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# Medical Errors for Mental Health Professionals 2022 (2CE)

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# **Medical Errors**

A **Medical Error** is a preventable adverse effect of care, whether or not it is evident or harmful to the patient (Grober, 2005). This might include an inaccurate or incomplete diagnosis or treatment of a disease, injury, syndrome, infection, or other ailment. Globally it is estimated that 142,000 people died in 2013 from adverse effects of medical treatment up from 94,000 in 1990 (Institute of Medicine, 2000).

Medical errors can be further defined as the failure of a planned action to be completed as intended or the use of a wrong plan to achieve an aim. Among the problems that commonly occur during the course of providing health care are adverse drug events and improper transfusions, surgical injuries and wrong-site surgery, suicides, restraint-related injuries or death, falls, burns, pressure ulcers, and mistaken patient identities (Carver 2020). High error rates with serious consequences are most likely to occur in intensive care units, operating rooms, and emergency departments.

Medical errors may occur along the continuum of medical care from diagnosis to treatment; they are not intentional, do not always arise to the level of malpractice or negligence, and do not necessarily result in injury to the patient. Those that do are referred to as adverse or sentinel events and must be subjected to rigorous root cause analysis and a response. Even under circumstances in which a medical error did not rise to the level of an adverse or sentinel event, there is often something to be gained by a thorough analysis of the error, why it occurred and what can be done to present a recurrence, and, quite possibly, an adverse or sentinel event.

Beyond their cost in human lives, preventable medical errors exact other significant tolls. Medical errors are estimated to cost between US\$17-billion and US\$29-billion per year in lost income, lost household production, disability and additional health care costs (Grober 2005). Errors also are costly in terms of loss of trust in the health care system by patients and diminished satisfaction by both patients and health professionals. Patients who experience a long hospital stay or disability as a result of errors pay with physical and psychological discomfort. Health professionals pay with loss of morale and frustration at not being able to provide the best care possible. Society bears the cost of errors as well, in terms of lost worker productivity, reduced school attendance by children, and lower levels of population health status.

# **Definitions**

Multiple similar definitions are available for each of these terms from various sources; the health practitioner should be aware of the general principles and probable meaning. The following are common terms and the varying definitions as found in literature:



# Active Error

- Active errors are those taking place between a person and an aspect of a larger system at the point of contact.
- Active errors are made by people on the front line such as clinicians and nurses. For example, operating on the wrong eye or amputating the wrong leg are classic examples of an active error.

# Adverse Event

Adverse events may be preventable when there is a failure to follow accepted practice at a system or individual level.

- An adverse event attributable to an error usually is a preventable adverse event.
- An adverse event is a type of injury that most frequently is due to an error in medical or surgical treatment rather than the underlying medical condition of the patient.
- Not all adverse outcomes are the result of an error; hence, only preventable adverse events are attributed to medical error.

# **Unintended Injury**

- An unintended injury is medical or surgical patient management that results in prolonged hospitalization, measurable physical disability, or both.
  - An unintended injury or complication is the result of prolonged hospitalization or disability or is caused by factors inherent in the healthcare system rather than a disease.

# Latent Error

- Errors in system or process design, faulty installation or maintenance of equipment, or ineffective organizational structure, and may go unnoticed for a long time with no ill effect.
- When a latent error occurs in combination with an active human error, some type of event manifests in the patient. The active human error triggers the hidden latent error, resulting in an adverse event.
- Latent errors are basically "accidents waiting to happen."
- A classic example is a hospital with several types of chest drainage sets, all requiring different connections and setups, yet not all frontline clinicians and nurses are familiar with the intricacies of each setup, creating the scenario for potential error.

# **Medical Error**



- The failure to complete the intended plan of action or implementing the wrong plan to achieve an aim.
- An unintended act or one that fails to achieve the intended outcome.
- Deviations from the process of care, which may or may not result in harm.
- When planning or executing a procedure, the act of omission or commission that contributes or may contribute to an unintended consequence.

