

This medicine protocol is a specific written instruction for the administration of Infanrix Hexa (6 in 1 vaccine), to refugees and applicants seeking protection in Ireland (hereafter referred to as vaccine recipients) and in the event of an outbreak by registered nurses and registered midwives. This medicine protocol is valid for the 2025/2026 Health Service Executive (HSE) Primary Childhood Immunisation Programme (PCIP), for National Immunisation Advisory Committee (NIAC) recommended catch-up vaccination and in the event of an outbreak as advised by public health. This medicine protocol enables registered nurses and registered midwives employed in the voluntary and statutory services of the HSE including HSE community vaccination clinics who have undertaken the required education and training programmes to administer Infanrix Hexa (6 in 1 vaccine), with reference to and guidance from Nursing & Midwifery Board of Ireland (NMBI), National Immunisation Advisory Committee (NIAC), National Immunisation Office (NIO), HSE and in accordance with the Summary of Product Characteristics (SmPC) for Infanrix Hexa (6 in 1 vaccine) as detailed by the European Medicines Agency (EMA) at www.ema.ie

- An Bord Altranais (2007) Guidance to Nurses and Midwives on Medication Management. Dublin: An Bord Altranais
- National Immunisation Advisory Committee (2023) Anaphylaxis: Immediate Management in the Community available at:https://www.higa.ie/areas-we-work/national-immunisation-advisory-committee/immunisationguidelines-ireland
- National Immunisation Advisory Committee Immunisation Guidelines for Ireland: National Immunisation Advisory Committee available at:https://www.higa.ie/areas-we-work/national-immunisation-advisorycommittee/immunisation-guidelines-ireland
- National Immunisation Office (2025)Children Who Have Come To Ireland From Another Country: Information For Healthcare Professionals On Catch-Up Vaccination available at https://www.hse.ie/eng/health/immunisation/hcpinfo/botpipa/hcwcatchupothercountries.pdf
- National Immunisation Office Supporting Information for Staff: Schools Immunisation Programme 2025/2026 available at: https://www.hse.ie/eng/health/immunisation/hcpinfo/schoolproghcp/supportingdoc.pdf
- National Immunisation Office (2025) Supporting Information for Vaccinations in General Practice available at: https://www.hse.ie/eng/health/immunisation/infomaterials/gpsupportingdocpci.pdf
- Nursing and Midwifery Board of Ireland (2025) Code of Professional Conduct and Ethics for Registered Nurses and Registered Midwives incorporating the Scope of Practice and Professional Guidance. Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/NMBI/media/NMBI/NMBI-Code-of-Professional-Conduct-and-Ethics.pdf?ext=.pdf
- Nursing and Midwifery Board of Ireland (2020) Guidance for Registered Nurses and Registered Midwives on Medication Administration. Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/NMBI/media/NMBI/NMBI-Medication-Administration-2020.pdf?ext=.pdf
- Nursing and Midwifery Board of Ireland (2022) Practice Standards for Midwives. Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/NMBI/media/NMBI/NMBI-Practice-Standards-for-Midwives.pdf?ext=.pdf

The Nursing and Midwifery Board of Ireland defines medicine protocols as "written directions that allow for the supply and administration of a named medicinal product by a registered nurse or midwife in identified clinical situations. A medication protocol involves the authorisation of the nurse or midwife to supply and administer a medication to groups of patients in a defined situation meeting specific criteria and who may not be individually identified before presentation for treatment (An Bord Altranais, 2007, page 37).



Document reference number	Version4-NIO-6in1-September 2025			
1.0 Critical elements				
Name of organisation/setting where medicine protocol applies	Health Service Providers across the voluntary and statutory services of the HSE including HSE community vaccination clinics. This Medicine Protocol applies to: • registered nurses and registered midwives employed in the voluntary and statutory services of the HSE including vaccination clinics, congregated settings, temporary clinics and mobile units			
Date the medicine protocol comes into effect	September 2025			
Date for review of medicine protocol	September 2026			
Document prepared by	HSE National Immunisation Office (NIO) in collaboration with the Office of the Nursing and Midwifery Services Director (ONMSD) HSE.			
Names and Signatures of the employing authority who is authorising the implementation of the medicine protocol	Name: Dr Éamonn O'Moore , Director of National Health Protection			
"On behalf of the authority employing professionals authorised to administer under this medicine protocol, I have read this medicine protocol and authorise its implementation"	Signature:			
	Name: Dr Colm Henry , Chief Clinical Officer, HSE			
	Signature:			
	Name: Dr Geraldine Shaw , Nursing and Midwifery Services Director, HSE			
	Signature:			



