Concise Incident Analysis Tool

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Co-Investigators

Julius Cuong Pham, MD, PhD Armstrong Institute for Patient Safety and Quality, Johns Hopkins University School of Medicine Carolyn Hoffman, RN, MN Quality & Healthcare Improvement, Alberta Health Services

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CHAPTER 1: BACKGROUND

Root Cause Analysis (RCA) investigations of patient safety incidents¹ have played an important role in

improving care. This rigorous methodology is designed to ensure that all relevant aspects of an incident are understood and that effective actions are taken to reduce the risk of recurrence.

Given the complexity of the healthcare environment and the significant resource requirements of an RCA (a form of comprehensive incident analysis), health care leaders and patient safety experts have begun to look for a more "concise" method of incident analysis to identify a more timely yet accurate approach. Examples of abbreviated incident analysis methodologies exist²³⁴⁵⁶ but evaluation of their effectiveness has been limited.²

Why would you want to use the concise method?

Concise Analysis is a less resource intensive approach to incident analysis that may contribute important knowledge regarding a larger number of incidents. A conscious and deliberate decision has been made to focus primarily on four aspects: the agreed-upon facts, key contributing factors and findings, actions for improvement (if any), and evaluation. The local learning can then flow into the higher organizational level for prioritization of risks and integration into a systematic quality improvement approach for improving patient safety. A concise incident analysis uses a systems approach and consideration of human factors.

Note: incident analysis should comply with all local policies and legislation. A Concise approach is not suitable for all types of reviews. It may be useful to transition from a Concise approach to some other type of analysis as new information is available.

Comprehensive incident analysis is defined as: analysis by an interdisciplinary group of staff and physicians that is facilitated by a person(s) with knowledge of the process, human factors and effective solutions development in healthcare. The process may take up to 90 days due to the depth and breadth of the Incidents resulting in analysis. none, mild, moderate, severe patient harm and/or death may receive **Comprehensive** Event Analysis.

as: analysis that is usually conducted by a person with knowledge of the incident analysis process, human factors and effective solutions development in healthcare with input gathered from staff and physicians local to the event. The process is often completed within hours or days due to the less intensive approach. Incidents with none, mild, and moderate patient harm may receive Concise Incident Analysis.

Concise incident analysis is defined

¹ A patient safety incident is an event or circumstance that could have resulted, or did result, in unnecessary harm to a patient (World Health Organization. Conceptual Framework for the International Classification for Patient Safety. 2009. Available from: http://www.who.int/patientsafety/implementation/taxonomy/icps_technical_report_en.pdf)

Ruddick P. et al. (2008) Using root cause analysis to reduce falls in rural health care facilities. NCBI Bookshelf. Accessed on July 14, 2011 at: http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=aps2v1&part=advances-ruddick_61

³ National Patient Safety Agency. (2008) Root Cause Analysis Investigation Tools: Three levels of RCA investigation – guidance. Retrieved on July 14, 2011 at: http://www.nrls.npsa.nhs.uk/resources/?EntryId45=75355

⁴ VA National Center for Patient Safety (2004). Retrieved on July 14, 2011 at:

http://www.patientsafety.gov/curriculum/TeachingMethods/PtSafety Case Conference Format/index.html

⁵ Johns Hopkins Quality and Safety Research Group (2007). On the cusp: Stop BSI. Retrieved on July 14, 2011 at: http://www.safercare.net/OTCSBSI/Resources.html

⁶ http://www.patientsafetyinstitute.ca/English/toolsResources/IncidentAnalysis/Pages/default.aspx

CHAPTER 2: OVERVIEW OF CONCISE INCIDENT ANALYSIS METHODOLOGY

A. Case selection

- Determine if incident analysis is appropriate.
- Determine if concise incident analysis is appropriate.

B. Understand What Happened

Obtain sufficient information to understand the incident.

- Identify a facilitator to conduct analysis.
- Gather facts from records and other applicable documents.
- If applicable, examine the equipment, product, or environment.
- Have informal discussions with patient/family, provider(s), manager(s), attending physician and/or expert(s) in the specific circumstance, equipment, and/or product.
- Develop a high-level timeline or narrative description.

