμο εναιώρημα σε	HU/H/0406/001	022511	SANOFI WINTHROP	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
προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).			INDUSTRIE	
Tetranas, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Tetranas, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Tetranas, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Tetranas, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Tetranas, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	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
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσοροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσοροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	HU/H/0406/001	022511	SANOFI WINTHROP INDUSTRIE	CY
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσοροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένης).	FR/H/0251/002	20232	GLAXOSMITHKLINE BIOLOGICALS S.A.	CY
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσοροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένης).	FR/H/0251/002	20232	GLAXOSMITHKLINE BIOLOGICALS S.A.	CY
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσοροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένης).	FR/H/0251/002	20232	GLAXOSMITHKLINE BIOLOGICALS S.A.	CY
Infanrix Tetra, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσοροφημένο) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας	FR/H/0251/002	20232	GLAXOSMITHKLINE BIOLOGICALS S.A.	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 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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	CZ
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 WINTHROP INDUSTRIE	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
poliomyelitidě (inaktivovaná).				
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 WINTHROP INDUSTRIE	CZ
Boostrix Polio injekční suspenze v předplněné injekční stříkačce Adsorbovaná vakcína proti difterii, tetanu, pertusi (acelulární komponenta) a poliomyelitide (inaktivovaná vakcína) se sníženým obsahem antigenu.	DE/H/0466/003	59/497/07-C	GLAXOSMITHKLINE S.R.O.	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	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
(adsorbovaná, se sníženým obsahem antigenů)				
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Adacel Polio injekční suspenze v předplněné injekční stříkačce Vakcína proti difterii, tetanu, pertusi (acelulární) a poliomyelitidě (inaktivovaná), (adsorbovaná, se sníženým obsahem antigenů)	DE/H/0215/001-002	59/265/19-C	SANOFI WINTHROP INDUSTRIE	CZ
Boostrix Polio Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis (azellulär, aus Komponenten)- und Poliomyelitis (inaktiviert)-Adsorbatimpfstoff (mit reduziertem Antigengehalt)	DE/H/0466/003	PEI.H.02950.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE
Boostrix Polio Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis (azellulär, aus Komponenten)- und Poliomyelitis (inaktiviert)-Adsorbatimpfstoff (mit reduziertem Antigengehalt)	DE/H/0466/003	PEI.H.02950.01.1	GLAXOSMITHKLINE GMBH & CO. KG	DE
Boostrix Polio Injektionssuspension in einer Fertigspritze Diphtherie-, Tetanus-, Pertussis (azellulär, aus	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
Komponenten)- und Poliomyelitis (inaktiviert)-Adsorbatimpfstoff (mit reduziertem Antigengehalt)				
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	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
Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)				
Repevax Injektionssuspension in einer Fertigspritze Diphtherie-Tetanus-Pertussis(azellulär, aus Komponenten)-Poliomyelitis(inaktiviert)-Impfstoff (adsorbiert, mit reduziertem Antigengehalt)	DE/H/0215/001-002	PEI.H.02354.01.1	SANOFI WINTHROP INDUSTRIE	DE
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i	SE/H/0154/001	30186	SANOFI WINTHROP	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
fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).			INDUSTRIE	
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Tetravac Injektionsvæske, suspension i fyldt injektionssprøjte. Difteri-, tetanus-, pertussis- (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet).	SE/H/0154/001	30186	SANOFI WINTHROP INDUSTRIE	DK
Boostrix Polio, injektionsvæske, suspension, fyldt injektionssprøjte	DE/H/0466/003	49761	GLAXOSMITHKLINE PHARMA A/S	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	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
(adsorberet, reduceret indhold af antigen(er)).				
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Repevax, injektionsvæske, suspension i fyldt injektionssprøjte. Difteri, tetanus, pertussis (acellulær, komponent) og poliomyelitis (inaktiveret) vaccine (adsorberet, reduceret indhold af antigen(er)).	DE/H/0215/001-002	33442	SANOFI WINTHROP INDUSTRIE	DK
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja	not available	430704	SANOFI WINTHROP INDUSTRIE	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)				
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE
Tetraxim, süstesuspensioon süstlis Difteeria, teetanuse, läkaköha (atsellulaarne komponent) ja poliomüelliidi (inaktiveeritud) vaktsiin (adsorbeeritud)	not available	430704	SANOFI WINTHROP INDUSTRIE	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
poliomüeliidi (inaktiveeritud) vaktsiin (adsorbeeritud)				
Infanrix Polio, süstesuspensioon süstlis. Difteria, teetanuse, läkaköha (atsellulaarne komponentvaktsiin) ja poliomüeliidi (inaktiveeritud) vaktsiin (adsorbeeritud).	FR/H/0251/002	483205	GLAXOSMITHKLINE BIOLOGICALS S.A.	EE
Infanrix Polio, süstesuspensioon süstlis. Difteria, teetanuse, läkaköha (atsellulaarne komponentvaktsiin) ja poliomüeliidi (inaktiveeritud) vaktsiin (adsorbeeritud).	FR/H/0251/002	483205	GLAXOSMITHKLINE BIOLOGICALS S.A.	EE
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomielititis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomielititis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomielititis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomielititis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	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
(componente acelular) y poliomieltis (inactivada) (adsorbida).				
