# Online Supplement to "A Perioperative Intervention to Prevent and Treat Emergence Delirium at a Veteran Affairs Medical Center"

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This supplementary material was provided by the authors to give readers additional information and resources to replicate their work.

### How to Cite the Original Article

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### Supplemental Materials S1–S15 Materials Supporting the Intervention to Prevent and Treat Emergence Delirium

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### Supplemental Material S1 Essentials of the Intervention

### How material is used: Clinical reference

**Purpose**: Identifies information integral to the safe and effective use of the intervention. This can be used as a handout during staff training

### **Essentials of the Intervention**

### **Clinical Components and Phases**

- The intervention consisted of 21 clinical components that were used across six phases, beginning with the *patient evaluation with individualized plan* phase and ending with the *actions following an episode of emergence delirium* phase.
- For a list of the 21 clinical components by phase, see supplemental material **S2**. The **S2** material identifies additional supplemental materials corresponding with the intervention and training of staff.

PAASO Form: Preoperative Anesthesia Assessment, Medication Strategy, and Outcome

- See supplemental material **S3**.
- Objectives for use of the form: identify (screen) patients who are at an elevated risk for emergence delirium, gather information to orient the patient upon emergence, facilitate patient hand-offs between staff, guide and document the medication strategy, and facilitate collection of information for the patient's record. Additionally, the form can be used to improve the continuity of care, track and trend the intervention integrity (i.e., implementation integrity) across staff, and monitor the overall effectiveness of intervention.
- Without routine screening for risk factors, anesthesia providers may not adjust their individualized anesthesia care plan to account for the patient's risk status and this may create an elevated risk of emergence delirium and patient/staff harm.

### **Communication Between Staff**

- Communication is important to the success of the intervention and is used in each of the phases identified in supplemental material **S2**.
- Most of the communication between staff related to the intervention occurs during transitions from one phase to another and during a response to emergence delirium.
- Make sure that all staff within the chain of care of the patient are informed of pertinent information regarding risk for ED, potential triggers, patient's preferred waking method, and any other information that may help orient the patient during emergence.

### **Adjust Environment**

- Create a low stimulation environment for the patient by placing them in a lower traffic area (e.g., private bay), dimming the lights, reducing noise, and consolidating/limiting unnecessary staff interactions (e.g., avoid interactions with trainees).
- Reduce the risk of patient harm by adjusting the table or bed (e.g., reduce height to increase leverage, lock wheels, pad side rails, secure arm boards and /or stirrups) and secure the intravenous (IV) site (e.g., extra tape and gauze).

### Medication Strategy for Patients at Elevated Risk for Emergence Delirium

- See supplemental material **S5** for details about the medication strategy (e.g., sequence, dose, rate) and see supplemental material **S6** for a literature review in support of the medication strategy.
- Based on previous studies of emergence delirium, we recommend avoidance of midazolam and volatile anesthetics, and, as an alternative, administration of propofol, dexmedetomidine, and ketamine.
- Although midazolam is a universally administered preop anxiolytic, it is also a risk factor for emergence delirium, which is why it is absent from this medication strategy.
- Dexmedetomidine, if not otherwise contraindicated for cardiac impairment, is an excellent alternative to midazolam.
- To lessen the likelihood of emergence delirium, we advocate for administration of ketamine under the following conditions: 1) given at induction, pre-incision, and/or post-incision; 2) at a low or "analgesic" dose of <0.5 mg/kg, and 3) paired with an active anesthetic agent (e.g., propofol, volatile agent) and/or an active sedative adjunct (e.g., dexmedetomidine). Additionally, we recommend no more than 50 mg IV maximum for ketamine. Finally, we do not recommend administering ketamine preemergence (late in the case), as this may increase the likelihood of emergence delirium.
- We recommend treating **low-to-moderate-risk behavior** (e.g., inconsolable, disinhibited, restless, anxious, hallucinating) with **dexmedetomidine**.
- We recommend treating **moderate-to-high-risk behavior** (aggressive, angry, agitated, thrashing, kicking, attempting to exit bed, and other dangerous behavior) with **propofol**.