Clinical condition for use of the medicine protocol	The clinical condition for which this medicine protocol has been developed is for the active immunisation against and prevention of Diphtheria, Tetanus, Pertussis, Hepatitis B, Poliomyelitis and Haemophilus type b disease.		
Circumstances in which the medicine protocol applies	To provide an Infanrix Hexa (6 in 1 vaccine) to vaccine recipients in Ireland and in the event of an outbreak.		
Inclusion criteria for	For primary immunisation:		
vaccine recipients receiving Infanrix Hexa (6 in 1 vaccine) under this	Children born on or before 30 th September 2024: Children are recommended a primary course of 3 doses of Infanrix Hexa given at 2, 4 and 6 months.		
medicine protocol	Children born on or after the 1st of October 2024: These children are recommended a primary course of 3 doses of Infanrix Hexa (6 in 1 vaccine) given at 2, 4, and 6		
	months. A booster dose of Infanrix Hexa (6 in 1 vaccine) is recommended at 13 months for these children.		
	If the 3 rd dose of 6in1 vaccine due at the 6-month visit is delayed to 12 months or more, the third dose of 6 in 1 vaccine should be given at the 13 month-visit or as soon as possible.		
	The 4th 6in1, (which is recommended at 13 months in the schedule), should be replaced by the 4in1 vaccine (Tetravac). The 4in1 vaccine should be given after an interval of 6 months from the 3rd dose of the 6in1 vaccine has been administered. Please refer to the NIAC chapter 2.		
	For catch-up programme: Children <u>under 10 years of age</u> with no history of receipt of these antigens previously as per the NIAC catch up schedule in the NIAC chapter 2. For outbreak response: As advised by public health		
	Precautions		
	Acute severe febrile illness, defer until recovery.		
	 Type III (Arthus) hypersensitivity reaction to a previous dose (see adverse reactions in NIAC Chapter 6). Persons experiencing these reactions usually have very high serum diphtheria or tetanus antitoxin levels; they should not be given further routine or emergency booster doses of tetanus or diphtheria containing vaccines more frequently than every 10 years. Note: COVID-19 vaccines (except for COVID-19 vaccine given to children aged 6 months to 4 years where a 14-day interval is recommended) and other vaccines may be administered at the same time or at any interval. 		
Exclusion criteria for vaccine recipients receiving Infanrix Hexa (6 in 1 vaccine) under this medicine protocol	A known history of anaphylactic or hypersensitivity reaction to Infanrix Hexa (6 in 1 vaccine) or any of the vaccine's constituents.		
Actions to be taken for those who are excluded	All recipients meeting exclusion criteria must be referred to the clinical lead for an individual clinical assessment.		
from the medicine protocol	 Document action in clinical notes Where Infanrix Hexa (6 in 1 vaccine) is prescribed following clinical assessment, the registered nurse or midwife may administer Infanrix Hexa (6 in1 vaccine) within their scope of practice. 		



		scope of practice, registered nu mpetency to carry out a role or		
Description of circumstances and referral arrangements when further advice or consultation is required	Discuss the vaccine recipient with the clinical lead in the event of: • Previous adverse reaction • Other clinical concerns			
Documentation required for the implementation of this medicine protocol 3.0 Name of Medicine	A consent form must be understood and completed by the parent/legal guardian for all children who receive the Infanrix Hexa (6 in 1 vaccine). Translation of consent undertaken with support of translator if required. Appropriate details should be recorded on the consent form. The following documents will be required at each vaccination session: Vaccination session report form Blank vaccine consent forms Vaccine Information Leaflets Patient held record cards/ Vaccine passport Post vaccination advice should be provided It is the responsibility of each registered nurse or midwife to be familiar with the appropriate documentation to support the safe administration of Infanrix Hexa (6 in 1 vaccine) vaccine which includes the following: This medicine protocol NIAC (2023) Anaphylaxis: Immediate Management in the Community https://www.hiqa.ie/areas-we-work/national-immunisation-advisory-committee/immunisation-guidelines-ireland Infanrix hexa, Powder and suspension for suspension for injection. Diphtheria (D), tetanus (T), pertussis (acellular, component) (Pa), hepatitis B (rDNA) (HBV), poliomyelitis (inactivated) (IPV) and Haemophilus influenzae type b (Hib) conjugate vaccine (adsorbed). Dose: 0.5 ml			
	Route: Intramuscular injection			
	Patients Age	Site	Needle length & Size	
	Birth to <12 months	Vastus lateralis muscle	25 mm (Use a 16 mm needle in infants under 2.5 - 3 kg)	
			23-25 gauge	
	12 to <36 months	Vastus lateralis or deltoid muscle (depending on muscle mass)	25 mm 23-25 gauge	
	3 years -9 years	Deltoid muscle	25 mm 23-25 gauge	
	Note: Vaccine must be reconstituted			
Link to Medicine Details of product information and other	Link to Summary of Proc Characteristics: https://ww hexa-epar-product-informa	ww.ema.europa.eu/en/documen	ts/product-information/infanrix-	