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomieltis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
TETRAXIM, suspensión inyectable en jeringa precargada, Vacuna de difteria, tétanos, tos ferina (componente acelular) y poliomieltis (inactivada) (adsorbida).	HU/H/0406/001	85337	SANOFI WINTHROP INDUSTRIE	ES
Boostrix Polio suspensión inyectable en jeringa precargada Vacuna antidiftérica, antitetánica, antitos ferina (componente acelular), antipoliomieltis (inactivada) (adsorbida, contenido antigénico reducido)	DE/H/0466/003	66.462	GLAXOSMITHKLINE, S.A.	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada. Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomieltis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada. Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomieltis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada. Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomieltis (inactivada) (adsorbida, contenido de antígeno(s)	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	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
reducido).				
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomiелitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomiелitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomiелitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
TRIAXIS POLIO, suspensión inyectable, en jeringa precargada.Vacuna de difteria, tétanos, pertussis (componente acelular) y poliomiелitis (inactivada) (adsorbida, contenido de antígeno(s) reducido).	DE/H/0215/001-002	84.886	SANOFI WINTHROP INDUSTRIE	ES
Infanrix-IPV, suspensión inyectable en jeringa precargada Vacuna (adsorbida) antidiftérica, antitetánica, antipertussis (componente acelular) y antipoliomiелítica (inactivada)	FR/H/0251/002	85.671	GLAXOSMITHKLINE, S.A.	ES
Infanrix-IPV, suspensión inyectable en jeringa precargada Vacuna (adsorbida) antidiftérica, antitetánica, antipertussis (componente	FR/H/0251/002	85.671	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
acelular) y antipoliomiélfica (inactivada)				
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	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
(soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)				
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	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
Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.				
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Tetravac, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu)	SE/H/0154/001	13676	SANOFI WINTHROP INDUSTRIE	FI
Boostrix Polio, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och poliomyelit (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0466/003	19479	GLAXOSMITHKLINE BIOLOGICALS S.A.	FI
Boostrix Polio, injektioneste, suspensio	DE/H/0466/003	19479	GLAXOSMITHKLINE	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
esitäytetyssä ruiskussa Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)			BIOLOGICALS S.A.	
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	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
(inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)				
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektioneste, suspensio,	DE/H/0215/001-002	17190	SANOFI WINTHROP	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
esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)			INDUSTRIE	
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektioneste, suspensio, esitäytetty ruisku Kurkkumätä, jäykkäkouristus, hinkuyskä (soluton, komponentti) ja polio (inaktivoitu) -rokote (adsorboitu, vähennetty antigeeni(en) pitoisuus)	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	FI
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll	DE/H/0215/001-002	17190	SANOFI WINTHROP INDUSTRIE	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
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et	not available	34009 302 801 8 8	SANOFI WINTHROP INDUSTRIE	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
poliomyéлитique (inactivé), adsorbé				
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 302 801 9 5	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 347 823 9 8	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 391 896 8 0	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 348 222 9 2	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 359 667 7 3	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 359 668 3 4	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et	not available	34009 368 751 7 3	SANOFI WINTHROP INDUSTRIE	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
poliomyéлитique (inactivé), adsorbé				
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 368 752 3 4	SANOFI WINTHROP INDUSTRIE	FR
TETRAXIM, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 391 897 4 1	SANOFI WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 360 647 6 8	SANOFI WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 348 221 2 4	SANOFI WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 360 644 7 8	SANOFI WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 360 645 3 9	SANOFI WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyéлитique (inactivé), adsorbé	not available	34009 348 223 5 3	SANOFI WINTHROP INDUSTRIE	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				
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 351 668 4 5	SANOFI WINTHROP INDUSTRIE	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 WINTHROP INDUSTRIE	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 WINTHROP INDUSTRIE	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 WINTHROP INDUSTRIE	FR
TETRAVAC-ACELLULAIRE, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire et poliomyélitique (inactivé), adsorbé	not available	34009 368 750 0 5	SANOFI WINTHROP INDUSTRIE	FR
BOOSTRIXETETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	6 240 479 3	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXETETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique,	DE/H/0466/003	6 240 479 3	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
coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)				
BOOSTRIXETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	6 240 479 3	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	6 240 479 3	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	6 240 479 3	LABORATOIRE GLAXOSMITHKLINE	FR
BOOSTRIXETRA, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en antigènes)	DE/H/0466/003	6 240 479 3	LABORATOIRE GLAXOSMITHKLINE	FR
REPEVAX, suspension injectable en seringue préremplie. Vaccin diphtérique, tétanique, coquelucheux (acellulaire, multicomposé) et poliomyélitique (inactivé), (adsorbé, à teneur réduite en	DE/H/0215/001-002	34009 368 740 5 3	SANOFI WINTHROP INDUSTRIE	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
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/0215/001-002	34009 368 739 7 1	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	34009 368 738 0 3	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	34009 368 737 4 2	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	34009 359 644 7 2	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	34009 359 643 0 4	SANOFI WINTHROP INDUSTRIE	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
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/0215/001-002	34009 359-642 4 3	SANOFI WINTHROP INDUSTRIE	FR
INFANRIXTETRA, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux acellulaire, poliomyélitique inactivé, adsorbé	FR/H/0251/002	6 977 770 6	LABORATOIRE GLAXOSMITHKLINE	FR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό,	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	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
συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).				