# Negligence

- Failure to meet the reasonably expected standard of care of an average, qualified healthcare worker looking after a patient in question within similar circumstances.
- For example, the healthcare worker may not check up on the pathology report which led to a missed cancer or the surgeon may have injured a nerve by mistaking it for an artery.

# Negligent Adverse Events

- A subcategory of preventable, adverse events that satisfy the legal criteria used in determining negligence.
- The injury caused by substandard medical management.

# **Near Miss**

- Any event that could have had an adverse patient consequence but did not.
- Potential adverse events that could have caused harm but did not, either by chance or because someone or something intervened.
- Near misses provide opportunities for developing preventive strategies and actions and should receive the same level of scrutiny as adverse events.

# Never Event

- Never events are errors that should not ever have happened. A classic example of a never event is the development of pressure ulcers or wrong-site surgery. The National Quality Forum has identified the following as Serious Reportable Events:
  - o Care Management
  - Device/Product
  - o Environmental
  - Patient Protective
  - o Surgical
  - o Radiological



# **Noxious Episode**

- Untoward events, complications, and mishaps that result from acceptable diagnostic or therapeutic measures deliberately instituted.
- For example, sending a hemodynamically unstable trauma patient for prolonged imaging studies instead of the operating room. The result could be a traumatic arrest and death.

# **Patient Safety**

The process of amelioration, avoidance, and prevention of adverse injuries or outcomes that arise as a result of the healthcare process.

# Potentially Compensable Event

- An error that could potentially lead to malpractice claims.
- An event due to medical management that resulted in disability, and, subsequently, a prolonged hospitalization.

# **Root Cause**

- A deficiency or decision that, if corrected or avoided, will eliminate the undesirable consequence.
- Common root causes include:
  - Changes in mental acumen including conducting healthcare in an automatic fashion, not seeking advice from peers, misapplying expertise, not formulating a plan, or not considering the most obvious diagnosis.
  - Communication issues, having no insight into the hierarchy, having no solid leadership, not knowing whom to report the problem, failing to disclose the issues, or having a disjointed system with no problem-solving ability.
  - Deficiencies in education, training, orientation, and experience.
  - Inadequate methods of identifying patients, incomplete assessment on admission, failing to obtain consent, and failing to provide education to patients.
  - o Inadequate policies to guide healthcare workers.
  - Lack of consistency in procedures.
  - o Inadequate staffing and/or poor supervision.
  - Technical failures associated with medical equipment.
  - No audits in the system.
  - No one prepared to accept blame or change the system.

# Sentinel Event



- Any unexpected occurrence involving death or serious physical or psychological injury, or the risk thereof...The phrase 'or the risk thereof' includes any process variation for which a recurrence would carry a significant chance of a serious adverse outcome (The Joint Commission, 2017).
- Sentinel events are so-called because once discovered, they frequently indicate the need for an immediate investigation, discovery of the cause, and response.

Definitions retrieved from (Rodziewicz & Hipskind, 2020).

# Types of Errors

A variety of factors have contributed to the nation's epidemic of medical errors. One oft-cited problem arises from the decentralized and fragmented nature of the health care delivery system--or "nonsystem," to some observers. When patients see multiple providers in different settings, none of whom has access to complete information, it becomes easier for things to go wrong.

There are **two types of errors: acts of commission and acts of omission**. An act of commission (doing something wrong) or omission (failing to do the right thing) that leads to an undesirable outcome or significant potential for such an outcome.

For instance, ordering a medication for a patient with a documented allergy to that medication would be an act of commission. Failing to prescribe a proven medication with major benefits for an eligible patient (e.g., low-dose unfractionated heparin as venous thromboembolism prophylaxis for a patient after hip replacement surgery) would represent an error of omission.<sup>3</sup>

According to a report by the Agency for Healthcare Research & Quality, errors of omission are more difficult to recognize than errors of commission but likely represent a larger problem (Kalisch, 2011). In other words, there are likely many more instances in which the provision of additional diagnostic, therapeutic, or preventive modalities would have improved care than there are instances in which the care provided quite literally should not have been provided. In many ways, this point echoes the generally agreed-upon view in the healthcare quality literature that underuse far exceeds overuse, even though the latter historically received greater attention. Clinicians commit acts of **commission** when they make mistakes, such as incorrectly diagnosing someone. Clinicians commit acts of **omission** when they fail to act in some way, such as a failure to report Vulnerable Adult Abuse.