nurse or registered midwife must remain with the vaccine recipient and closely monitor them for any adverse reactions. Vital signs should be recorded and the vaccine recipient should be reviewed by the registered nurse/midwife and/or medical practitioner. The incident must be reported to the relevant line manager as soon as possible. The incident and all actions taken must be promptly recorded in the recipient's documentation/notes and the relevant National Incident Management Report Form completed, available at: https://www.hse.le/eng/about/who/napsd/qps-incident-management/nims/nirf-01-v12-person-interactive.pdf The child's parent and/or legal guardian must be informed of the incident. Any suspected adverse reactions associated with medicine errors should be reported to the HPRA as outlined below. Any errors and near misses not involving medications/not directly involving the vaccine recipient (needle stick injuries etc.), the incident and all actions taken must be promptly recorded on the relevant National Incident Management Report Form and forwarded to the relevant line manager as per local policy. Refer to https://www.hpsc.ie/a-z/emi/algorithms/EMISharpsAlgo.pdf Procedure for reporting Adverse Drug Reactions to the HPRA unursing or midwifery staff should report to the HPRA any suspected adverse reactions, in accordance with criteria outlined by the HPRA. This reporting may be carried out online at https://www.hpra.ie Infanrix Hexa (6 in 1 vaccine) Fridge/cool box with minimum/maximum temperature recording device to monitor the cold chain temperature (between +2°C and +8°C) Disposable kidney dishes/coloured trays Gauze swabs/plasters Sharps bins, and bags for disposal of healthcare risk and non-risk material Accobol hand sanitiser Handwashing facilities Access to telephone	data including instructions for supply and administration is available from the Health Products Regulatory Authority at www.ema.ie	Link to Patient Information Leaflet: https://www.ema.europa.eu/en/documents/product-information/infanrix-hexa-epar-product-information_en.pdf	
adverse Drug Reactions to the HPRA adverse Practions, in accordance with criteria outlined by the HPRA. This reporting may be carried out online at https://www.hpra.ie Infanrix Hexa (6 in 1 vaccine) Infanrix Hexa (6 in 1 vaccine) Fridge/cool box with minimum/maximum temperature recording device to monitor the cold chain temperature (between +2°C and +8°C) Disposable kidney dishes/coloured trays Gauze swabs/plasters Sharps bins, and bags for disposal of healthcare risk and non-risk material Alcohol hand sanitiser Handwashing facilities Access to telephone Resuscitation equipment and drugs in accordance with the NIAC (2023) Anaphylaxis: Immediate management in the Community available at: https://rcpi.access.preservica.com/uncategorized/IO a36f9e4b-4c80-432d-826-546089359925/	reporting and documentation of errors and near misses involving the vaccine	medicine/patient/dose/route being administered or another medicine error, the registered nurse or registered midwife must remain with the vaccine recipient and closely monitor them for any adverse reactions. Vital signs should be recorded and the vaccine recipient should be reviewed by the registered nurse/midwife and/or medical practitioner. The incident must be reported to the relevant line manager as soon as possible. The incident and all actions taken must be promptly recorded in the recipient's documentation/notes and the relevant National Incident Management Report Form completed, available at: https://www.hse.ie/eng/about/who/nqpsd/qps-incident-management/nims/nirf-01-v12-person-interactive.pdf The child's parent and/or legal guardian must be informed of the incident. Any suspected adverse reactions associated with medicine errors should be reported to the HPRA as outlined below. Any errors and near misses not involving medications/not directly involving the vaccine recipient (needle stick injuries etc.), the incident and all actions taken must be promptly recorded on the relevant National Incident Management Report Form and forwarded to the relevant line manager as per local policy. Refer to https://www.hpsc.ie/a-	
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Current medicine protocol for Infanrix Hexa (6 in 1 vaccine). Audit process to identify appropriate use of the medicine protocol or All documentation will be held for review and audit purposes as per local policy.	required for the administration of Infanrix Hexa (6in1) Vaccine Audit process to identify appropriate use of the	 Fridge/cool box with minimum/maximum temperature recording device to monitor the cold chain temperature (between +2°C and +8°C) Disposable kidney dishes/coloured trays Gauze swabs/plasters Sharps bins, and bags for disposal of healthcare risk and non-risk material Alcohol hand sanitiser Handwashing facilities Access to telephone Resuscitation equipment and drugs in accordance with the NIAC (2023) Anaphylaxis: Immediate management in the Community available at: https://rcpi.access.preservica.com/uncategorized/IO a36f9e4b-4c80-432d-8264-546089359925/ Safe storage areas for medicines and equipment Current medicine protocol for Infanrix Hexa (6 in 1 vaccine). 	