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Tetranac, ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Προσροφημένο εμβόλιο διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό, συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	SE/H/0154/001	12029/31-1-2025	SANOFI WINTHROP INDUSTRIE	GR
Boostrix Polio ενέσιμο εναιώρημα σε προγεμισμένη σύριγγα Εμβόλιο (προσροφημένο, μειωμένο περιεχόμενο σε αντιγόνο(α) διφθερίτιδας, τετάνου, κοκκύτη (ακυτταρικό συστατικό) και πολιομυελίτιδας (αδρανοποιημένο).	DE/H/0466/003	2650601	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	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
Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))				
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	GR
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	GR
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	GR
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	GR
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	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
Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))				
REPEVAX, ενέσιμο εναιώρημα, σε προγεμισμένη σύριγγα Εμβόλιο Διφθερίτιδας, Τετάνου, Κοκκύτη (ακυτταρικό, συστατικό) και Πολιομυελίτιδας (αδρανοποιημένο), (προσροφημένο, μειωμένης περιεκτικότητας αντιγόνου (ων))	DE/H/0215/001-002	1041/13-01-2025	SANOFI WINTHROP INDUSTRIE	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
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
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
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
TETRAXIM, suspenzija za injekciju u napunjenoj štrcaljki	HU/H/0406/001	HR-H-185201369-07	SANOFI WINTHROP INDUSTRIE	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano), adsorbirano				
TETRAXIM, suspenzija za injekciju u napunjenoj štrcaljki cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano), adsorbirano	HU/H/0406/001	HR-H-185201369-08	SANOFI WINTHROP INDUSTRIE	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-185201369-06	SANOFI WINTHROP INDUSTRIE	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-185201369-01	SANOFI WINTHROP INDUSTRIE	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-185201369-02	SANOFI WINTHROP INDUSTRIE	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-185201369-05	SANOFI WINTHROP INDUSTRIE	HR
TETRAXIM, suspenzija za injekciju u	HU/H/0406/001	HR-H-185201369-03	SANOFI WINTHROP	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
napunjenoj štrcaljki cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano), adsorbirano			INDUSTRIE	
TETRAXIM, suspenzija za injekciju u napunjenoj štrcaljki cjepivo protiv difterije, tetanusa, hripavca (nestanično, komponentno) i poliomijelitisa (inaktivirano), adsorbirano	HU/H/0406/001	HR-H-185201369-04	SANOFI WINTHROP INDUSTRIE	HR
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/08	SANOFI WINTHROP INDUSTRIE	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/06	SANOFI WINTHROP INDUSTRIE	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/07	SANOFI WINTHROP INDUSTRIE	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/01	SANOFI WINTHROP INDUSTRIE	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
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/02	SANOFI WINTHROP INDUSTRIE	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/03	SANOFI WINTHROP INDUSTRIE	HU
Tetraxim szuszpenziós injekció előretöltött fecskendőben diftéria, tetanusz, pertusszisz (sejtmentes komponens) és poliomielitisz (inaktivált) vakcina (adszorbeált)	HU/H/0406/001	OGYI-T-9452/04	SANOFI WINTHROP INDUSTRIE	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 WINTHROP INDUSTRIE	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 BIOLOGICALS S.A.	HU
Boostrix Polio szuszpenziós injekció előretöltött fecskendőben diphtheria, tetanus, pertussis (acelluláris összetevő) és (inaktivált) poliomyelitis vakcina (adszorbeált, csökkentett antigén	DE/H/0466/003	OGYI-T-20497/03	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
tartalmú)				
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 BIOLOGICALS S.A.	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-20497/07	GLAXOSMITHKLINE BIOLOGICALS S.A.	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 BIOLOGICALS S.A.	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 BIOLOGICALS S.A.	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertussisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/01	SANOFI WINTHROP INDUSTRIE	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertussisz (sejtmentes alegység) és poliomielitisz (inaktivált)	DE/H/0215/001-002	OGYI-T-23632/02	SANOFI WINTHROP INDUSTRIE	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
vakcina (adszorbeált, csökkentett antigéntartalmú)				
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/03	SANOFI WINTHROP INDUSTRIE	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/04	SANOFI WINTHROP INDUSTRIE	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/05	SANOFI WINTHROP INDUSTRIE	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/06	SANOFI WINTHROP INDUSTRIE	HU
ADACEL POLIO szuszpenziós injekció előretöltött fecskendőben Diftéria, tetanusz, pertusszisz (sejtmentes alegység) és poliomielitisz (inaktivált) vakcina (adszorbeált, csökkentett antigéntartalmú)	DE/H/0215/001-002	OGYI-T-23632/07	SANOFI WINTHROP INDUSTRIE	HU
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis,	FR/H/0251/002	OGYI-T-20352/02	GLAXOSMITHKLINE BIOLOGICALS S.A.	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 poliomyelitis adszorbeált vakcina				
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina	FR/H/0251/002	OGYI-T-20352/05	GLAXOSMITHKLINE BIOLOGICALS S.A.	HU
