



'Good Catch' Initiative

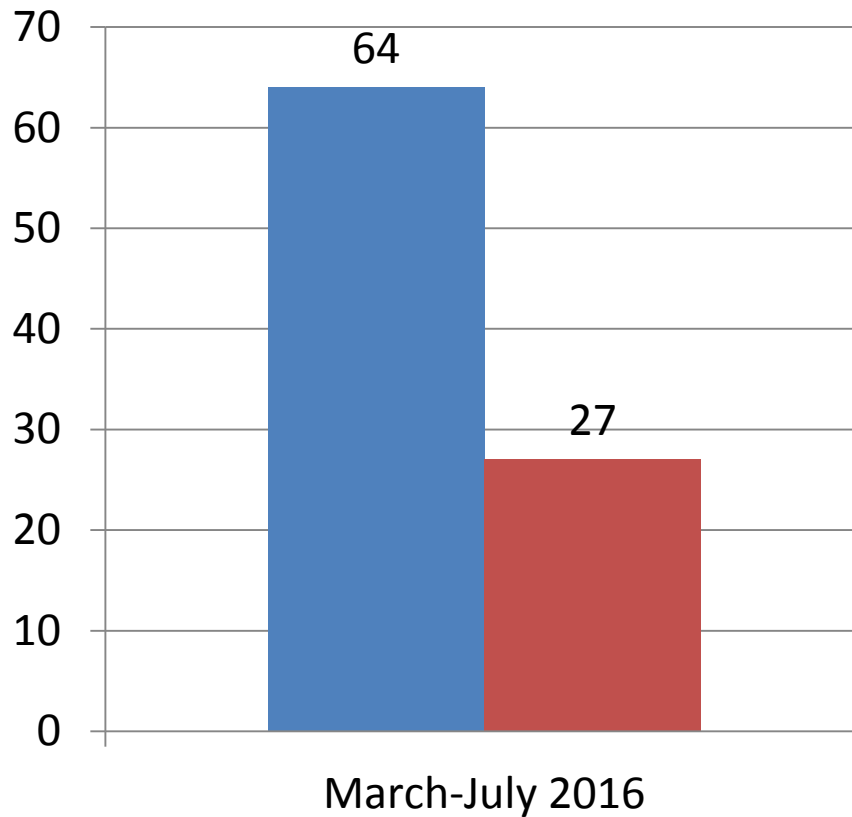
Promoting reporting of medication safety near misses