### **Patient Orientation Upon Emergence**

- Mention familiar people and places. E.g., patient's name and nickname, names of immediate family members, pet's name, and location.
- Use patient's preferred physical interactions during emergence. E.g., tap on right shoulder and do not touch legs. Many staff have reported that patients will often express concern about being startled and not wanting to hurt anyone.
- Choose staff gender for the nursing care team consistent with the patient's preference. Some patients may have a history of sexual abuse/trauma and may have a strong gender preference of staff on the nursing care team. For example, a patient who has a history of sexual assault by a male perpetrator may prefer a female nursing team.
- Speaking in a softer tone of voice may help when assisting a patient during the emergence and reorientation process.

### Actions in Response to an Episode of Emergence Delirium

- Call "Anesthesia stat" for additional support.
- Monitor for medical complications and use caution related to the surgical site.
- Monitor and protect IV site and airway.
- Administer medication, if necessary, to help manage patient behavior. See supplemental material S5.
- Manual (hands-on) restraint, if necessary
  - The purpose of manual restraint is to prevent the patient, who is engaging in dangerous behavior, from harming themselves and staff.
  - Manual restraint refers to staff placing their hands directly on the patient to secure a limb or area of the body. As a part of this intervention, staff must use specific techniques taught by experts through the lecture and hands-on (i.e., guided practice and simulation) training.
    - Techniques that may have been learned elsewhere are not permitted and are deemed unsafe.
    - Note that mechanical restraint, such as belts, straps, and cuffs, are different from manual restraint techniques.
    - Restraints are clinical interventions and therefore must be authorized by the appropriate level of clinical staff.
  - A minimum of four staff should be involved in managing and treating patient behavior. Work as a team; no one person should be responsible for manually restraining a patient. There must be an observer (a medically trained staff member who is not physically participating in the restraint) present to assess the safety of the patient and alert the responding team to any safety issue or medical distress occurring with the patient.
  - The manual restraint techniques focus on management of the patients' arms, legs, head, oxygen apparatus, and IV location. This section is not intended to be a "how-to" restraint application guide.
    - Arms Pressure will be applied to the shoulder area and below the elbow (forearm) of both arms. At no time should pressure be applied to the elbow joint. The arms will remain in line with the body and will not be manipulated in a way that produces pain to force compliance. The arms will not be maneuvered away from the body or exceed the patient's natural range of motion.
    - Legs Pressure will be applied to the thighs and shin areas. The legs can typically support pressure that
      is placed above or below the knees, but not directly on the knees.
    - Head It may be necessary to support the head of the patient in a way that will prevent the patient from
      removing or dislodging the oxygen apparatus. Pressure may be placed on the patient's forehead and
      under the chin, but at no time should weight be applied to the neck (i.e., do not use a choke hold).
    - IV site The area around the IV site will need to be secured to prevent IV removal. This will be
      accomplished by cupping a hand over the IV site to prevent dislodgment.
    - General guidelines –Please note that learners may have difficulty gauging the difference between
      adequate and excessive force when restraining a patient; therefore, this topic should be emphasized
      during training.
  - Patient positions
    - Recommended
      - Supine
      - Side patients who are pregnant or obese
      - Standing escort technique apply only if patient exits the table or bed
    - Not recommended
      - Prone
  - Risks with manual restraint
    - Positional asphyxiation To prevent asphyxiation, weight should never be applied to the patients' thoracic cavity/abdominal area, lower/upper back, or neck area. Do not lie across the patient's upper body, which may restrict the patient's respiration.
    - Aspiration The risk of aspiration should be low due to patient's nothing by mouth (NPO) status, but if the patient does aspirate during restraint, medical intervention must take priority over the restraint in order to clear the airway and ensure the safety of the patient.