In addition to commission vs. omission, three other dichotomies commonly appear in the literature on errors: active failures vs. latent conditions, errors at the sharp end vs. errors at the blunt end and slips vs. mistakes (AHRQ 2019). Active errors involve frontline personnel and occur at the point of contact between a human and some aspect of a larger system (e.g., a human-machine interface). By contrast, latent errors are accidents waiting to happen—failures of organization or design that allow the inevitable active errors to cause harm. Personnel at the sharp end may literally be holding a scalpel when the error is committed, or figuratively be administering any kind of treatment. The blunt end refers to the many layers of the health care system not in direct contact with patients, but which influence the personnel that come into contact with patients. Slips represent failures of schematic behaviors and occur in the face of competing sensory distractions (i.e. fatigue, stress). Mistakes reflect incorrect choices, and more often reflect lack of experience, insufficient training, or outright negligence (AHRQ 2019).

# **The Joint Commission**

The Joint Commission is a national organization with a mission to, "to continuously improve health care for the public, in collaboration with other stakeholders, by evaluating health care organizations and inspiring them to excel in providing safe and effective care of the highest quality and value"(Joint Commission, 2020). Accreditors play an important role in encouraging and supporting actions within healthcare organizations by holding them accountable for ensuring a safe environment for patients. Healthcare organizations should actively engage in a cooperative relationship with the Joint Commission through this accreditation process and participate in the process to reduce risk and facilitate desired outcomes of care.

The Joint Commission defines a sentinel event as "an unexpected occurrence involving the death or serious physical or psychological injury, or the risk thereof" (Joint Commission, 2020). Serious injury specifically includes loss of limb or function. The phrase 'or the risk thereof' includes any process variation for which a recurrence would carry a significant chance of a serious adverse outcome.

# Root Cause Analysis

Root cause analysis (RCA) is a structured method used to analyze serious adverse events. Initially developed to analyze industrial accidents, RCA is now widely deployed as an error analysis tool in health care. A central tenet of RCA is to identify underlying problems that increase the likelihood of errors while avoiding the trap of focusing on mistakes by individuals (AHRQ 2019). RCA thus uses the systems approach to identify both active errors (errors occurring at the



point of interface between humans and a complex system) and latent errors (the hidden problems within health care systems that contribute to adverse events). It is one of the most widely used retrospective methods for detecting safety hazards.

The Joint Commission defines Root Cause Analysis as "a process for identifying the basic or causal factors that underlie variation in performance, including the occurrence or possible occurrence of a sentinel event" (Joint Commission, 2020).

The sentinel event has resulted in an unanticipated death or major permanent loss of function not related to the natural course of the patient's illness or underlying condition

### or

The event is one of the following (even if the outcome was not death or major permanent loss of function unrelated to the natural course of the patient's illness or underlying condition):

- Suicide of any patient receiving care, treatment, and services in a staffed around-the-clock care setting or within 72 hours of discharge
- Abduction of any patient receiving care, treatment, and services
- Discharge of an infant to the wrong family
- · Rape or sexual assault

RCAs should generally follow a prespecified protocol that begins with data collection and reconstruction of the event in question through record review and participant interviews. A multidisciplinary team should then analyze the sequence of events leading to the error, with the goals of identifying how the event occurred (through identification of active errors) and why the event occurred (through systematic identification and analysis of latent errors.

As part of the accreditation standards, the Joint Commission requires that healthcare organizations have a process in place to recognize these sentinel events, conduct thorough and credible root cause analyses that focus on process and system factors, and document a risk-reduction strategy and internal corrective action plan that includes measurement of the effectiveness of process and system improvements to reduce risk. This process must be completed within 45 days of the organization having become aware of the sentinel event. The Joint Commission will consider a root cause analysis acceptable for accreditation purposes if it focuses primarily on systems and processes, not individual performance. In other words, the healthcare organization should minimize the individual blame or retribution for involvement in a medical error.