C. Determine How and Why It Happened

Analyze information to identify key contributing factors and the relationships among them. Use systems approach and human factors.

- Use the guiding questions to BRIEFLY explore all the domains of contributing factors.
- Select some specific guiding questions.
- Identify and map the contributing factors as well as the relationship between them.
- Summarize findings as summary statements.

D. Develop and Manage Actions for Improvement

If there is sufficient evidence to formulate actions for improvement to reduce the risk of recurrence and make care safer:

- List actions for improvement (evidence-based where possible and always striving to select the most rigorous action possible on the Hierarchy of Effectiveness);
- Include proposed persons accountable for implementation, timeline and an evaluation strategy for each action;
- Discuss the list with applicable decision maker for decision and action.

E. Evaluate the effect of the actions for improvement

The implementation status of each action is followed through and its effect evaluated based on the strategy identified in the previous phase.

The general lessons and findings should be disseminated within and, where applicable, outside the organization to prevent a recurrence of the incident.

A. Case Selection

Determine if incident analysis is appropriate

Because the concise analysis method is not suitable for all types of analyses, the first step in the process is to determine if systems-based incident analysis is appropriate. The following types of incidents are not recommended for a systems-based incident analysis:

- 1. Events thought to be the result of a criminal act;
- 2. Purposefully unsafe acts (an act where care providers intend to cause harm by their actions);
- 3. Acts related to substance abuse by provider/staff; and
- 4. Events involving suspected patient abuse of any kind.

These situations should be referred to suitable administrative, professional, or regulatory bodies for resolution.

Determine if concise incident analysis is appropriate.

If a systems-based incident analysis is suitable, the following attributes may be used to determine if concise incident analysis is appropriate for that case.

- Incidents that resulted in no or low harm to the patient.
- Incidents primarily limited to one work area, division, or department.
- New incidents for which a comprehensive analysis was recently completed.
- Initial review to determine whether or not a comprehensive incident analysis is warranted.

Note: not all information regarding the incident may be available during the case selection process, therefore the facilitator selects the optimal method and anticipates the potential for changing the method as new information emerges.

Table 2.1 describes the characteristics of concise and comprehensive methods to assist with case selection.

Table 2.1: Characteristics of Concise and Comprehensive Incident Analysis⁷

Characteristic	Concise	Compre- hensive?
Should include person(s) with knowledge of incident analysis, human factors, systems approach and effective solutions development	Yes	Yes
Often facilitated by an individual with input gathered from the patient, family, staff, and physicians local to the incident as well as organizational or external experts	Yes	No
Conducted by an inter-disciplinary medium to large ad hoc group (may include patients, family members, staff, and physicians local to the incident as well as recognized independent internal or external experts/consultants not involved in the incident)	No	Yes
Time taken for analysis	Short timeframe (hours to days)	Long timeframe (up to 90 days)
Identifies contributing factors as well as remedial actions(s) taken (if any)	Focus on key factors	Yes
Recommendations for improvement	Yes (if applicable)	Yes
Principles of incident analysis (begins as soon as possible, includes all involved in the incident [including patient/family] and leadership of the organization, is objective and impartial, is thorough, considers relevant literature and evidence)	Reflects the intent, but may not address all	Incorporates all principles
Evaluation strategy	Yes (if applicable)	Yes

B. Understand what happened

Identify a facilitator

A facilitator (analyst/reviewer) with knowledge and skills in incident analysis, human factors, systems approach and effective solution development performs the concise analysis. The facilitator usually gains this expertise through a formal education program and/or mentored experience. The individual may be a healthcare provider or other professional such as a process improvement expert; this individual does not necessarily have to be a risk manager or quality improvement consultant.