Louise Hendrick, Lead NCHD

Reena Patel, Chief Pharmacist

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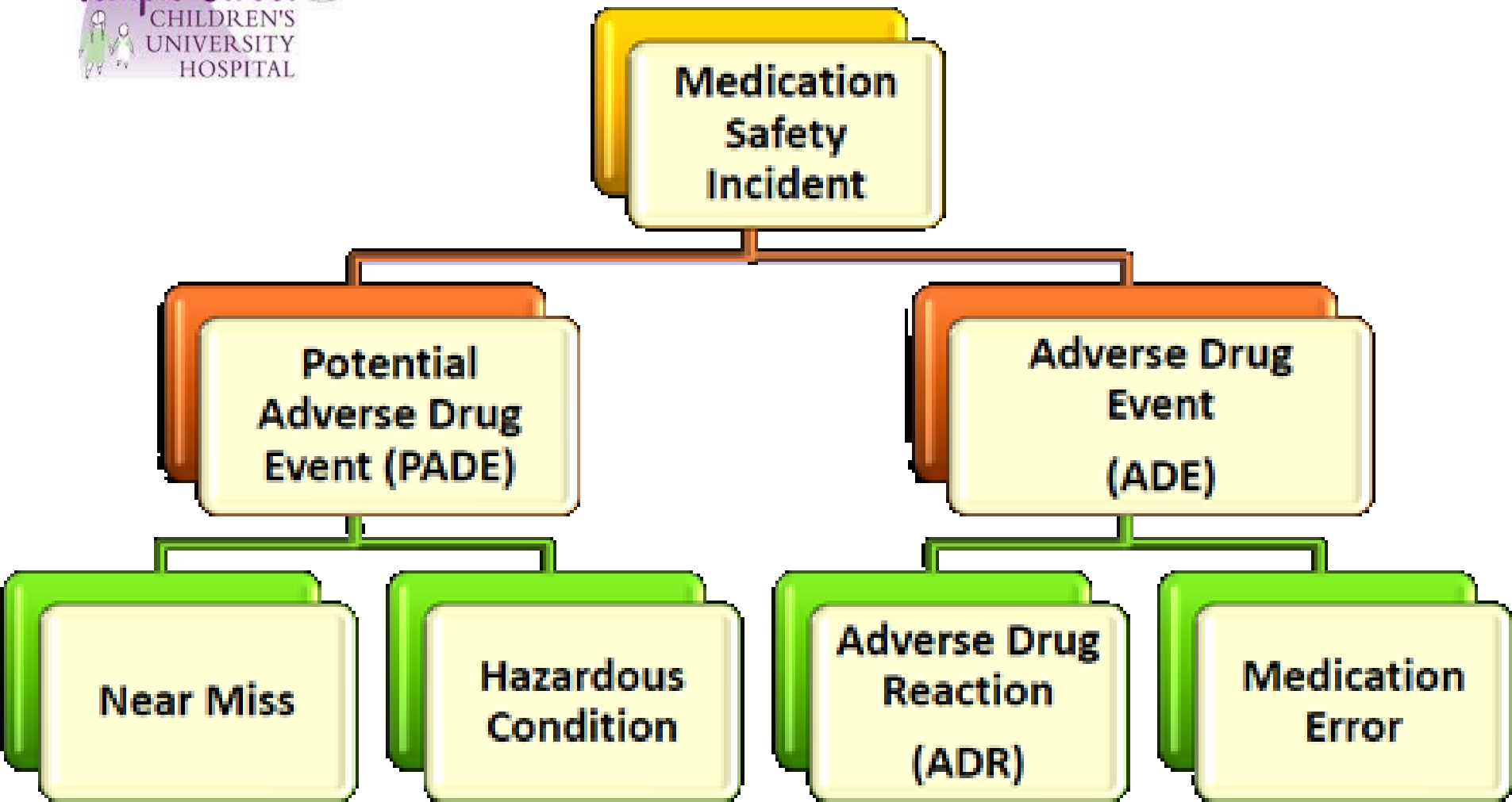
Paula Day, Risk & Legal Management