- Orthopedic injuries Do not exceed the patient's range of motion and do not use joint lock or pain compliance techniques.
- Skin integrity Apply a measured amount of force during restraint and do not default to full force.

### Actions Following an Episode of Emergence Delirium

- Debrief meeting with patient and family
  - Helping the patient process the experience is important. The concern is that patients who experience emergence delirium may be less likely to schedule a future procedure, which may have a long-term negative impact on their health (e.g., delayed diagnosis, treatment, and/or therapy).
  - Provide the patient with a prepared document explaining emergence delirium.
  - Prior to discharge provide the patient with a referral or contact information to a mental health service.
- Debrief meeting among staff
  - Discuss alternative explanations for the delirium.
  - Do we know what triggered the episode?
  - Could the episode have been prevented?
  - Discuss team response and what should be improved.
- Documentation
  - File an event report with the patient safety office or risk management.
  - Add a detailed note to the patient's record.

### **Supplemental Material S2**

### Clinical Components of the Perioperative Intervention to Prevent and Treat Emergence Delirium

How material is used: Clinical reference

**Purpose**: This table outlines the use of 21 clinical components across six phases, beginning with the *patient evaluation with individualized plan* phase and ending with the *actions following an episode of emergence delivium* phase

Phases		<b>Clinical Components</b>	Supplemental Material
Patient	1.	Use the PAASO Form to conduct a preoperative anesthesia assessment	<b>▲</b> S1–S3, S9
evaluation		A. Review patient's record and interview patient to assess (screen) for	
with		emergence delirium (ED) risk factors	
individualized		B. Meet with the high-risk patient and family to gather information to	
plan		facilitate wake-up	
	2.	Communication; call ahead to procedure room to initiate intervention	
		protocol	<b>◄</b> S1, S2, S9
Preoperative	3.	Communication; patient wears a unique colored surgical cap and bracelet to	<b>◄</b> S1, S2, S9
actions		help staff recognize them as high-risk for ED	
	4.	Adjust environment	<b>◄</b> S1, S2, S9
		A. Low stimulation	
		B. Prepare equipment/apparatuses to mitigate risk in the event of	
		dangerous behavior	
	5.	Follow medication strategy and document in the PAASO form	<b>◀</b> S1-S6, S9
	6.	Communication during time-out	<b>◄</b> S1, S2, S9
		A. Remind staff of patient's high-risk for ED	
		B. Inform staff of IV location and type of airway (e.g., O2 mask,	
		endotracheal tube)	
Intraoperative	7.	Follow medication strategy and document in the PAASO form	<b>◀</b> \$1-\$6, \$9
actions	8.	Communication; as case concludes, call the recovery room/post-anesthesia	<b>◄</b> S1, S2, S9
		care unit to initiate intervention protocol	
Postoperative	9.	Adjust environment	<b>◄</b> S1, S2, S9
actions		A. Low stimulation	
		B. Prepare equipment/apparatuses to mitigate risk in the event of	
		dangerous behavior	
	10.	Communication; consider proactively requesting additional staff due to	<b>◄</b> S1, S2, S9
		anticipated risk for ED	
	11.	Staff should be prepared to protect IV access and maintain the airway	<b>◄</b> S1, S2, S9, S10, S12, S13, S15
	12.	Follow medication strategy and document on the PAASO form	
	13.	Use the PAASO form to document the patient's outcome and note in	<b>◀</b> S1–S6, S9
		patient's record	<b>▲</b> S1–S3, S9
Actions in	14.	Communication; if needed, call for immediate staff support by using a	<b>◀</b> S1, S2, S9, S10, S13, S15
response to an		standardized code (e.g., "Anesthesia stat")	
episode of	15.	If needed, staff should apply manual restraint (hands-on) to the patient to	<b>◀</b> \$1, \$2, \$9-\$15
emergence		mitigate risk of harm to the patient and staff (note: special training and	
delirium		technique are required to ensure safety)	
	16.	If needed, follow medication strategy	<b>◀</b> S1, S2, S4–S6, S9, S10, S13,
			S15
	17.	Attempt to orient the patient by stating familiar people and places (e.g.,	<b>◀</b> \$1–\$3, \$9, \$11, \$13–\$15
		patient name, partner name, location of hospital)	
Actions	18.	Debrief meeting with patient and family, which includes providing them	<b>◄</b> S1, S2, S9
following an		with a prepared brochure about emergence delirium and a referral to	
episode of		behavioral health, if desired	
emergence	19.	Debrief meeting among staff to discuss alternative explanations for the ED	<b>◄</b> \$1, \$2, \$9
delirium		and the effectiveness of their intervention	
	20.	If applicable, in the PAASO form document any medications administered in	<b>◀</b> \$1–\$3, \$9
		response to emergence delirium	
	21.	If applicable, file an event report with the patient safety office or risk	<b>◀</b> \$1–\$3, \$9
		management, and add a detailed note to the patient's record	