Gather facts

The facilitator should gather facts from different sources to understand what happened and to develop a high-level timeline or narrative of the incident from:

⁷ Canadian Patient Safety Institute. Canadian Incident Analysis Framework. Edmonton, AB: Canadian Patient Safety Institute; 2012. Available

http://www.patientsafetyinstitute.ca/English/toolsResources/IncidentAnalysis/Documents/Canadian%20Incident%20Analysis%20Framework.P_DF

- Records (health record, incident report) and other documents
- Discussions (interviews) with the healthcare providers, managers, experts, patients, and/or family members directly involved in the incident.⁸
- Equipment/ products examination (if applicable)

Interview principles

- Interviews should be conducted as soon as reasonably possible after the incident for two reasons. First, memories fade quickly and important details may be lost over time. Second, as individuals involved in the incident discuss their recollections with one another, versions may blur together and the opportunity to obtain unique perspectives and details may be missed.
- Informal interviews should be conducted one person at a time so that individual perspectives about the incident are well understood.
- A cooperative approach is encouraged, using open-ended questions. Individuals should be asked to "tell their story" and possibly re-enact the incident or portions of the incident. Ask individuals if there are any factors that contributed to the incident as well as factors that mitigated the outcome of the incident (e.g. "what went well").
- Sincerely thank people for helping and ensure that their questions about the process are answered.

Gathering equipment/products/items

Gather materials such as the equipment and any product/care items used during or close to the time of the incident that may have directly or indirectly contributed to the circumstances. They can be secured for testing and review. They include, but are not limited to, biomedical equipment, IV solutions, medications, packaging, garments, etc. Photographs of the items and workspace are often helpful.

Develop a timeline or narrative description

Document key factual information in the form of a high-level timeline or narrative description. It is common to provide this information in the form of a narrative chronological description. This understanding will collate information from various sources, including the health record and informal interviews with key individuals. It is important that the timeline include only the actual facts or processes as they occurred, and not what was supposed to happen.

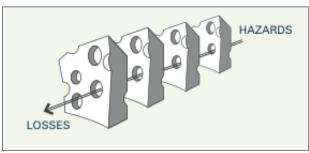
⁸ Investigative Interview Guidance. NRLS Root Cause Analysis (RCA) Investigation Guidance. http://www.nrls.npsa.nhs.uk/resources/?entryid45=75355

C. Determine how and why it happened

Concepts

There are two key concepts to consider when ensuring that the analysis reflects the complexities of the current healthcare system while remaining practical: the systems approach (as illustrated by the Swiss Cheese Model in Figure 2.1) and the domain of human factors. These concepts support a deeper understanding of how and why incidents occur in healthcare, including the identification of specific contributing factors.

Figure 2.1: The Swiss Cheese Model⁷



James Reason's Swiss Cheese Model⁷ provides a framework for understanding and analyzing the complex and dynamic nature of patient care within a systems perspective. The model explains how the defenses, barriers and safeguards that exist in a system are not impermeable and can be penetrated. This occurs when active failures (unsafe acts) and latent conditions (dormant system conditions) combine to create the opportunity for an incident. Latent conditions can be identified and corrected. Targeted strategies can also mitigate the frequency and severity of unsafe acts. It also points to the fact that humans are fallible and errors are to be expected even in the best organizations because people are incapable of perfect performance every time.

The questions to ask when an incident happens are how and why the defenses in the system failed and in the case of a near miss, how did they succeed; in other words, look at the system as a whole, rather than just at the actions of individuals.

At its core, the science of human factors examines how humans interact with the world around them. This specialized knowledge is used to help determine how and why incidents occur as well as help design efficient, human-centered processes to improve reliability and safety.

Historically, when an incident occurred, the tendency was to look for the most obvious explanation of what and why it happened. In most cases, individual human error was identified as the cause, primarily because it was easy to identify (frequently referred to as the "sharp end" of the system) and appeared to be easy to fix. Patient safety experts advocate a way of thinking that views human error as a symptom of broader issues within a poorly designed system (often referred to as the "blunt end" of the system), such as an adverse physical or organizational environment. A deeper inquiry into the circumstances will yield

system-based contributing factors. Recommended actions for improvement vary significantly and may range from physically changing the design of a software interface, sign, form, or medical device to changing the entire design of a room in a facility to optimize safety and efficiency.

Identify key contributing factors

Use the information gathered to identify key factors that contributed to the incident occurring. Two key questions that assist in this process are: "how did this happen?" and "what else influenced the circumstances?" The facilitator continues to ask "how" and "what influenced it" questions until no further information can be generated for the key contributing factors.

