

Enhance Your Medication Error Reporting Program to Improve Global Medication Safety

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Objectives

- Identify factors that may cause healthcare practitioners to be reluctant to report an error.
- State how the underreporting of errors as well as the lack of a thorough investigation prevents effective system enhancement and improvement of patient outcomes.
- Identify useful methods of collecting, analyzing, and using data about medication safety.



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In order to convince staff to report, we need to know.....

Why are we reporting potential and actual medication errors?

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Why do we have a reporting program?

- —"Learn why errors are occurring"
- "Identify trends within our organization"
- "To determine how well we are doing"

Is this happening in your organization?



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Purpose of Reporting Systems

- Support a culture of open communication
- Promote the concept that each employee is an important contributor to improvements
- Identify latent and active failures
- Provide a record of the event
- Ineffective way to collect quantitative "data" or "trends"



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What can a reporting system do?

- Identify local system hazards;
 - Most valuable lessons can occur from a <u>single</u> report
 - Tools can provide a systematic analytic framework to learn from these events (e.g., ISMP Assess-Err)
- Aggregate experiences for uncommon conditions
 - Patient Safety Organizations (PSOs)



https://www.ismp.org/resources/assess-err-worksheet

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What can a reporting system do?

- Improve patient safety culture
 - How YOUR organization views patient safety
 - Communication with staff could change culture
- Share lessons within and across organizations
 - Lessons learned can be used to prevent the same types of adverse events
 - Use of external sources of information
 - Failure to examine potential for errors
 - "It's never happened here"



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Ok, that was nice. But...

Why are we not <u>getting</u> many medication error reports?

Why does that same event keep happening?

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Limitations to Reporting Systems

- Rarely generates in-depth analyses or result in strong interventions
 - Unrealistic expectations of what staff will know
 - Staff often have limited training in adverse event investigation
 - Lack of time to report
 - Filling out a report takes too long
 - Inappropriately designed forms
 - Too much emphasis on front line staff to fill out the entire form



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Limitations to Reporting Systems

- Rarely generates in-depth analyses or result in strong interventions to reduce risk con't
 - Error investigations and analysis are often superficial
 - Superficial analysis = no meaningful change
 - Majority of changes being informing staff and education/training (low level strategy)
 - Lack of meaningful change diminishes value of your reporting system



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Limitations to Reporting Systems

- May generate too many reports;
 - Reports usually include many that are incomplete and/or inaccurate "Nurse gave the wrong drug"

"Pharmacist dispensed wrong dose"

- No resources to read or analyze <u>all</u> of these reports.
- Provides minimal data about medication-use system
- Reporting used to complain
- Dissatisfaction from users when "nothing's changed"



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Limitations to Reporting Systems

- Can't be used to measure safety (error rates);
 - Events are under-reported
 - Why are they under-reported?
 - What is reportable? Definition?
 - Rely on the vigilance, time, honesty, and whim of healthcare providers to detect and report adverse events
 - Some provider types report adverse events with regularity (nurses), some don't
 - Lack of feedback to staff
 - Fear of punishment or ridicule



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Voluntary Reporting

- "We found that less than 4% of all adverse drug events involving use of rescue drugs were reported."
- "Studies of medical services suggest that only 1.5% of all adverse events result in an incident report and only 6% of adverse drug events are identified through traditional incident reporting or a telephone hotline."



Schade, Am J Med Qual. 2006 Sep-Oct;21(5):335-41 O'Neil A et al. Physician reporting compared with medical-record review to identify adverse medical events. Ann Intern Med. 1993;119:370-376 www.ismp.org | 13

Voluntary Reporting Systems

- "Routine reporting system implemented in a large hospital missed incidents identified by case note review and detected only 5% of incidents that resulted in patient harm."
- "Of the events experienced by Medicare beneficiaries discharged in October 2008, hospital incident reporting systems captured only an estimated 14 percent."