In addition, the root cause analysis should progress from special causes in clinical processes to common causes in organizational processes, and the analysis should repeatedly dig deeper by asking why, then when answered, why again, and so on. The analysis should also identify changes that can be made in systems and processes, either through redesign or development of new systems or processes, which would reduce the risk of such events occurring in the future (AHRQ, 2019). The Joint Commission requires that the analysis be thorough and credible.<sup>2</sup>

To be considered **thorough**, the root cause analysis must include:

- 1. Determination of the human and other factors most directly associated with the event and the process(es) and systems related to its occurrence
- 2. Analysis of the underlying systems and processes through a series of "why" questions to determine where redesign might reduce risk
- 3. Inquiry into all areas appropriate to the specific type of event
- 4. Identification of risk points and their potential contributions to this type of event
- 5. A determination of potential improvement in processes or systems that would tend to decrease the likelihood of such events in the future or a determination, after analysis, that no such improvement opportunities exist.

(Oberoi, 2004)

To be considered **credible**, the root cause analysis must meet the following standards:

- 1. The organization's leadership and the individuals most closely involved in the process and systems under review must participate in the analysis.
- 2. The analysis must be internally consistent; that is, it must not contradict itself or leave obvious questions unanswered.
- 3. The analysis must provide an explanation for all findings of "not applicable" or "no problem."
- 4. The analysis must include consideration of any relevant literature.

(Oberoi, 2004)\_

# **Root Cause Analysis and Action Plan Tool Template**

The Joint Commission Root Cause Analysis and Action Plan tool has 24 analysis questions. The following framework is intended to provide a template for



answering the analysis questions and aid organizing the steps in a root cause analysis. All possibilities and questions should be fully considered in seeking "root cause(s)" and opportunities for risk reduction. Not all questions will apply in every case and there may be findings that emerge during the course of the analysis. Be sure however to enter a response in the "Root Cause Analysis Findings" field for each question #. For each finding continue to ask "Why?" and drill down further to uncover why parts of the process occurred or didn't occur when they should have. Significant findings that are not identified as root causes themselves have "roots".

Sentinel Event Settings 2004 through 2014	#	%
Hospital	5749	66.5%
Psychiatric hospital	880	10.2%
Ambulatory care	324	3.8%
Psych unit in general hospital	443	5.1%
Emergency department	478	5.5%
Behavioral health facility	320	3.7%
Home care	165	1.9%
Long term care facility	97	1.1%
Other***	111	1.3%
Office-based surgery	73	0.8%

As an aid to avoid "loose ends," the two columns on the right are provided to be checked off for later reference:

"Root cause" should be answered "Yes" or "No" for each finding. A root cause is typically a finding related to a process or system that has a potential for redesign to reduce risk. If a particular finding is relevant to the event is not a root cause, be sure that it is addressed later in the analysis with a "Why?" question such as "Why did it contribute to the likelihood of the event" or "Why did it contribute to the severity of the event?" Each finding that is identified as a root cause should be considered for an action and addressed in the action plan



• "Plan of action" should be answered "Yes" for any finding that can reasonably be considered for a risk reduction strategy. Each item checked in this column should be addressed later in the action plan.

The Joint Commission Template is Continued on the next page and can be downloaded by clicking here.

# When did the event occur?

Date:	Day of the	Time:	
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Detailed Event Description Including Timeline:

Diagnosis:

Medications:

**Autopsy Results:** 

Past Medical/Psychiatric History:

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	cause	of
			Analysis		Action
			Findings		

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1	What was	List the		
	the	relevant		
	intended	process		
	process	steps as		
	flow?	defined by		
		the policy,		
		procedure,		
		protocol, or		
		guidelines		
		in effect at		
		the time of		
		the event.		
		You may		
		need to		
		include		
		multiple		
		processes.		
		Note: The process steps as they occurred in the event will be entered in the next question. Examples of defined process steps may include, but are not limited to: • Site		
		verification		
		protocol		
		•		
		Instrument,		
		sponge,		
		sharps		
		count		
		procedures		
		•		
		Patient		
		identificatio		