Use the guiding questions (workbook) to explore different domains (task, equipment, work environment, patient characteristics, care team, organization, other) of factors that may have contributed to this incident. Briefly explore each domain. For domains that are relevant, further explore each specific question.

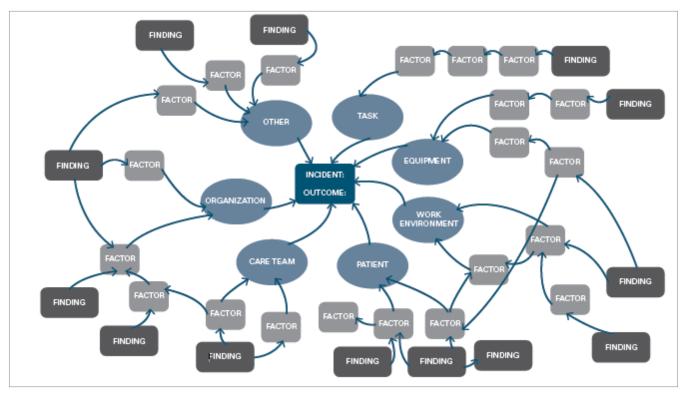
Identify the relationship between contributing factors using a diagram

Diagramming is a helpful exercise in understanding the relationship between contributing factors. The Tree (Figure 2.2) and Constellation Diagram (Figure 2.3) are two potential tools to accomplish this. The benefits of the constellation diagram include a visual description of the cascading aspects of each contributing factor. This allows for a better understanding of the relationship between contributing factors and identification of clusters of factors where they directly impact one another. These clusters are most often the basis for the development of recommended actions. The overall goal is to determine if an action

Caused by Caused by Caused by ROOT ACTION OF CONDITION ACTION OR CONDITION CAUSE Caused by Caused by Caused by HARMFUL INCIDENT ACTION OR CONDITION ACTION OR CONDITION ROOT CAUSE Caused by Caused by Caused by ACTION OR CONDITION ACTION OR CONDITION ACTION OF CONDITION ROOT Caused by CAUSE ACTION OR CONDITION Caused by

Figure 2.2 Tree diagram⁷

Figure 2.3 Constellation Diagram⁷



or a small number of actions can be taken to address all key contributing factors identified.

Prioritize and Summarize Findings

Once the team has completed the analysis, contributing factors should be prioritized in terms of their importance. Several attributes might be used to help in prioritizing:

How important is this factor in contributing to this incident? ⁹
Factors that, if corrected, would likely have prevented the incident. ⁸
Factors that, if corrected, would likely have mitigated the harm.
Factors that if corrected, would not have prevented the incident or mitigated the harm, but are important for patient/staff safety or safe patient care in general
Factors that didn't allow the incident to have more serious consequences and represent solid safeguards that should be kept in place
How important is this factor in contributing to future incidents? ⁸

A summary of the findings is prepared to clearly articulate the contributing factors related to the incident and provide the backbone for development of recommended actions. This summary is provided as a series

⁹ Pham JC et al. ReCASTing the RCA. AJMQ 2010

of "statements of findings". The statements of findings describe the relationships between the contributing factors and the incident and/or outcome. The suggested statement format is as follows: "The contributing factor(s), within the context of the incident, increased/decreased the likelihood that this outcome would occur."

D. Develop Recommended Actions

After summarizing the findings, the facilitator should determine what can be done to reduce the risk of recurrence. Note that in some instances, analyses may not generate any new recommended actions. A few well thought out high-leverage recommendations may ultimately be more effective than a lengthy list of low impact actions. The recommended actions should address the risks and contributing factors identified during the analysis.

Develop Recommended Actions

In order to develop robust, credible, and more precise recommended actions, try to include as many of the features of effective recommended actions below.