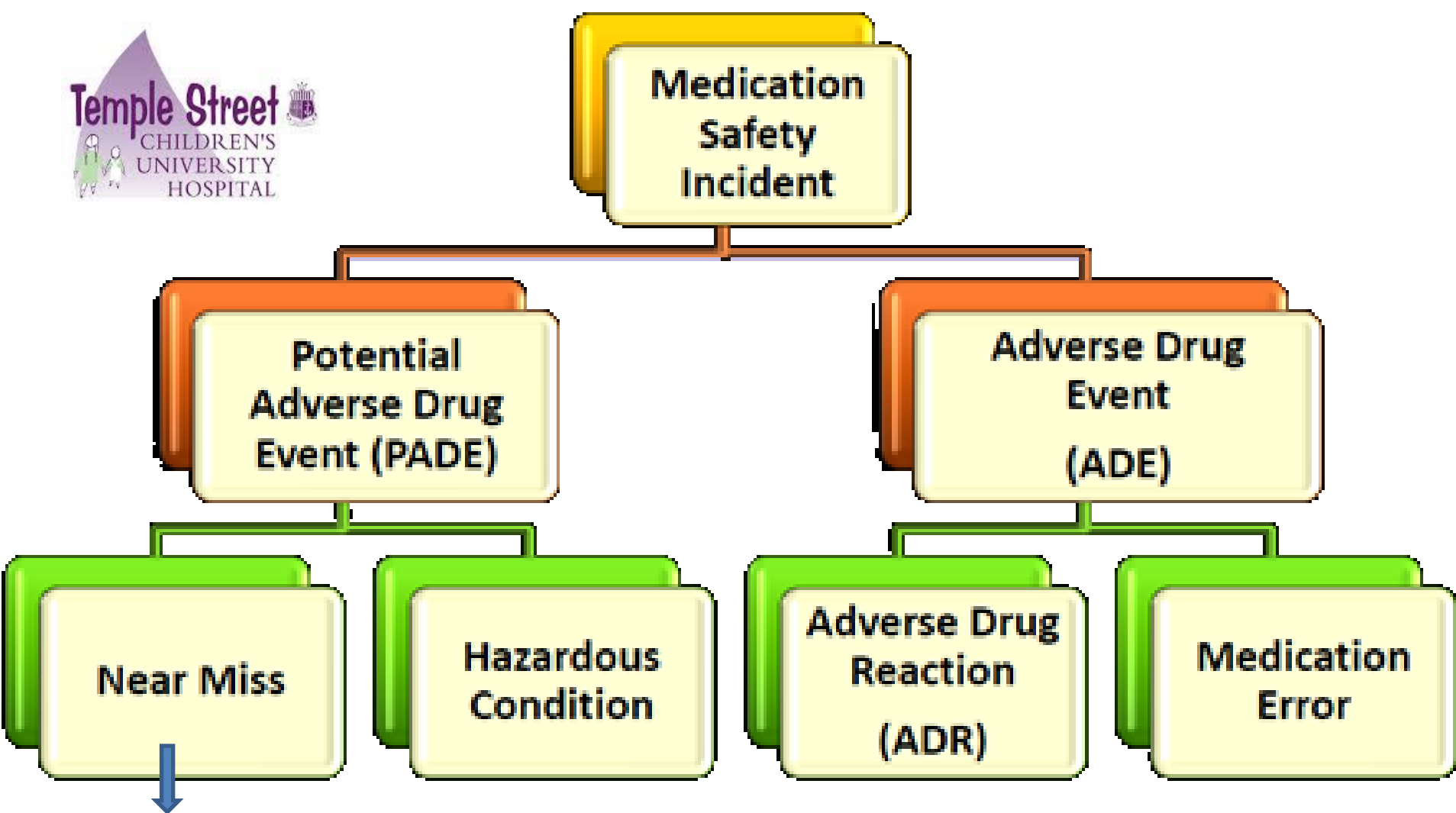


“We have very few near misses”

19,751 Admissions (day cases and inpatients)

Put Children First





never reached the patient

or

reached the patient but did not cause harm



Background

Problems

- Passive surveillance
- Under-reporting common

Aim

- Assess barriers to and promote increased reporting of near misses

Barriers

- Difficulty navigating through RESPOND reporting system, requiring information not readily to hand
- Length of time to complete report: minimum of 8 minutes per case
- Assumption that another staff member would complete the report
- Lack of feedback on submitted reports

Actions taken

1. Medication Safety Working Group
2. Paper-based alternative developed & launched
April 2017
3. Simple 'tick the box' design, 1-2 minutes to complete
4. Low cost with booklet placed on all wards & in the res
5. Near misses rebranded as good catches
6. Form anonymised
7. NCHD and Nurse champion



Form

Temple Street Children's University Hospital MEDICATION SAFETY REPORT FORM

Affix a Patient Identification Label

Patient Name: _____

Hospital Number: _____

Date of Birth: _____

Consultant: _____ Speciality: _____

Date of Report (DD/MM/YYYY): _____

Date of Incident (if different): _____

Time of Incident (24 hr clock): _____

Ward/Dept: _____

Grade of person reporting incident: _____

CATEGORY OF EVENT

Actual Good catch (Near miss)

STAGE OF PROCESS WHERE EVENT OCCURRED

Prescription Ordering/Supply Dispensing Administration Storage Monitoring

TYPE OF EVENT

Adverse Drug Reaction (ADR)	<input type="checkbox"/>	Incorrect storage	<input type="checkbox"/>
Allergic reaction to known allergen	<input type="checkbox"/>	Incorrect time	<input type="checkbox"/>
Contraindication to use of medication	<input type="checkbox"/>	Incorrect patient weight / BSA / BMI recorded	<input type="checkbox"/>
Drug omission (no. of episodes.....)	<input type="checkbox"/>	Infusion pump incident	<input type="checkbox"/>
Drug Security Issue	<input type="checkbox"/>	Interaction: (Drug – drug, drug-food)	<input type="checkbox"/>
Expired drug	<input type="checkbox"/>	IV infusion problem (diluent type / volume, rate of administration)	<input type="checkbox"/>
Incorrect anatomical site	<input type="checkbox"/>	Medication on admission / transfer / discharge Incorrect	<input type="checkbox"/>
Incorrect diluent / method of reconstitution	<input type="checkbox"/>	Monitoring issue	<input type="checkbox"/>
Incorrect dose (over/under/extra)	<input type="checkbox"/>	Non-compliance with hospital policy	<input type="checkbox"/>
Incorrect drug / drug formulation	<input type="checkbox"/>	Therapeutic duplication	<input type="checkbox"/>
Incorrect duration of treatment	<input type="checkbox"/>	Unauthorised self-administration / administration by parent	<input type="checkbox"/>
Incorrect frequency	<input type="checkbox"/>	Unclear / incomplete documentation	<input type="checkbox"/>
Incorrect patient	<input type="checkbox"/>	Unclear / incomplete / incorrect dispensary labelling	<input type="checkbox"/>
Incorrect route	<input type="checkbox"/>	Unclear / incomplete prescription	<input type="checkbox"/>