2	Were there	Explain in		
	any steps	detail any		
	in the	deviation		
	process	from the		
	that did not	intended		
	occur as	processes		
	intended?	listed in		
3	What	Discuss		
	human	staff-related		
	factors	human		
	were	performanc		
		e factors		
		that		

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n
	relevant to	contributed to the event.			
	the	Examples may include, but			
	outcome?	are not limited to:			
		<ul> <li>Boredom</li> <li>Failure to follow</li> <li>established policies/</li> <li>procedures</li> <li>Fatigue</li> <li>Inability to focus on task</li> <li>Inattentional blindness/</li> <li>confirmation bias</li> <li>Personal problems</li> <li>Lack of complex critical thinking skills</li> <li>Rushing to complete task</li> <li>Substance abuse</li> <li>Trust</li> </ul>			



4	How did the	Consider all medical
	equipment	equipment and devices used
	equipment	in the course of patient care,
	performanc	including AED devices, crash
	e affect the	carts, suction, oxygen,
		instruments, monitors,
	outcome?	iniusion equipment, etc. in
		information on the following
		as applicable:
		Descriptions of
		hismodical checks
		Diomedical checks
		Availability and
		condition of equipment
		Descriptions of
		equipment with multiple or
		removable pieces
		Location of
		equipment

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n
		and its accessibility to staff			
		and patients			
		Staff knowledge			
		of or education on equipment including			
		applicable competencies			
		Correct			
		operation of alarms.			
		displays, and controls			



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5	What	What environmental factors
	controllable environment	within the organization's
	al factors	control affected the
	affected this	outcome?
	outcome?	Examples may include, but
		are not limited to:
		Overhead paging
		that cannot be heard
		Safety or security
		risks
		Risks involving
		activities of visitors
		Lighting or space
		issues
		The response to this
		question may be addressed
		more globally in Question
		#17. This response should
		be specific to this event.
6	What	Identify any factors the
	uncontrollabl	organization cannot change
	e external	that contributed to a
	factors	breakdown in the internal
	influenced	process, for example
	this	natural

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Finding s		n
	outcome?	disasters.			
7	Were there	List any other factors not yet			
	any other	discussed.			
	factors that				
	directly				
	influenced				
	this				
	outcome?				



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8	What are the	List all other areas in which
	other areas	the potential exists for similar
	in the	circumstances. For example:
	organization	Inpatient surgery/
	where this	outpatient surgery
	could	Inpatient
	happen?	psychiatric care/outpatient
		psychiatric care
		Identification of other areas within the organization that have the potential to impact patient safety in a similar manner. This information will help drive the scope of your action plan.
9	Was the staff	Include information on the
	properly	following for all staff and
	qualified and	providers involved in the
	currently	event. Comment on the
	competent	processes in place to ensure
	for their	staff is competent and
	responsibiliti	qualified. Examples may
	es at the	include but are not limited to:
	time of the event?	<ul> <li>Orientation/ training</li> <li>Competency assessment (What</li> </ul>

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of

Analysis

Findings

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Actio

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		competencies do the staff		
		have and how do you		
		evaluate them?)		
		<ul> <li>Provider and/or staff scope of practice concerns</li> <li>Whether the provider was credentialed and privileged for the care and services he or she rendered</li> <li>The credentialing and privileging policy and procedures</li> <li>Provider and/or</li> </ul>		
10	How did	stan performance issues		
10	How did actual staffing	and actual staffing ratios		
	compare with	along with unit census at the		
	ideal levels?	time of the event. Note any		
		unusual circumstance that		
		occurred at this time. What		
		process is used to		
		determine the care area's staffing		
		ratio,		
		experience level and skill		
		mix?		
11	What is the plan for	Include information on what the organization does		
	dealing with staffing	a staffing crisis, such as call- ins, bad weather or increased		
	contingencie s?	patient acuity.		
		Describe the organization's		
		use of alternative staffing.		
		Examples may include, but are		
		not limited to:		