- Whenever possible, review the literature for the most evidence-based actions. Aim to use the
 highest level of evidence available (randomized controlled trials are the highest, followed by
 controlled observational studies, uncontrolled studies, opinion of experts and opinion of peers).
 Consider "best practices" that are recommended by professional organizations. In the absence of
 evidence-based recommendations, consider best practices within your organization or at other
 reputable organizations.
- Utilize the most effective solution on the Hierarchy of Effectiveness (see Table 2.2).
- Offer a long-term solution to the problem.
- Actions should have a greater positive than negative impact on other processes, resources, and

Table 2.2 Hierarchy of effectiveness¹⁰

Type of Intervention	Effectiveness
Forcing Function and Constraints	
(e.g. different connectors for oxygen and air)	High leverage – Most effective
Automation/Computerization	
(e.g. automated alerts for drug allergies)	
Simplification/Standardization	
(e.g. standard dosing of antibiotics)	Medium leverage
Reminders, Checklists, Double-Checks	Medium leverage
(e.g. central venous catheter insertion checklist)	
Rules and Policies	Low loverage Least offective
(e.g. policy on patient rounding to assess fall risk)	Low leverage – Least effective (while these are important, they will not result in sustained
Education and Information	practice change when used alone)
(e.g. educate staff on high-alert medications)	practice change when used arolle)

¹⁰ Institute for Safe Medication Practices (ISMP). Medication error prevention "toolbox". ISMP Med Saf Alert. 1999 Jun; 4(11): 1-2.

schedules. Balancing measures should be in place to ensure that unintended consequences are known and understood.

- Provide enough context (explanation and facts) to ensure that if the action is implemented, those responsible for implementing it will understand the rationale behind it.
- Actions are written using the "SMART" format:
 - o Specific tackle a clearly defined issue and have a clear scope;
 - o Measurable can demonstrate impact on process and outcomes;
 - o Assignable can be allocated to one individual to be accountable for implementation;
 - o Realistic ensure that the action is possible; and
 - Timely have a timeframe for implementation.

Where possible, a consultation step may be beneficial in order to ensure that the recommendations are appropriate, the identified risks have been addressed, and there is a high probability to reduce the reoccurrence of this or similar incidents. Providers from the area where the incident occurred, experts, and in some cases patients/families may be consulted.

Discuss recommended actions with leadership/administration

The facilitator discusses the recommended actions with key local decision makers and experts. An assessment of the risk, benefits, costs, and logistics of implementation of the recommended actions are discussed. It is an opportunity to consider the potential for introducing unintended consequences to processes (e.g. creating unnecessary steps or added workload, possibly leading to unsafe workarounds).

Recommended actions that are accepted by organizational leadership become action items for implementation.

E. Implement and evaluate the effect of actions items

Implement the action items

The facilitator, or other person(s) designated by the organization, oversees the implementation status of each action item. Since this is the output of the CIA, it is important that action items are fully implemented as intended.

Evaluate the effect of action items

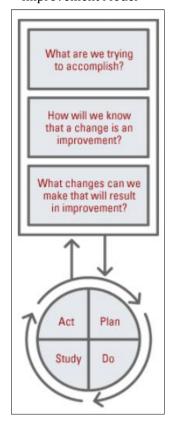
At times, monitoring the effect of action items on patient safety might seem like "just more work." Some action items, though well intentioned and planned, may not have the desired effect in practice. Thus the effectiveness of the implemented action items should be monitored. This will determine if the changes helped make the system safer, had no or limited impact, or actually made the system less safe.

Use of a change management¹¹ or improvement tool can help to facilitate implementation of action items in a way that will support success.¹²¹³ The Improvement Model (Figure 2.4) is a commonly used and effective tool that can be used to provide guidance for establishing measures and tracking progress.

Share learning

The general lessons and findings should be disseminated within, and where applicable, outside the organization to prevent harm recurrence. This is the final objective of the analysis. Without learning and sharing, the organization is still vulnerable as the same or similar incidents could happen again and no other external systems or organizations have the benefit of the learning. Results of analyses should roll up into organization-wide reporting and be shared with the senior leadership, board and the public.

Figure 2.4. Improvement Model



¹¹ Kotter JP, Schlesinger LA. Choosing strategies for change. *Harvard Business Review*. 1979; 57(2): 106-114.

