

Quality Improvement and Today's Technician

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Objectives:

- Why do we have Quality Improvement
- Steps of a Quality Improvement Program
- How does the role of the technician fit into this

I have no conflicts of interest with this presentation.

By show of hands:

1. How many of you know if where you work has a quality improvement program?
2. How many of you are involved with your peer review?
3. How many of you are actively involved with Quality Improvement?

Why do we have Quality Improvement?

Why do we have Quality Improvement?

1. Patient Safety
No news is **not** good news.
We can not manage what we cannot measure.
"I am really careful and don't make mistakes.."
If a pharmacy fills 99.9% accurate
-If they fill 1,000 prescriptions each week
-How many errors each week?
-52,000 rx's in a year - what is the error rate of a very careful pharmacy?

Why do we have Quality Improvement?

1. Patient Safety
2. State Board Regulations
3. Medicare Part D – FWA and Insurance Contracts
4. Freedom from Discovery
 1. Information must be uploaded to an approved Patient Safety Organization Website.
 2. All paperwork must be labeled as **Patient Safety Work Product**. Mark your QI data as confidential and treat it as such.

5 Steps to a Quality Improvement Program



Step 1 – Establish Pharmacy Workflow

1. Divides the dispensing process into manageable parts, allowing us to concentrate on 1 process at a time.
2. Individuals within the workflow need to understand his/her role within the process.
3. Each process becomes a check on the other.
4. Increases efficiency.

Step 2 – Collect Quality Related Event Data

1. Responsibility of EVERY employee
2. Report QRE's to online website
3. Report dispensing errors using Process Improvement Report
 - Meets ND Board of Pharmacy requirements

Step 3 – Analyze QRE's

1. FHCP Peer Review Committee
 1. Meets monthly
 2. Consists of Pharmacy Director, Quality Supervisor, RPh, and Technician
 3. Review Process Improvement Forms

Step 3 – Analyze QRE's

2. Discussing Errors
 - Negative:
 - Dwells on the past
 - Threatening, insulting, punitive
 - Focuses on WHO made an error
 - Concludes erring individual has character flaws they need to correct
 - Individuals won't willingly participate

Step 3 – Analyze QRE's

2. Discussing Errors

Positive:

- Looks to the future
- Open, Blame-free, non-punitive
- Focuses on How or What in the system allowed the error to occur
- Realizes error is a reality and recognizes the opportunity to improve for the future
- Creates blame-free, shame-free environment

Step 4 – Formulate Improvement Plan

1. Propose workflow changes
2. Identify the need for training
3. Recommit to using existing workflow processes

Step 5 – Implementation

1. Communicate changes to pharmacy processes during staff meetings
2. Provide training when indicated
3. Allow staff to provide feedback, both positive and negative
4. Reassess changes and need for additional intervention

Annual Staff Training

1. Staff complete annual training on QI process
2. Provide monthly/quarterly newsletter from ISMP (Institute of Safe Medication Practices) as well as PQC (Pharmacy Quality Commitment)
3. PQC provides free QI related CE

Reporting of Errors

Two different ways to report

1. Errors that do not get past the pharmacist (near miss)
2. Errors that do get past the pharmacist (incident)

Errors that do not get past the pharmacist (near miss):

1. Errors still need to get entered into your PQC program
2. Allows us to generate reports to see where/when different types of errors occur
3. Allows us to proactively look at our system to find flaws before errors occur

Who Discussed the QDE? **Type required** **How/Why required**

When was the QDE discussed? **required** **When did the QDE start? required**

What Type of QDE was used? **required** **What time was the QDE used? required** **Received by patient? required**

Drug Prescribed **required** **Drug Dispensed? If different from prescribed**

Routing Date **required**

What happened and why? Why did you think this step had to be taken to avoid future incidents?

Why? Why not? What was your confidence level? Example: Did the sign-in the wrong patient?

Remember: unless stated otherwise, all reports are for the PICU. The QDE data entered for a patient should not be reported for patients that the breast feed is not. If you select your membership in the above chart, it will include the patient's health information only and not the patient.