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n



		Agency nurses		
		Cross training		
		Float pool		
		Mandatory		
		overtime		
10	Mara avala	PRN pool		
12	vvere sucn	It alternative staff were used describe their		
	factor in this	orientation to the area,	-	
			~	
	event?	verification of competency		
		and environmental		
10	D'1	familiarity.		
13	Did staff	Describe whether staff		
	during the	within or outside of the	-	
	event meet	processes. To what extent		
	expectations	was leadership aware of		
	?	any	_	
		performance deviations at		
		the time? What proactive		
		surveillance processes are in	-	
		place for leadership to	-	
		identify deviations from	-	
		expected processes?	-	
		Include	m n	
		omissions in chicar trinking	-	
		and/or performance	-	
		variance(s) from defined		
		policy, procedure, protocol	-	
		and guidelines in effect at		
		the time.		
14	To what	Discuss whether patient		
	degree was	assessments were		
	all	completed,	-	
	the necessary	shared and accessed by	-	
	information	members of the treatment		
	available when	team, to include providers,		
	needed?	according to the		
	Accurate?	organizational processes.		



Complete?	Identify the information			
-----------	--------------------------	--	--	--

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysi	е	Actio
			s		n
			Finding		
			S		
	Unambiguou s?	systems used during patient			
		care.			
		Discuss to what extent the available patient information			
		(e.g. radiology studies, lab			
		results or medical record) was clear and sufficient to			
		provide an adequate			
		condition, treatment and			
		response to treatment.			
		Describe staff utilization and			
		procedure, protocol and			
		guidelines specific to the			
		patient are provided			
15	To what	Analysis of factors related to			
	degree was	communication should			
	the	include evaluation of verbal,			
	communicati	written, electronic			
	on among	communication or the lack			
	participants	thereof. Consider the			
	adequate for	following in your response,			
	this	as appropriate:			
	situation?	• The timing of			
		communication of key			
		information			
		Misunderstandings			
		related to language/cultural			
		barriers, abbreviations,			
		terminology, etc.			
		Proper completion of			
		internal and external hand-			
		off communication			
		Involvement of patient,			
		family and/or significant			



#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n
16	Was this the	Consider processes that			
	appropriate	proactively manage the			
	physical	patient care environment.			
	environment	This response may			
	for the	correlate to the response in			
	processes	question 6 on a more global			
	being carried	scale.			
	out for this	What evaluation tool or			
	situation?	method is in place to			
		and mitigate physical and			
		patient care environmental			
		How are these process			
		needs addressed			
		organization-wide?			
		Examples may include, but			
		are not limited to:			
		alarm audibility			
		testing			
		evaluation of			
		egress points			
		patient acuity			
		level and setting of care			
		managed across the			
		continuum,			
		• preparation of			
		medication outside of			
		pharmacy			
17	What	Identify environmental risk			
	systems are	assessments.			
	in place to	Does the			
	identify	current environment meet codes, specifications.			
	environment	regulations?			
	al risks?	<ul> <li>Does staff know</li> <li>how to report environmental</li> </ul>			
		risks?			



#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n
		Was there an			
		environmental risk involved			
		in the event that was not			
		previously identified?			
18	What	Describe variances in			
	emergency	expected process due to an			
	and failure-	actual emergency or failure			
	mode	mode response in			
	responses	connection to the event.			
	have been	Related to this event, what			
	planned and	safety evaluations and drills			
	tested?	have been conducted and at what frequency (e g			
		mock code blue, rapid			
		response, behavioral			
		abduction or patient			
		elopement)?			
		Emergency responses may			
		include, but are not limited			
		to:			
		• Fire			
		External disaster			
		• Mass casually $\Delta$ Medical emergency			
		Failure mode responses			
		may include, but are not			
		limited to:			
		Computer down			
		time			
		Diversion			
		● Facility			
		construction			
		Power loss     Utility issues			
19	How does	How does the overall			
	the	culture?			
	organization' s	encourage change?			