Infanrix IPV szuszpenziós injekció előretöltött fecskendőben Kombinált diphtheria, tetanus, acellularis pertussis, inaktivált poliomyelitis adszorbeált vakcina	FR/H/0251/002	OGYI-T-20352/06	GLAXOSMITHKLINE BIOLOGICALS S.A.	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 BIOLOGICALS S.A.	HU
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	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
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine, (adsorbed).	SE/H/0154/001	PA 23458/007/001	SANOFI WINTHROP INDUSTRIE	IE
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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
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
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
REPEVAX, suspension for injection in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	IE
REPEVAX, suspension for injection in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed,	DE/H/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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
reduced antigen(s) content)				
REPEVAX, suspension for injection in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PA 23458/004/001	SANOFI WINTHROP INDUSTRIE	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
IPV Infanrix, suspension for injection in	FR/H/0251/002	PA 1077/108/001	GLAXOSMITHKLINE	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
pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed)			(IRELAND) LIMITED	
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
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
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
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	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
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkrampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkrampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkrampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Tetravac, stungulyf, dreifa. Barnaveiki-, stífkrampa-, kíghósta (frumulaus þáttur)- og aðsogað mænusóttarveirubóluefni (dautt).	SE/H/0154/001	970045	SANOFI WINTHROP INDUSTRIE	IS
Boostrix Polio stungulyf, dreifa í áfylltri sprautu Bóluefni gegn barnaveiki, stífkrampa, kíghósta (frumulaust, hlutar) og mænusótt (deyðdar veirur), (aðsogað, skert mótefnavakainnihald)	DE/H/0466/003	IS/1/07/027/01	GLAXOSMITHKLINE PHARMA A/S	IS
Boostrix Polio stungulyf, dreifa í áfylltri sprautu Bóluefni gegn barnaveiki, stífkrampa, kíghósta (frumulaust, hlutar) og mænusótt (deyðdar veirur), (aðsogað, skert mótefnavakainnihald)	DE/H/0466/003	IS/1/07/027/01	GLAXOSMITHKLINE PHARMA A/S	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkrampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	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
mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).				
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
REPEVAX stungulyf, dreifa í áfylltri sprautu Bóluefni (aðsogað, minnkað mótefnisvakainnihald) gegn barnaveiki, stífkampa, kíghósta (frumulaust, samsett) og mænusótt (dautt).	DE/H/0215/001-002	IS/1/02/020/01	SANOFI WINTHROP INDUSTRIE	IS
Tetravac sospensione iniettabile in siringa preriempita	SE/H/0154/001	034127011	SANOFI WINTHROP INDUSTRIE	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 in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127023	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127035	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127047	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127062	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127050	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127074	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita	SE/H/0154/001	034127086	SANOFI WINTHROP INDUSTRIE	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 in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127098	SANOFI WINTHROP INDUSTRIE	IT
Tetravac sospensione iniettabile in siringa preriempita Vaccino (adsorbito) antidifterico, antitetanico, antipertossico (componente acellulare) e antipolio (inattivato).	SE/H/0154/001	034127100	SANOFI WINTHROP INDUSTRIE	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,	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
pertossico (componente acellulare) e 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
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001-002	048290011	SANOFI WINTHROP INDUSTRIE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001-002	048290023	SANOFI WINTHROP INDUSTRIE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001-002	048290035	SANOFI WINTHROP INDUSTRIE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico,	DE/H/0215/001-002	048290047	SANOFI WINTHROP INDUSTRIE	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
antipertossico (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/0215/001-002	048290050	SANOFI WINTHROP INDUSTRIE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001-002	048290062	SANOFI WINTHROP INDUSTRIE	IT
TRIAXIS POLIO, sospensione iniettabile in siringa pre-riempita. Vaccino antidifterico, antitetanico, antipertossico (componenti acellulari) e antipoliomielitico (inattivato) (adsorbito, contenuto antigenico ridotto).	DE/H/0215/001-002	048290074	SANOFI WINTHROP INDUSTRIE	IT
PolioInfranix – 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
PolioInfranix – 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
PolioInfranix – sospensione iniettabile in siringa preriempita Vaccino (adsorbito) anti difterico, tetanico, pertossico (componente acellulare) e anti-	FR/H/0251/002	037157070	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
poliomielite (inattivato).				