Other (please specify) _____

Description of Event / Good catch (near miss)

OUTCOME

Did event result in harm (e.g. pain, injury, development or worsening of symptoms)? Yes No Uncertain

Details:

ACTION

Was action required to treat the patient? Yes No

Antidote administered	<input type="checkbox"/>	Oxygen	<input type="checkbox"/>
Drug therapy	<input type="checkbox"/>	CPR administered	<input type="checkbox"/>
Laboratory tests performed	<input type="checkbox"/>	Intensive care	<input type="checkbox"/>
Observation of vital signs increased	<input type="checkbox"/>	Reported to HPRA	<input type="checkbox"/>
Other (please specify below)	<input type="checkbox"/>		<input type="checkbox"/>

PERSONS INFORMED

Consultant / Team Yes No Not Applicable Doctor notified _____ Bleep no. _____

Patient/Parent aware Yes No Not Applicable Informed by _____

Nursing admin Yes No Person Informed _____

Documented in the health care record Yes No

CONTRIBUTING FACTORS

Identify any apparent contributing factor(s) to event			
Calculation error	<input type="checkbox"/>	Medication not cancelled	<input type="checkbox"/>
Communication factors	<input type="checkbox"/>	Medication unavailable	<input type="checkbox"/>
Documentation error	<input type="checkbox"/>	Multiple kardexes for patient	<input type="checkbox"/>
Drug look-alike/sound-alike confusion	<input type="checkbox"/>	Order misinterpreted/misread	<input type="checkbox"/>
Equipment malfunction	<input type="checkbox"/>	Procedure/policy not available	<input type="checkbox"/>
Illegible/unclear prescription	<input type="checkbox"/>	Procedure/policy not followed	<input type="checkbox"/>
Information missing on prescription	<input type="checkbox"/>	Transcription error	<input type="checkbox"/>
Knowledge/skills deficit	<input type="checkbox"/>	Work environment factors	<input type="checkbox"/>
Other (please specify): _____			

MEDICATION(S) INVOLVED

Generic name(s): (e.g. prednisolone) _____

Brand name(s) if applicable: (e.g. Delfacortri) _____

Strength(s) /concentration (s); dosage form (s); route(s); (e.g. 5mg tablets po) _____

Batch Number: _____ Expiry Date: _____

(Batch no. and Expiry Date are required if reporting an adverse drug reaction or a flaw with a particular medication)

INFUSION PUMP DETAILS

Brand/Make of pump: _____ Other: _____

Pump Asset No: _____ Other: _____

Has Clinical Engineer been notified? Yes No

Name of Reporter _____ Contact phone number/bleep/e-mail _____

Note: Name of reporter and contact details are optional; however it can facilitate follow-up and feedback if they are provided.

ENTERED ON RESPOND Yes No RESPOND REF NUMBER _____

If applicable, please describe any follow-up or actions identified or taken to reduce the chance of this incident recurring. Do not delay submission of report to complete this section - action taken can be notified at a later date.

Please fill out the form as completely as possible, and send in confidence (by hand, internal mail) to: The Risk Management Department. The Risk Manager may be contacted on extension 1800 or at paula.day@cuh.ie