¹² Safer Healthcare Now! *Improvement Frameworks: Getting Started Kit.* Edmonton, AB: Safer Healthcare Now!; 2011 Oct. Available from: https://www.patientsafetyinstitute.ca/English/toolsResources/ImprovementFramework/Documents/Improvement%20Frameworks%20GSK%20EN.PDF

¹³ Boston Consulting Group. *DICE: How to beat the odds in program execution.* 2012. Available from: http://dice.bcg.com/dice.html

CHAPTER 3: WORKBOOK

A. Case selection criteria

Determine if an incident analysis framework is appropriate.

	Yes/No
Is the event thought to be the result of a criminal act?	
Was the event a purposefully unsafe act?	
Was the event related to substance abuse by provider/staff?	
Did the event involve suspected patient abuse?	

If the answers to any of these are clearly yes, do not proceed with an incident analysis framework; rather refer to suitable administrative, professional, or regulatory bodies for resolution.

Determine if a concise incident analysis is appropriate.

	Yes/No
Did the incident result in no or low harm to the patient?	
Is the incident primarily limited to one work area, division, or department?	
Is this a new incident for which a comprehensive analysis was recently completed?	
Is this an incident where you have insufficient information as to whether to perform a	
comprehensive or concise incident analysis?	

If the answer to any of these questions is yes, consider using the concise incident analysis tool.

B. Interview sheets

Interviewee:	Date of Interview
What happened?	
Factors that may have contributed to the incident	
Factors that may have mitigated severity of the incide	nt
How might an incident like this be prevented in the fu	ture?
0 · · · · · · · · · · · · · · · · · · ·	

C. Timeline

Date/Time	Information	Source/Comment

D. Guiding questions

Factor	Relevant?
Task (care/work process):	
Were there previous or predicted failures for this task or process?	
Were specialized skills required to perform the task?	
Was a fixed process or sequence of steps required (e.g. order sets, checklists)?	
Did it exist, and was it followed?	
Was a protocol available, was it up-to-date, and was it followed in this case?	
Were there constraints or pressures (e.g. time, resources) when performing the task?	
Was the information required to make care decisions available and up-to-date (e.g. test results,	
documentation, patient identification)?	
Was there a risk assessment/audit/quality control program in place for the task/process?	
Other?	
Equipment (including information and communication systems):	
Were the displays and controls understandable?	
Did the equipment automatically detect and display problems?	
Was the display functional?	
Were the warning labels, reference guide and safety mechanisms functional and readily visible/accessible?	
Were the maintenance and upgrades up-to-date?	
Was the equipment standardized?	
Would the users describe this equipment as "easy to use?"	
Were the communication systems (phone, pager, software, hardware, etc.) available and operational?	
Other?	
Work environment:	
Did noise levels interfere with the alarms?	
Was the lighting adequate for the task?	
Was the work area adequate for the task(s) being performed (e.g. space, layout, location and accessibility of	
resources)?	
Other?	
Patient(s) characteristics: (Considered in the context of how well the system identified, understand, and acted	
upon these factors. It should not be the only factor considered)	
Did the patient(s) have the information to assist in avoiding the incident?	
If not, what would have supported the patient in assisting their care team?	
Did factors like age, sex, medications, allergies, diagnosis, other medical conditions, contribute to the incident?	
How did they contribute?	
Did any social or cultural factors contribute to the incident?	
What factors? In which way?	
Was language a barrier?	
Other?	
Care team - Caregiver(s):	
Were the education, experience, training and skill level appropriate?	
Was fatigue, stressors, health or other factors an issue?	
Was the workload appropriate?	

Were appropriate and timely help or supervision available?	
Other?	
Care team – Supporting team (all involved in care process):	
Was there a clear understanding of roles and responsibilities?	
Was the quality and quantity of communication (verbal and/or written) between team members appropriate	
(clear, accurate, free of jargon, relevant, complete and timely)?	
Were there regular team briefings/debriefings about important care issues?	
Was team morale good? Do team members support each other?	
Were the communication channels available and appropriate to support the needs of the team (e.g. email, pager, and phone)?	
Other?	
Organization – Policies and priorities:	
Were the relevant policies and procedures available, known, accessible, and did they meet the needs of users?	
Were there workarounds to the documented policy/procedure?	
Was there a mechanism in place to identify and resolve gaps between policy and practice?	
Were the strategic priorities of the organization clear to all?	
Other?	
Organization - Culture:	
Was everyone (patients, clinicians, other staff) comfortable to speak-up about safety concerns?	
Was there visible support from leadership and board for safe patient care?	
Was communication between staff and management supportive of day-to-day safe patient care?	
Were incidents considered system failures with people not blamed?	
Other?	
Organization - Capacity (resources):	
Did scheduling influence the staffing level, or cause stress, fatigue?	
Was there sufficient capacity in the system to perform effectively (e.g. access to resources)?	
Were formal and/or incentives appropriate?	
Other?	
Other – consider:	
Were there any local conditions or circumstances that may have influenced the incident and/or an outcome?	
Were there any sector specific conditions or circumstances that may have influenced the incident and/or	
outcome?	
Other?	