PolioInfranix – 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
PolioInfranix – 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
TETRAXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir poliomieliito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/001	SANOFI WINTHROP INDUSTRIE	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir poliomieliito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/002	SANOFI WINTHROP INDUSTRIE	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir poliomieliito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/003	SANOFI WINTHROP INDUSTRIE	LT
TETRAXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (neląstelinė, komponentinė) ir poliomieliito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/004	SANOFI WINTHROP INDUSTRIE	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
TETRIXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelastelinė, komponentinė) ir poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/005	SANOFI WINTHROP INDUSTRIE	LT
TETRIXIM injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelastelinė, komponentinė) ir poliomielito (inaktyvuota), (adsorbuota)	not available	LT/1/99/2298/006	SANOFI WINTHROP INDUSTRIE	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelastelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/002	GLAXOSMITHKLINE BIOLOGICALS S.A.	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelastelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/005	GLAXOSMITHKLINE BIOLOGICALS S.A.	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelastelinė, komponentinė) ir nuo poliomielito (inaktyvuota), (adsorbuota, sumažinto antigenų kiekio)	DE/H/0466/003	LT/1/08/0971/006	GLAXOSMITHKLINE BIOLOGICALS S.A.	LT
BOOSTRIX POLIO injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos, stabligės, kokliušo (nelasteline, komponentine) ir nuo poliomielito (inaktyvuota), (adsorbuota,	DE/H/0466/003	LT/1/08/0971/001	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
sumažinto antigenų kiekio)				
Infanrix Polio injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos (D), stabligės (T), kokliušo (nelastelinė, komponentinė) (Pa) ir nuo poliomiélito (inaktyvuota) (IPV), (adsorbuota)	FR/H/0251/002	LT/1/05/0301/003	GLAXOSMITHKLINE BIOLOGICALS S.A.	LT
Infanrix Polio injekcinė suspensija užpildytame švirkšte Vakcina nuo difterijos (D), stabligės (T), kokliušo (nelastelinė, komponentinė) (Pa) ir nuo poliomiélito (inaktyvuota) (IPV), (adsorbuota)	FR/H/0251/002	LT/1/05/0301/001	GLAXOSMITHKLINE BIOLOGICALS S.A.	LT
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactif) (adsorbé)	SE/H/0154/001	0432975	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactif) (adsorbé)	SE/H/0154/001	0433003	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique (inactif) (adsorbé)	SE/H/0154/001	0432961	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant)	SE/H/0154/001	0432992	SANOFI WINTHROP INDUSTRIE	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
et poliomyélique (inactivé) (adsorbé)				
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélique (inactivé) (adsorbé)	SE/H/0154/001	0432989	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélique (inactivé) (adsorbé)	SE/H/0154/001	0434631	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélique (inactivé) (adsorbé)	SE/H/0154/001	0253876	SANOFI WINTHROP INDUSTRIE	LU
Tetravac, suspension injectable en seringue préremplie Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélique (inactivé) (adsorbé)	SE/H/0154/001	0434627	SANOFI WINTHROP INDUSTRIE	LU
Infanrix-IPV, 0,5 ml/Dosis Injizierbare Suspension	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Infanrix-IPV, 0,5 ml/dose, suspension injectable	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Infanrix-IPV, 0,5 ml/Dosis Injizierbare Suspension Impfstoff gegen Diphtherie, Tetanus, Keuchhusten und Poliomyelitis	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
Infanrix-IPV, 0,5 ml/dose, suspension injectable	not available	2007069308	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
Boostrix Polio, suspension injectable en seringue préremplie Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélitique (inactive)	DE/H/0466/003	2009020265	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
Boostrix Polio, suspension injectable en seringue préremplie Vaccin (adsorbé, contenu réduit en antigènes) diphtérique, tétanique, coquelucheux (composant acellulaire) et poliomyélitique (inactive)	DE/H/0466/003	2009020265	GLAXOSMITHKLINE BIOLOGICALS S.A.	LU
Boostrix Polio, Injektionssuspension in einer Fertigspritze Impfstoff (adsorbiert, Inhalt antigenreduziert) gegen Diphtherie, Tetanus, Pertussis (azellulär) und Poliomyelitis (inaktiviert)	DE/H/0466/003	2009020265	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))	DE/H/0215/001-002	0433048	SANOFI WINTHROP INDUSTRIE	LU
TRIAXIS POLIO, suspension injectable en seringue préremplie	DE/H/0215/001-002	0433034	SANOFI WINTHROP INDUSTRIE	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
Vaccin diphtérique, tétanique, coquelucheux (acellulaire, composant) et poliomyélitique inactivé (contenu réduit en antigène(s) adsorbé(s))				
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/0215/001-002	0433021	SANOFI WINTHROP INDUSTRIE	LU
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/0215/001-002	0433017	SANOFI WINTHROP INDUSTRIE	LU
TETRAXIM suspensija injekcijai pilnšjircē Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivētā poliomiēlīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšjircē Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivētā poliomiēlīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšlirce Adsorbēta difterijas, stingumkrampju, acelulārā gara klepus un inaktivēta poliomiēlīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšjircē Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivētā poliomiēlīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšjircē Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivētā poliomiēlīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	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
poliomielīta vakcīna.				
