



10:55 am

Vaccination errors: your go-to guide

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Declarations



Advisory Board Member – Seqirus, Pfizer, GSK, Sanofi, MSD, Moderna

Honoraria received from the mentioned companies

Immunisation Coalition - Member

Immunisation Coordinator - Adelaide PHN

Learning Objectives



At the end of the session, participants will be able to:

- 1. Explain the steps to take when a vaccine program error occurs
- 2. Implement strategies to reduce the risk of vaccination errors

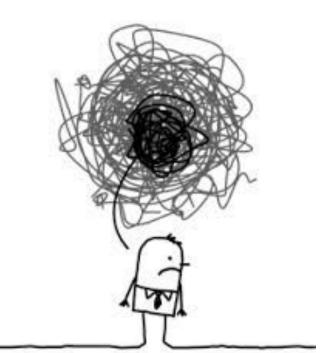


Why do vaccination errors occur?

I'M NOT A ROBOT



I'M HUMAN







Infanrix Infanrix hexa Infanrix IPV

HIB and HepB HIB and influenza HPV and HIV



What are the most common vaccination errors?

1. Skipping parts of the pre-vaccination process:

- The patient was not adequately identified
- The pre-vaccination checklist was not undertaken
- Risk and benefits were not explained
- VALID consent was not obtained
- The AIR was not checked, and other previous vaccination evidence was not sought





How to reduce the risk

- Ask the patient their full name
- Complete the pre-vaccination checklist and ask about medical conditions, allergies, pregnancy
- Ensure they know about possible adverse events, how to look for fever and manage fever in a child, how to report an AEFI
- Follow valid consenting steps:
 - It must be given by a person with legal capacity, and of sufficient intellectual capacity to understand the implications of receiving a vaccine.
 - It must be given voluntarily in the absence of undue pressure, coercion or manipulation.
 - It must cover the specific procedure that is to be performed.
 - It can only be given after the potential risks and benefits of the relevant vaccine, the risks of not having it, and any alternative options have been explained to the person.



2. Vaccine administration errors:

- The wrong vaccine was selected
- The vaccine had expired
- The vaccine was given for the wrong reason (Tetanus vaccine)
- The vaccine was given at the wrong time (seasonal, pregnancy gestation)
- The minimum interval between vaccines was not observed
- The vaccine was not reconstituted, or the incorrect diluent was used
- The incorrect vaccine for their age as administered (DTPa or dTpa, paed Hep B/adult Hep B)
- The wrong dose was administered (COVID vaccines and multidose vials)
- The vaccine was administered via the wrong route....oral, sub-cut, IM
- The patient was not checked for immediate AEFI and was not recommend to wait 15-minutes post vaccination
- Poor documentation: Insufficient information in clinical record, not recorded on the AIR, take home record was not provided

How to reduce the risk?

- You have the right vaccine for the right reason, at the right time, and with the right interval (dTpa vaccine, pregnancy, at 20 weeks)
- The vaccine is not expired
- You reconstitute the vaccine if required using the correct diluent
- You have the right vaccine for their age (DTPa or dTpa, paed Hep B/adult Hep B, MMR or MMRV)
- You give the right dose (COVID vaccines and multidose vials)
- You administer via the right route....oral, sub-cut, IM
- You check for immediate AEFI and recommend 15-minute wait post vaccination
- You document correctly: Clinical record, AIR, take home record











3. Administering an inappropriately stored vaccine

- Temperatures not checked and recorded twice daily, and data logger not downloaded weekly
- Vaccines stored outside the manufacturer's recommended temperature range (usually +2°C to +8°C) and appropriate action not taken
 - Signage on the fridge: "Possible cold chain breach: Do not use vaccines until further notice"
 - Isolate affected vaccines (do not discard them and keep them refrigerated between +2°C to +8°C)
 - Download the fridge logger
 - Investigate cause (power failure, door left ajar, mechanical failure)
 - Notify State/Territory Health Department/Vaccine supplier and seek advice



How do I reduce the risk?

- Ensure all staff, clinical and non-clinical, have received vaccine cold chain training
- Ensure vaccine fridge is in good working order regular servicing
- Read temperatures twice daily, morning and night......document the readings on the temperature chart
- Download the logger weekly, or whenever there is suspicion of a cold chain excursion
- Protect the power supply to the fridge
- Seek advice if unsure
 - State and Territory Health Departments if NIP vaccines
 - Pharmaceutical manufacturer if non-NIP vaccines







Immediate actions to take if an error is made:

- Admit it!
- Inform the patient (or the parent) and apologize
- If the wrong vaccine was administered explain risks and benefits of the vaccine
- If an expired vaccine, or a vaccine stored outside recommended temperature range was administered –
 seek advice:
 - NIP vaccines State and Territory Health Department
 - non-NIP vaccines vaccine manufacturer
- Advise State and Territory Health Department
- DOCUMENT, DOCUMENT, DOCUMENT





Follow-up actions to take if an error is) made:

- Contact the patient (or the parent) in 24 hours and assess if an adverse event has occurred if it has,
 assist them with reporting the event
- Meet with your team discuss the event, examine why the error happened, develop and implement strategies to reduce the risk of recurrence
- Contact the patient (or the parent)
 - check they are OK
 - apologize again
 - inform them that strategies have been implemented to prevent reoccurrence



Take home messages:

- Vaccine administration has become more complicated due to new vaccines, combination vaccines, and the increasing complexity of the immunisation schedule.
- Know your limitations and never assume you know everything about vaccine administration. Know your own limitations and your profession's scope of practice.
- Don't feel obligated to perform functions (including administering different vaccines) if you are not skilled and knowledgeable enough to do so safely.
- If you do make a vaccine administration error, follow the immediate and follow-up process, inform your State or Territory Health Department, and learn from it so that you don't repeat it