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Sari et al. Sensitivity of routine system for reporting patient safety incidents in an NHS hospital: retrospective patient case note review. BMJ 2007;334:79

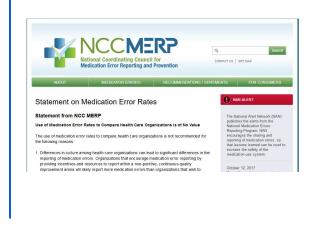
Levinson DR. Washington, DC: US Department of Health and Human Services, Office of the Inspector General;

January 2012. Hospital Incident Reporting Systems Do Not Capture Most Patient Harm. Report No. OEI-06-09-00091. @2020 ISMP | www.ismp.org | 14

Available from Internet: https://oig.hhs.gov/oei/reports/oei-06-09-00091.pdf

Limitations to Reporting Systems

- Can't be used to measure change
 - Increase in reports may be due to increased reporting (reporting bias) from increased awareness
 - Decrease may be typical challenges of reporting
- Can't be used to compare organizations





http://www.nccmerp.org/statement-medication-error-rates https://www.ismp.org/faq.asp#Question_1

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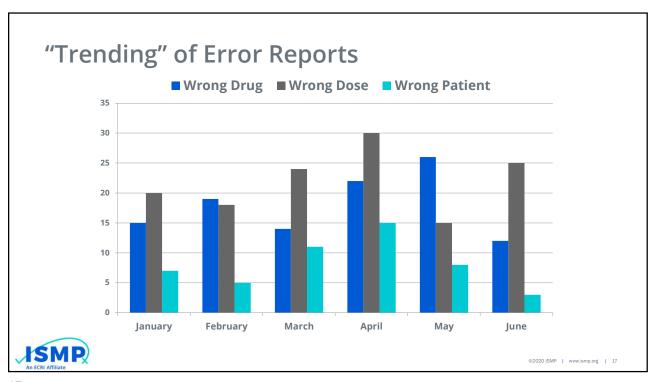
What is the national medication error rate?

- Numerator = Number of error reports received, but.....you just told me you don't get all the reports?!?!
- Therefore...cant "benchmark" a reliable "error-rate" to compare against anyone else
- Actually calculating an error reporting rate



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Which drug classes cause the most amount of harm in your organization?

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What does aggregated data really mean?

- Shows interesting "trends"
 - Reporting trend or event trend?
- No detail to work with
- Cause unclear at the macro level
 - What really happened?
 - Why did it happen?
- Potential false conclusions
- Event descriptions will tell you what happened



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Why do practitioners report to ISMP?

(and how does this apply to me?)

- Motivation is altruism
 - To promote change, prevent recurrences
- Evidence your information will be put to use
- Feedback on product and practice changes
- Trust that identity won't be revealed
- Confidential, non-punitive, can be anonymous at reporter's option



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Strategies to Maximize Error Reporting

- Make reporting easier
 - Make reporting easy and less burdensome
 - Quick and readily accessible reporting mechanism
 - Requires minimal training
 - Limit number of questions



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Strategies to Maximize Error Reporting

- Make reporting meaningful to the reporter
 - Won't report when identified problems are not remediated
- Supports a culture of open communication
- Promote the concept that each employee is an important contributor to improvements
- More reports = better culture



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Make Reporting Meaningful to the Reporter

- Communication
 - Provide feedback
 - Share reports with staff
 - Leaders should devote resources to collect AND analyze events AND mediate risk.
 - If staff observe change based upon their feedback, real changes in safety culture start to occur.



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Make Reporting Meaningful to the Reporter

- Communication
 - Measure the number of successful system changes, not events reported;
 - Ultimate measure of success is the amount of harm prevented, not the number of reports received



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Strategies to Maximize Error Reporting

- Prioritize which events to investigate
- Focused reporting of a finite set of high-yield events
 - Which drug is YOUR greatest concern?



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Error reporting is only a starting point...

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Risk Identification

- Many sources of information to detect potential for errors and actual errors
- Many methods to detect potential for errors and actual errors
- Using multiple methods of detecting risk will identify different types of risk that are not commonly reported



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Methods of Data Collection

- Proactive Risk Assessment
 - Self Assessments
 - Failure Mode and Effects Analysis (FMEA)
 - External Sources of Data
 - Walkrounds™
 - Staff Meetings, Safety Briefs
- Concurrent Risk Assessment
 - Pharmacy Interventions (clinical or dispensing staff)
 - Nursing Interventions
 - Triggers and Markers



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Methods of Data Collection

- Retrospective Risk Assessment
 - Observational methodology
 - · Data from technology
 - Chart reviews
 - · Internal, voluntary reporting
 - Medication errors
 - Adverse drug reactions (ADR)



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Conclusion

- Change your culture so that your staff WANTS to report to you
- Avoid collecting reports for the sake of counting reports
- One error report could be an indication that you have a bigger problem.
- Be realistic about what you are getting and what it means
- Consider a report to be the starting point, not the end point.



https://ismp.org/report-medication-error

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Questions?

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