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n



	<u> </u>			
	culture	suggestions and warnings		
	support risk	from staff regarding risky		
	reduction?	situations or problematic		
		areas?		
		<ul> <li>How does leadership demonstrate the organization's culture and safety values?</li> <li>How does the organization measure culture and safety?</li> <li>How does leadership establish methods to identify areas of risk or access employee suggestions for change?</li> <li>How are changes implemented?</li> </ul>		
20	What are the	Describe specific barriers to		
	barriers to	effective communication		
	communicati	among caregivers that have		
	on of	been identified by the		
	potential risk	organization. For example,		
	factors?	residual intimidation or		
		reluctance to report co-		
		worker activity.		
		Identify the measures being taken to break down barriers (e.g. use of SBAR). If there are no barriers to communication, discuss how this is known.		
21	How is the prevention of	Describe the organization's adverse outcome procedures		

#	Analysis	Prompts	Root	Root	Plan
	Question		Cause	caus	of
			Analysis	е	Actio
			Findings		n
	adverse	and how leadership plays a			
	outcomes	role within those			
	communicat	procedures.			
	ed as a high				
	priority?				



#### Online Continuing Education for Professionals 22 How can Describe how orientation orientation and ongoing education and inneeds of the staff are service evaluated and discuss its training be relevance to event. (e.g. revised to competencies, critical reduce the thinking skills, use of risk of such simulation labs, evidenceevents in the based practice, etc.) future? 23 Was Examples may include, but available are not limited to: CT scanning technology • equipment used as Electronic • intended? charting Medication delivery system Tele-radiology ٠ services 24 How might Describe any future plans for implementation or technology be redesign. introduced or Describe the ideal technology system that can redesigned to reduce help mitigate potential risk in the adverse events in future? the future.

Action Plan	Organization Plan of Action Risk Reduction Strategies	Position/ Tit le Responsi ble Party	Method: Policy, Educati on, Audit, Observ ation on & Implem ent action
For each of the findings identified	Action Item #1:		
in the analysis as needing an	Action Item #2:		



Online Continuing Education	on for Professionals	
action, indicate	Action Item #3:	
the planned		
action expected,		
implementation		
date and		
associated		
measure of		
OR		
If after		
consideration of		
such a finding, a		
decision is made		
not to implement		
an associated risk		
reduction		
strategy, indicate		
not taking action		
at this time		
Check to be sure		
that the selected		
measure will		
provide data that		
will permit		
assessment of		
the		

Action item #4.		
Action Item #5:		
Action Item #6:		
Action Item #7:		
Action Item #8:		
	Action Item #4: Action Item #5: Action Item #6: Action Item #7: Action Item #8:	Action Item #4:



# Bibliography: Cite all books and journal articles that were considered in developing this root cause analysis and action plan.

# Misdiagnosis of Mental Health/ Psychiatric Disorders

Regarding mental illnesses, sufferers of dissociative identity disorder usually have psychiatric histories that contain three or more separate mental disorders and previous treatment failures. The disbelief of some doctors around the validity of dissociative identity disorder may also add to its misdiagnosis.<sup>5</sup>

Studies have found that bipolar disorder has often been misdiagnosed as major depression. Its early diagnosis necessitates that clinicians pay attention to the features of the patient's depression and also look for present or prior hypomanic or manic symptomatology.

The misdiagnosis of schizophrenia is also a common problem. There may be long delays of patients getting a correct diagnosis of this disorder. The DSM-5 field trials included "test-retest reliability" which involved different clinicians doing independent evaluations of the same patient—a new approach to the study of diagnostic reliability.

# **Patient Suicide**

The Joint Commission issued a Sentinel Event Alert on preventing inpatient suicides; this Alert updates the prevention strategies presented in that Alert with a focus on general hospitals and prevention of suicide in medical/surgical units and the emergency department. The goal of this Alert is to assure that patients outside of psychiatric units are appropriately screened and cared for. In addition to non-psychiatric settings, the Sentinel Event Database includes reports of suicide in psychiatric hospitals, behavioral health units of general hospitals, and residential treatment facilities. While psychiatric settings are designed to be safe for suicidal individuals and have staff with specialized training, typically, medical/surgical units and emergency departments are not designed or assessed for suicide risk and do not have staff with specialized training to deal with suicidal individuals. Not surprisingly, suicidal individuals often are admitted to general hospital emergency departments often at the urging of families or friends – when they are most desperate.