Official Use Only

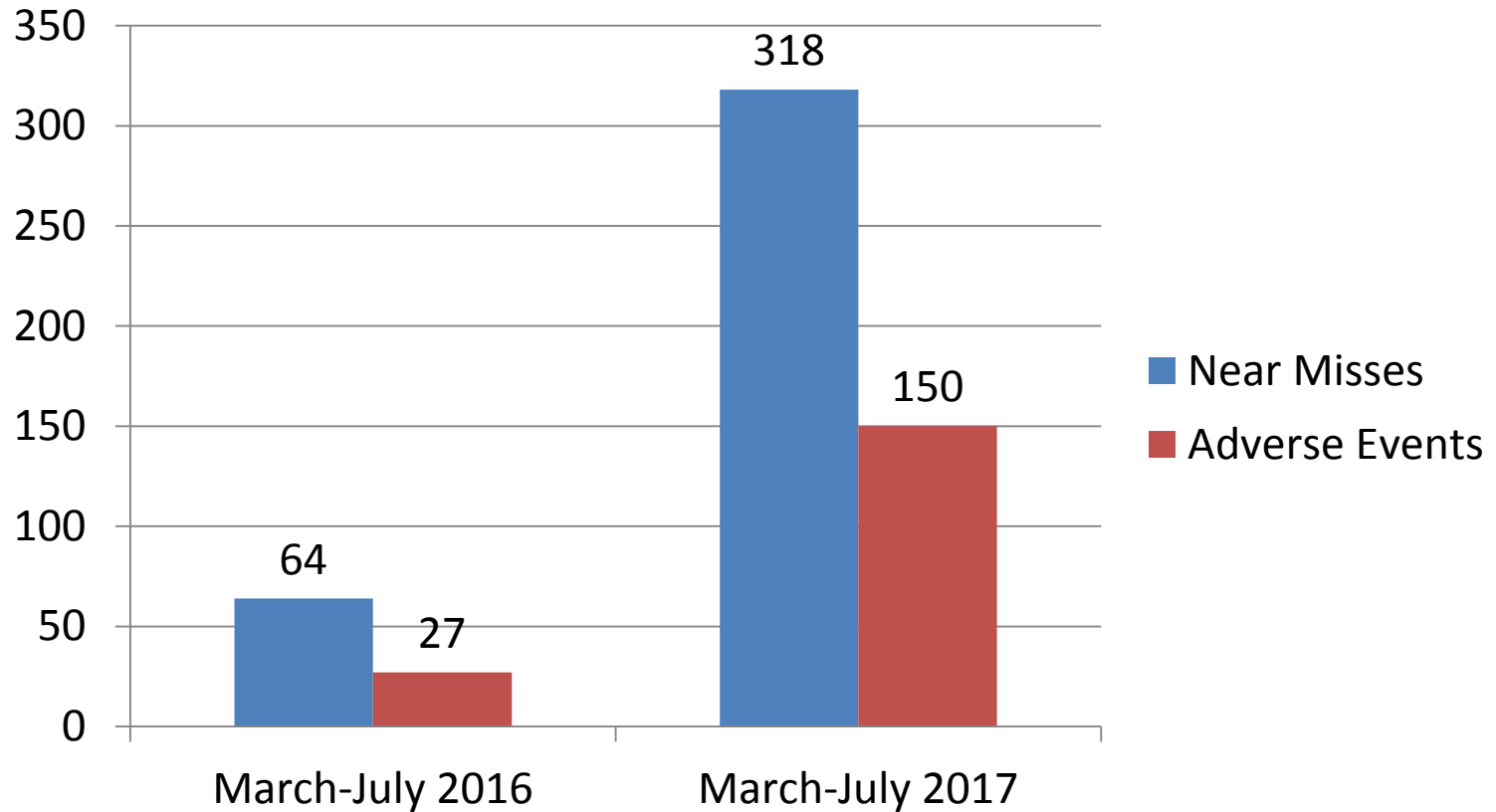
Date received: _____

NCCMERP Category: _____ Nursing Administration Notified: _____

Severity _____ CIS Notified _____

Likelihood of recurrence _____

Results



Put Children First

Sustainability

- Feedback
- NCHD champion
- Redesign of online reporting module
- Still under-reporting?
- Consider high risk medications card
- New Clinical transfer task module within Clinical portal linked to BNFC and Crumlin formulary
 - Warning box for high risk medications



Thank you...

To the incredibly amazing team at Temple Street

Caroline O'Connor

Reena Patel

Paula Day

Karen Cunningham

Tips

- You are not alone
- Think small to go big
- Front line ownership is key
 - Engage the engaged
 - Try and win over the middle group
 - Recognise some people will never come along....
- It's good to fail, but pick yourself up...
- Don't be hard on yourself
- Remember you are here to advocate for NCHDs
- Don't forget interns and non-scheme trainees