4.0 Information for vaccine recipient/parent/legal guardian

Advice to be given to the vaccine recipient before treatment

The information must be supplied with the consent form to each parent/legal guardian prior to administration of the vaccine. Reiterate the information provided in the HSE patient information leaflet and translator support if required.

Advice to be given to the vaccine recipient after treatment

After Treatment

The vaccine recipient must be advised to remain seated in the post vaccination observation area for 15 minutes to allow monitoring of any immediate reaction including possible anaphylactic reaction and must be advised to report any side effects to the registered nurse or registered midwife who is present.

Details of any necessary follow-up, action and referral arrangements

In the event of an adverse reaction the registered nurse or midwife must ensure that all procedures are adhered to as outlined in Section 3.

5.0 Staff authorised to use this medicine protocol

Professional qualifications, training, and competence required prior to using this medicine protocol

Registered nurse or registered midwife must have completed all of the following:

- 1) Be a Registered Nurse or Registered Midwife, on the active register maintained by the NMBI
- 2) Primary Childhood Immunisation Programme accessible on www.HSeLanD.ie
- 3) Education programme for nurses and midwives on National Vaccination Education Programme for Registered Nurses/Midwives/Public Health Nurses: 2025/2026 and any updates for nurses and midwives accessible on www.HSeLanD.ie
- 4) An approved Basic Life Support for Health Care Providers Course within the last two years (i.e. Irish Heart Foundation (IHF))
- 5) Initial National Anaphylaxis Education Programme for Health Care Professionals accessible on www.HSeLanD.ie followed by a two hour classroom based skills workshop. Recertification is required every two years by completing the on-line National Anaphylaxis Education Programme for Health Care Professionals accessible on www.HSeLanD.ie
- 6) Children having vaccinations and healthcare procedures: Clinical Holding (Professor Lucy Bray/ONMSD, 2023) available at www.HSeLanD.ie The registered nurse/midwife must complete the Competency Self-Assessment Form to vaccinate Refugees and Applicants Seeking Protection in Ireland available at www.immunisation.ie



References

An Bord Altranais (2007) Guidance to Nurses and Midwives on Medication Management Dublin: An Bord Altranais

Infanrix Hexa Summary of Product Characteristics and Patient Information Leaflet, available at: https://www.ema.europa.eu/en/medicines/human/EPAR/infanrix-hexa#product-information-section

HSE Policy on the Management of Sharps and Prevention of Sharp Injuries (2022) available at: https://healthservice.hse.ie/filelibrary/staff/policy-on-the-management- of-sharps-and-prevention-of-sharp-injuries.pdf

National Clinical Guideline No. 30 (2023) - Infection Prevention and Control (IPC) available at: https://www.gov.ie/en/publication/a057e-infection-prevention-and-control-ipc/.

National Immunisation Advisory Committee (2023) Anaphylaxis: Immediate Management in the Community available at: https://www.higa.ie/areas-we-work/national-immunisation-advisory-committee/immunisation-guidelines-ireland

National Immunisation Advisory Committee Immunisation Guidelines for Ireland: Royal College of Physicians of Ireland National Immunisation Advisory Committee available at: https://www.higa.ie/areas-we-work/nationalimmunisation-advisory-committee/immunisation-guidelines-ireland

National Immunisation Office Supporting Information for Staff: Schools Immunisation Programme 2025/2026 available at: https://www.hse.ie/eng/health/immunisation/hcpinfo/schoolproghcp/supportingdoc.pdf

National Immunisation Office (2025) Children Who Have Come To Ireland From Another Country: Information For Healthcare Professionals On Catch-Up Vaccination available at https://www.hse.ie/eng/health/immunisation/hcpinfo/botpipa/hcwcatchupothercountries.pdf

National Immunisation Office (2025) Supporting Information for Vaccinations in General Practice available at: https://www.hse.ie/eng/health/immunisation/infomaterials/gpsupportingdocpci.pdf

Nursing and Midwifery Board of Ireland (2025) Code of Professional Conduct and Ethics for Register Nurses and Registered Midwives incorporating the Scope of Practice and Professional Guidance. Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/NMBI/media/NMBI/NMBI-Code-of-Professional-Conduct-and-Ethics.pdf?ext=.pdf

Nursing and Midwifery Board of Ireland (2020) Guidance for Registered Nurses and Registered Midwives on Medication Administration. Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/NMBI/media/NMBI/NMBI-Medication-Administration-2020

Nursing and Midwifery Board of Ireland (2022) Practice Standards for Midwives Dublin: Nursing and Midwifery Board of Ireland available at: https://www.nmbi.ie/Standards-Guidance/Midwives-Standards