- Contributing factors and systems
- Check all that apply:
- 1. Allergist/OTC use rules
 - 2. Incomplete medication reconciliation
 - 3. Incomplete reconciliation during medication reconciliation
 - 4. Human error (misreading, calculation, transcription)
 - 5. Inappropriate prescription
 - 6. No or incomplete patient verification or check against orders
 - 7. Patient education/consultation, education errors, changes, requests to add
 - 8. Pharmacy inventory & Culture/medication stock/shortage
 - 9. Patient Characteristics (allergies, look-alike/sound-alike and high-risk medications)
 - 10. Technology/Equipment or software defect
 - 11. Training or supervision
 - 12. Incomplete/Inconsistent medication reconciliation
 - 13. Medication
 - 14. Workflow or process issues
- Corrective Actions/Recommendations
- Check all that apply:
- 1. Revised policy or packaging
 - 2. Dispensing procedure or process change
 - 3. Medication/Supply (e.g. medication signing, physical count, check-in/hold, etc.) other
 - 4. Medication product change (order for other or inactive ingredients)
 - 5. No action planned
 - 6. Other
 - 7. Patient profile updating (Allergies, history, other)
 - 8. Pharmacy management factors (allergies, culture, policies & procedures)
 - 9. Policy Update to medical Linc.commission
 - 10. Staff Education/Training - equipment independent basic check back/backpack training staff
 - 11. Staff product knowledge update
 - 12. Software change - add workflow alert or other equipment requirement
 - 13. Staffing - change check-in management/monitor
 - 14. Incomplete medication reconciliation/medication change or adjustment
- Save QDE

Reporting Home Reports

Available Reports: Select A Report, Date Range: Today, Graph Type: Pie 3D, Location: All Locations

QDE REPORTS

- All Reports
- QDE Type
- QDE By Day of Week
- Where Was QDE Discussed
- Where Was QDE Used
- How It is Used
- Routing Date
- Contributing Factors
- Corrective Actions/Recommendations
- Possible in Error
- QDE Success Rate
- Error Percentage
- Which Report
- Received the Report
- DATA IN CASE REPORTS
- All Reports

Reporting Home Reports

Available Reports: Select A Report, Date Range: Today, Graph Type: Pie 3D, Location: All Locations

QDE REPORTS

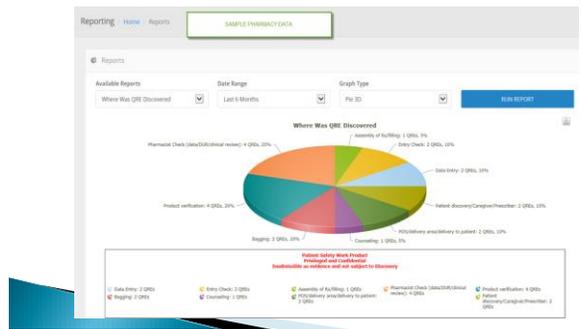
- Today
- Last 2 Days
- Last 7 Days
- Last 15 Days
- Last 30 Days
- Last Month
- Last 3 Months
- Custom Range

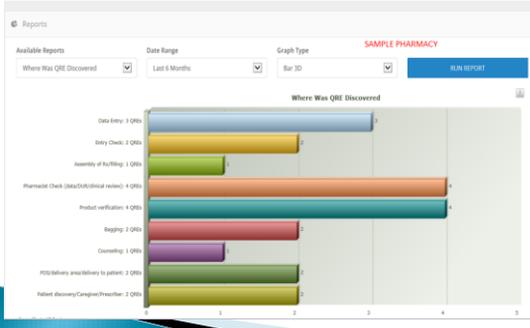
Reporting Home Reports

Available Reports: Select A Report, Date Range: Today, Graph Type: Pie 3D, Location: All Locations

QDE REPORTS

- QDE 3D
- Category 3D
- Category 2D
- Line 3D
- Area 3D
- Bar 3D
- Donut 3D
- Donut 2D
- Area 2D
- Bar 2D





Errors that get past the pharmacist (incident):

1. Need to be entered into your PQC website
2. Need additional report filled out (incident report)
3. All parties involved need to do both of the above steps
4. These reports are to be analyzed at the Peer Review Meeting