TETRAXIM suspensija injekcijai pilnšīrce Adsorbēta difterijas, stingumkrampju, acelulāra gara klepus un inaktivēta poliomielīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšīrce Adsorbēta difterijas, stingumkrampju, acelulāra gara klepus un inaktivēta poliomielīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
TETRAXIM suspensija injekcijai pilnšīrce Adsorbēta difterijas, stingumkrampju, acelulārā garā klepus un inaktivēta poliomielīta vakcīna.	not available	02-0343	SANOFI WINTHROP INDUSTRIE	LV
Boostrix Polio suspensija injekcijām pilnšīrce Difērijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna (adsorbēta, ar samazinātu antigēnu saturu)	DE/H/0466/003	07-0238	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Boostrix Polio suspensija injekcijām pilnšīrce Difērijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna (adsorbēta, ar samazinātu antigēnu saturu)	DE/H/0466/003	07-0238	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Boostrix Polio suspensija injekcijām pilnšīrce Difērijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna (adsorbēta, ar samazinātu antigēnu saturu)	DE/H/0466/003	07-0238	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Boostrix Polio suspensija injekcijām pilnšīrce Difērijas, stingumkrampju, garā klepus (acelulārais komponents) un poliomielīta (inaktivēta) vakcīna	DE/H/0466/003	07-0238	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
(adsorbēta, ar samazinātu antigēnu saturu)				
Infanrix Polio suspensija injekcijām pilnšjircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomiēlīta (inaktivēta) vakcīna (adsorbēta) Vaccinum diphtheriae, tetani, pertussis sine cellulis ex elementis preparatum et poliomyelitidis inactivatum adsorbatum	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Infanrix Polio suspensija injekcijām pilnšjircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomiēlīta (inaktivēta) vakcīna (adsorbēta) Vaccinum diphtheriae, tetani, pertussis sine cellulis ex elementis preparatum et poliomyelitidis inactivatum adsorbatum	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Infanrix Polio suspensija injekcijām pilnšjircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomiēlīta (inaktivēta) vakcīna (adsorbēta) Vaccinum diphtheriae, tetani, pertussis sine cellulis ex elementis preparatum et poliomyelitidis inactivatum adsorbatum	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
Infanrix Polio suspensija injekcijām pilnšjircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomiēlīta (inaktivēta) vakcīna (adsorbēta) Vaccinum diphtheriae, tetani, pertussis sine cellulis	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
ex elementis preparatum et poliomyelitidis inactivatum adsorbatum				
Infanrix Polio suspensija injekcijām pilnšjircē Difterijas, stingumkrampju, garā klepus (acelulāra komponenta) un poliomiēlīta (inaktivēta) vakcīna (adsorbēta) Vaccinum diphtheriae, tetani, pertussis sine cellulis ex elementis preparatum et poliomyelitidis inactivatum adsorbatum	FR/H/0251/002	05-0635	GLAXOSMITHKLINE BIOLOGICALS S.A.	LV
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	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
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	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
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	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
IPV-Boostrix suspension for injection in pre-filled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed,	DE/H/0466/003	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
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	MA170/00104	GLAXOSMITHKLINE BIOLOGICALS S.A.	MT
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerde antigeeninhoud)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerde antigeeninhoud)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerde antigeeninhoud)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin	DE/H/0466/003	RVG 35123	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
(geadsorbeerd, gereduceerde antigeeninhoud)				
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerde antigeeninhoud)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Boostrix Polio, suspensie voor injectie in voorgevulde spuit Difterie, tetanus, pertussis (acellulaire component) en poliomyelitis (geïnactiveerd) vaccin (geadsorbeerd, gereduceerde antigeeninhoud)	DE/H/0466/003	RVG 35123	GLAXOSMITHKLINE B.V.	NL
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i	SE/H/0154/001	21-14086	SANOFI WINTHROP	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
ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).			INDUSTRIE	
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	NO
Tetravac injeksjonsvæske, suspensjon i ferdigfylte sprøyter Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbert).	SE/H/0154/001	21-14086	SANOFI WINTHROP INDUSTRIE	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
REPEVAX suspensjon til injeksjon i en ferdigfylt sprøyte Vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert)	DE/H/0215/001	2002-01069	SANOFI WINTHROP INDUSTRIE	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
(adsorbert, redusert antigeninnhold)				
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	2002-01069	SANOFI WINTHROP INDUSTRIE	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	2002-01069	SANOFI WINTHROP INDUSTRIE	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	2002-01069	SANOFI WINTHROP INDUSTRIE	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	2002-01069	SANOFI WINTHROP INDUSTRIE	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	2002-01069	SANOFI WINTHROP INDUSTRIE	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	2002-01069	SANOFI WINTHROP INDUSTRIE	NO
Infanrix Polio injeksjonsvæske, suspensjon i ferdigfylt sprøyte vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt	FR/H/0251/002	05-3312	GLAXOSMITHKLINE AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(inaktivert), (adsorbent)				
Infanrix Polio injeksjonsvæske, suspensjon i ferdigfylt sprøyte vaksine mot difteri, tetanus, kikhoste (acellulær, komponent) og poliomyelitt (inaktivert), (adsorbent)	FR/H/0251/002	05-3312	GLAXOSMITHKLINE AS	NO
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	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
adsorbowana.				