These patients are "known at risk" for suicide. It is noteworthy that many patients who kill themselves in general hospital inpatient units do not have a psychiatric history or a history of suicide attempt – they are "unknown at risk" for suicide. Compared to the psychiatric hospital and unit, the general hospital setting also presents more access to items that can be used to attempt suicide – items that are either already in or may be brought into the facility and more opportunities for the patient to be alone to attempt or re-attempt suicide. This Alert presents strategies that can be used and suggested actions that can be taken by general hospitals to help better prepare their staffs and their facilities for suicidal patients and to care for both their physical and mental needs.

The location of the events included bathroom, bedroom, closet, shower and other locations, or they occurred after discharge or leaving the hospital against medical advice. The methods of suicide included hanging, asphyxiation by other than hanging, gunshot, jumping from a height, drug overdose, laceration, drowning, other methods (e.g., jumping in front of a moving vehicle, ingestion of poison, stabbing or burning).

Risk factors for suicide: The risk factors common across health care settings include having previously attempted suicide; recent suicide attempt; suicidal thoughts or behaviors; a family history of suicide or psychiatric illness; on antidepressants; physical health problems, including central nervous system disorders such as traumatic brain injury; diagnosis of delirium or dementia; chronic pain or intense acute pain; poor prognosis or prospect of certain death; social stressors such as financial strain, unemployment or loss of financial independence; disability; trauma; divorce or other relationship problems; hopelessness; and substance abuse. Substance abuse may also exacerbate psychological symptoms such as depression, and the disinhibitory effects of alcohol may contribute to impulsive suicidal behavior. Older adults are prone to additional suicide risk factors including declining health, loneliness and recent bereavement.

Warning signs that are associated with increased desperation and imminent risk include: irritability, increased anxiety (in addition to panic), agitation, impulsivity, decreased emotional reactivity, complaining of unrelenting pain, refusing visitors, crying spells, declining medications, and requesting early discharge. In addition, the following warning signs are diagnostic criteria for depression: hopelessness or helplessness, decreased interest in treatment or prognosis, feelings of worthlessness, and refusing to eat.

There are numerous medications that are associated with an increased risk of suicidal thoughts and behaviors, including antidepressants, antiepileptic or anticonvulsant medicines, and antipsychotic agents. The risk of suicidal thoughts and behaviors applies to both psychiatric and non-psychiatric uses of these



medications. The American Society of Health-System Pharmacists (ASHP) maintains a list of such medications based on FDA alert. Certain other medicines have also been associated with increased risk of suicide, such as some smoking-cessation drugs, anti-infectives (e.g. Mefloquine, interferons, amantadine), and others (e.g., isotretinoin). The increased risk can be associated with use of the medicine either as an inherent risk of an underlying psychiatric or other illness (e.g., epilepsy) or a side effect of the medication itself.





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B. Jones, has a Bachelors in Psychology (Cum Laude) and a Juris Doctor in Law. She is an attorney licensed by the Florida Bar with a focus on healthcare/ mental health law. While in law school she received numerous accolades such as: Pro Bono Honor Program, Gold Level (300+ Pro Bono Hours), ILSA Journal of International & Comparative Law, Junior Staff Editor Dean's List (Fall 2011; Winter 2014. B. Jones was the research assistant to Nova Southeastern, Law Professor, Elena Langan and assisted with nomenclature changes to family law statutes around the country, searched relevant court decisions, statutory changes and journal articles.

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Fiona Gain has a Bachelors Degree in environmental science from the University of Florida, a Masters. Degree from Johns Hopkins University. She has experience researching policy-based topics and has been working with ACE classes to gain expertise in medical policy and information.