PQC Quality Management Tools Process Improvement Form for Peer Review Meetings
STEP ONE – IDENTIFY and DISCUSS

IDENTIFY ISSUES: Below record findings and trends from the data that you have reported investigated. Date: _____

Identified Incidents/Patterns (Date of incident must be one reported by Board of Pharmacy or other regulatory authority. *Do not report from PDCs for another incident.)	Notes/Findings	Process Issues Identified	Root Cause Identified
Incidents or patterns of errors that reached the patient			
Incidents or patterns of Adverse Events			

*Reportable Adverse Events (ADEs) include but are not limited to: "Incidents or patterns of errors that reached the patient" and "Incidents or patterns of Adverse Events".

PQC Quality Management Tools Process Improvement Form for Peer Review Meetings
STEP TWO – CORRECTIVE ACTION/ PREVENTATIVE ACTION AND ASSESSMENT

Use chart below to track quality improvement changes made as a result of Quality Improvement Peer Review meetings.

Implementation Date	Corrective Action/ Preventative Action/ Process Improvement plan	Essential elements of quality improvement effort	How successful results/ Assessment of patient safety	Follow up and Assessment of Action Plan - Follow up needed

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Incidents or patterns of errors that reached the patient	The pharmacist has incorrectly filled. Checked during control of Subsequent filling	Errors in stock took not have to be PDC reviewed on their list and then you have to make sure the PDC has the NDC number in the system	NDC was not verified during control and there was an incomplete label
Incidents or patterns of Adverse Events			

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8/17/2018	A. Not have another looking B. Make sure of the drugs to the stock side C. Ensure that NDCs are being looked on errors to stock side at the time they are being entered	To make sure all medications are following all work flow process		As follow up date, use of corrective actions made are being followed, as if any of the corrective actions need adjustments

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REVIEW:

1. Is this a positive or negative entry into PQC for an incident?

I went to fill the posted prescription and the posting tech had entered the day supply in wrong.

Review:

2.What should happen in this situation?

- a. Posting tech should enter into PQC what lead them to post the rx with the wrong day supply
- b. Filling tech should enter into PQC why they did not catch the incorrect day supply
- c. Person counting the rx should enter into PQC why they did not catch the incorrect day supply
- d. If it got past the pharmacist, the pharmacist should enter into PQC why they did not catch the incorrect day supply and an incident report should be filled out.

REVIEW:

3.Why do we have Quality Improvement?

- A.Patient Safety
- B.State Regulations
- C.Medicare Part D – FWA and Insurance Contracts
- D.Freedom from Discovery
- E.All of the above

REVIEW:

3.Why do we have Quality Improvement?

- A.Patient Safety
- B.State Regulations
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- D.Freedom from Discovery
- E.All of the above**

Review:

4.What are the steps of a Quality Improvement Program?

- A.Establish Pharmacy Workflow
- B.Collect QRE's
- C.Analyze QRE's
- D.Formulate plan for improvement
- E.Implement New Process or Training
- F.All of the Above

Review:

4.What are the steps of a Quality Improvement Program?

- A.Establish Pharmacy Workflow
- B.Collect QRE's
- C.Analyze QRE's
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- E.Implement New Process or Training
- F.All of the Above**

Review:

5. How does the Technician fit into Quality Improvement?
- A. Always keep in mind patient safety
 - B. It is the responsibility of every employee, including the technician
 - C. Technician input, both prospectively and retrospectively allows us to analyze our system for flaws
 - D. Employee feedback, including technicians, helps us to determine if our trainings or implementations were successful or need to be re-evaluated.
 - E. All of the above
- 

Review:

5. How does the Technician fit into Quality Improvement?
- A. Always keep in mind patient safety
 - B. It is the responsibility of every employee, including the technician
 - C. Technician input, both prospectively and retrospectively allows us to analyze our system for flaws
 - D. Employee feedback, including technicians, helps us to determine if our trainings or implementations were successful or need to be re-evaluated.

E. All of the above



References

ND Board of Pharmacy Administrative Code 61-02-09
Alliance for Patient Medication Safety (APMS) Pharmacy
Quality Commitment (PQC+) Program (powerpoint)