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
TETRAXIM, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana.	HU/H/0406/001	23545	SANOFI WINTHROP INDUSTRIE	PL
Boostrix Polio, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0466/003	14241	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	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
Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów				
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	PL
ADACEL POLIO, zawiesina do wstrzykiwań w ampułko-strzykawce	DE/H/0215/001-002	25842	SANOFI WINTHROP INDUSTRIE	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
Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana, o zmniejszonej zawartości antygenów				
Infanrix-IPV, zawiesina do wstrzykiwań w ampułkostrzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana	FR/H/0251/002	11884	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Infanrix-IPV, zawiesina do wstrzykiwań w ampułkostrzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana	FR/H/0251/002	11884	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Infanrix-IPV, zawiesina do wstrzykiwań w ampułkostrzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana	FR/H/0251/002	11884	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Infanrix-IPV, zawiesina do wstrzykiwań w ampułkostrzykawce Szczepionka przeciw błonicy, tężcowi, krztuścowi (bezkomórkowa, złożona) i poliomyelitis (inaktywowana), adsorbowana	FR/H/0251/002	11884	GLAXOSMITHKLINE BIOLOGICALS S.A.	PL
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	5376389	SANOFI WINTHROP INDUSTRIE	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
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	2782480	SANOFI WINTHROP INDUSTRIE	PT
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	5458583	SANOFI WINTHROP INDUSTRIE	PT
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	5458781	SANOFI WINTHROP INDUSTRIE	PT
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	5376488	SANOFI WINTHROP INDUSTRIE	PT
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	2782589	SANOFI WINTHROP INDUSTRIE	PT
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular)	SE/H/0154/001	5458682	SANOFI WINTHROP INDUSTRIE	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
e a poliomielite (inativada).				
Tetravac, suspensão injetável em seringa pré-cheia Vacina (adsorvida) contra a difteria, o tétano, a tosse convulsa (componente acelular) e a poliomielite (inativada).	SE/H/0154/001	5458484	SANOFI WINTHROP INDUSTRIE	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 antígeno(s))	DE/H/0466/003	5212683	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 antígeno(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 antígeno(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 antígeno(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	DE/H/0466/003	5271085	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 antígeno(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 antígeno(s))	DE/H/0466/003	5271184	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	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
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	5835236	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	5835244	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-poliomielitic inactivat, adsorbit	not available	4490/2012/01	SANOFI WINTHROP INDUSTRIE	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelular-	not available	4490/2012/02	SANOFI WINTHROP INDUSTRIE	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
poliomielitice inactivate, adsorbite				
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelu- poliomielitice inactivate, adsorbite	not available	4490/2012/03	SANOFI WINTHROP INDUSTRIE	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelu- poliomielitice inactivate, adsorbite	not available	4490/2012/04	SANOFI WINTHROP INDUSTRIE	RO
TETRAXIM suspensie injectabilă în seringă preumplută Vaccin diftero-tetano-pertussis acelu- poliomielitice inactivate, adsorbite	not available	4490/2012/05	SANOFI WINTHROP INDUSTRIE	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelu- componente) și poliomielitice (inactivate) (adsorbite, cu conținut redus de antigen(e))	DE/H/0466/003	8900/2016/03	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelu- componente) și poliomielitice (inactivate) (adsorbite, cu conținut redus de antigen(e))	DE/H/0466/003	8900/2016/02	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelu- componente) și poliomielitice (inactivate) (adsorbite, cu conținut redus de antigen(e))	DE/H/0466/003	8900/2016/01	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelu- componente) și poliomielitice (inactivate)	DE/H/0466/003	8900/2016/04	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
(adsorbit, cu conținut redus de antigen(e))				
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat) (adsorbit, cu conținut redus de antigen(e))	DE/H/0466/003	8900/2016/05	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Boostrix-IPV suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat) (adsorbit, cu conținut redus de antigen(e))	DE/H/0466/003	8900/2016/06	GLAXOSMITHKLINE BIOLOGICALS S.A.	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/01	SANOFI WINTHROP INDUSTRIE	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/02	SANOFI WINTHROP INDUSTRIE	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/03	SANOFI WINTHROP INDUSTRIE	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis	DE/H/0215/001-002	15540/2024/04	SANOFI WINTHROP INDUSTRIE	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 poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))				
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/05	SANOFI WINTHROP INDUSTRIE	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/06	SANOFI WINTHROP INDUSTRIE	RO
Adacel Polio suspensie injectabilă în seringă preumplută Vaccin difteric, tetanic, pertussis (acelular, componente) și poliomieltic (inactivat), (adsorbit, cu conținut redus de antigen(e))	DE/H/0215/001-002	15540/2024/07	SANOFI WINTHROP INDUSTRIE	RO
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	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, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.				