E. Diagramming contributing factors and their interconnection



F. Prioritize and Summarize Findings:

Priority #	Domain (task, equipment, etc.)	Contributing factor	Comment

G. Developing action items

Priority #	Contributing Factor	Recommended Action	Hierarchy of Effectiveness	Strength of Evidence	Costs

H. Oversee implementation of action items

Priority #	Action Item	Measure of effectiveness	Responsible Person	Target Completion Date	Status

I. Incident report template (NOTE: OPTIONAL RESOURCE FOR CONFIDENTIAL HOSPITAL USE ONLY)

1.	Date of Incident (/) 2. Time of Incident (24 HOUR CLOCK) (:)
3.	Date of Incident (/)
4.	Date that the Incident Analysis process was initiated: (//) MM DD YYYY
5.	Date that the Incident Analysis process was completed: (//) MM DD YYYY
6.	Patient's demographic information: Date of Birth: (//) MM DD YYYYY
	Gender
7.	Medical Record Number
8.	What was the patient's principal diagnosis? (Please use ICD code and note country specific ICD modification if possible)
9.	If applicable, what was the patient's principal procedure related to the admission? (Please use ICD code and note country specific ICD modification if possible)

- 10. Select the first applicable category below (in descending order) that best describes the extent of harm to the patient as assessed 24 hours post event.
- a. Death
- b. **Severe permanent harm.** Severe life-long bodily or psychological injury or disfigurement that interferes significantly with functional ability or quality of life
- c. Permanent harm. Life-long bodily or psychological injury or increased susceptibility to disease
- d. Temporary harm. Bodily or psychological injury, but likely not permanent
- e. **Additional treatment.** Injury limited to additional intervention during admission or encounter and/or increased length of stay, but no other injury
- f. **Emotional distress or inconvenience**. Mild and transient anxiety or pain or physical discomfort, but without the need for additional treatment other than monitoring (such as by observation, physical examination, laboratory testing, including phlebotomy, and/or imaging studies).
- g. **No harm.** Event reached patient, but no harm evident

(PATIENT OUTCOME HARM SCALE - Based on a preliminary version of the Agency for Healthcare Research and Quality Harm Scale)

11.	Location of Incident (more than on	Location of Incident (more than one may be selected if applicable)				
	a. Emergency Department	e. Intensive Care Unit / Coronary Care Unit (other high intensity unit)				
	b. Inpatient Unit	f. Long Term Care / Skilled Nursing Facility				
	c. Outpatient Unit	g. \square Other (please specify)				
	d. Pharmacy					
	·					
12.	Was a device or product directly involved in the event?					
	Yes* If Yes, describe the devise or produce and how it was involved					
						
	☐ No					
13.	List the medication(s) directly related to the event. Specify the generic drug name, dose and how the					
	medication was related to the ever	nt (if any).				
	1					
	2					
	2.					
	3					
	4					
	_					
	5					
	6					
	6					
	(add to medication list if required)					
14.	Narrative of EventWhat happened	Narrative of EventWhat happened? Do not include provider or patient identifiable information.				
	(add additional pages as required)					

	Please indicate all that apply and provide a short description of the selected factors and how they contributed to the event (see Guiding Questions)
Con	clusions of incident analysis
16.	a. Following the Incident Analysis, are there recommendations for improving the safety and reliability of
	the applicable care process? Yes
	□ No
	If so, please specify

15. What were the factors that contributed to the occurrence of this event?