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Tetravac, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, stelkramp och kikhosta, acellulärt, komponent samt mot polio, inaktiverat, adsorberat.	SE/H/0154/001	13464	SANOFI WINTHROP INDUSTRIE	SE
Boostrix Polio, injektionsvätska,	DE/H/0466/003	21068	GLAXOSMITHKLINE AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
suspension i förfylld spruta. Vaccin mot difteri, stelkramp, kikhosta (acellulärt, komponent) och polio (inaktiverat), adsorberat, med reducerat antigeninnehåll				
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
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
Repevax, injektionsvätska, suspension i förfylld spruta	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	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)				
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
Repevax, injektionsvätska, suspension i förfylld spruta Vaccin mot difteri, tetanus, pertussis (acellulärt, komponent) och (inaktiverat) poliomyelit (adsorberat, lägre antigenhalt)	DE/H/0215/001-002	59538	SANOFI WINTHROP INDUSTRIE	SE
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
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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
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
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
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 BIOLOGICALS S.A.	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/10/00651/002	GLAXOSMITHKLINE BIOLOGICALS S.A.	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/10/00651/003	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
antigenov)				
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskeemu kašlju – acelularno in otroški paralizi – inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/003	H/10/00651/004	GLAXOSMITHKLINE BIOLOGICALS S.A.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskeemu kašlju – acelularno in otroški paralizi – inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/003	H/10/00651/007	GLAXOSMITHKLINE BIOLOGICALS S.A.	SI
Boostrix Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi cepivo proti davici, tetanusu, oslovskeemu kašlju – acelularno in otroški paralizi – inaktivirano (adsorbirano, z zmanjšano vsebnostjo antigenov)	DE/H/0466/003	H/10/00651/008	GLAXOSMITHKLINE BIOLOGICALS S.A.	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/0215/001-002	H/20/02707/001	SANOFI WINTHROP INDUSTRIE	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed,	DE/H/0215/001-002	H/20/02707/002	SANOFI WINTHROP INDUSTRIE	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
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/0215/001-002	H/20/02707/003	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	H/20/02707/004	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	H/20/02707/006	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	H/20/02707/005	SANOFI WINTHROP INDUSTRIE	SI
Adacel Polio suspenzija za injiciranje v napolnjeni injekcijski brizgi Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed,	DE/H/0215/001-002	H/20/02707/007	SANOFI WINTHROP INDUSTRIE	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
reduced antigen(s) content)				
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	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
(acelulárna zložka), detskej obrne (inaktivovaná).				
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
TETRAXIM Injekčná suspenzia v naplnenej injekčnej striekačke Očkovacia látka (adsorbovaná) proti záškrtu, tetanu, čiernemu kašľu (acelulárna zložka), detskej obrne (inaktivovaná).	HU/H/0406/001	59/0291/16-S	SANOFI WINTHROP INDUSTRIE	SK
Boostrix Polio Injekčná suspenzia v naplnenej 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 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/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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	DE/H/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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
(inaktivovaná)				
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/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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	DE/H/0215/001-002	59/0035/20-S	SANOFI WINTHROP INDUSTRIE	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
(inaktivovaná)				
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	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
(inaktivovaná)				
Infanrix Polio injekčná suspenzia v naplnenej injekčnej striekacke Ockovacia látka (adsorbovaná) proti diftérii, tetanu, pertussis (acelulárna zložka) a poliomyelitíde (inaktivovaná)	FR/H/0251/002	59/0417/08-S	GLAXOSMITHKLINE BIOLOGICALS S.A.	SK
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetrvac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
component) and poliomyelitis (inactivated) vaccine (adsorbed).				
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	XI
Tetravac, suspension for injection in prefilled syringe Diphtheria, tetanus, pertussis (acellular, component) and poliomyelitis (inactivated) vaccine (adsorbed).	SE/H/0154/001	PL 23228/0004	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
REPEVAX, suspension for injection, in pre-filled syringe Diphtheria, Tetanus, Pertussis (acellular, component) and Poliomyelitis (inactivated) Vaccine (adsorbed, reduced antigen(s) content)	DE/H/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	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/0215/001-002	PL 23228/0009	SANOFI WINTHROP INDUSTRIE	XI