Note: The supplemental materials consist of information that either directly pertains to staff skill development (training or clinical reference) or are a tool integral to the clinical component.

### **Supplemental Material S3**

### PAASO Form: Preoperative Anesthesia Assessment, Medication Strategy, and Outcome

### How material is used: Clinical documentation

**Purpose**: The form will be used to facilitate identification of patients who are at an elevated risk for emergence delirium, gather information to orient the patient upon emergence, facilitate hand-offs, guide and document the medication strategy, facilitate collecting information for patient record, and monitor effectiveness of the intervention across providers

Durana	PAASO Form (v 6/17/2022)
Freope	Date: Procedure:
	Anosthosia Dlan (airala as annlias):
	Local MAC General Regional
	History of Difficult Airway (circle one)? No Yes describe:
[Patient Sticker]	History of Difficult All way (choice one). No res, describe.
	Allergies (circle one)? NKDA Yes, describe:
	ASA Status (circle one): I II III IV V (E)
Part 1: Preoperative Anesthesia Assessment	<b>Staff who complete part 1</b> : Anesthesia provider <b>Purpose</b> : Identify patients who are at elevated risk for emergence delirium, further refine the medication strategy, gather information to orient the patient upon emergence, and facilitate
	hand-offs
1.1. Based on chart review and patie	ent interview, is there a history of the following symptoms, conditions, or experiences
(potential risk factors for emergenc	e delirium)?
NONE     Previous occurrence of emerges	Depression     Depression
flashbacks hallucinations scree	$\Box$ Anxiety
else)	Sleep or mood disorder
□ Post-traumatic stress disorder (]	2TSD)
□ Hypervigilance	$\Box$ Traumatic event (e.g., sexually assaulted.
□ Flashbacks	domestic violence. combat)
□ Irritability	$\square$ Substance use
□ Exaggerated startle response	□ Other:
12 Deced on about unions and not	nt interview what is now activate of the metions? wish for an encourse a delivious?
1.2. Based on chart review and path	$\Box Low \Box Moderate \Box High$
1.3. Based on chart review and patie	ent interview, are there any current medications and/or substances used (drugs and
alcohol) that may impact the effecti	veness of analgesic or sedative agents? $\Box No$ $\Box Yes$ . list:
Questions to Facilitate Wake-Up	
1.4A. At home, do you often wa	ke up confused, angry or violent?
$\Box No  \Box Yes$ , list:	
<b>1.4B. Is there anything specific</b> touch me, only touch your left sh	I should do or not do ( <i>triggers</i> ) to wake you up at the end of your procedure? E.g., Don't oulder, gender preference for nursing care team, etc.
1.4C. Are there any names that	I can use to help orient you when you wake up? E.g., your nickname, family member, pet
Part 2: Medication Strategy	<b>Stall who complete part 2</b> : Anesthesia provider <b>Purpose</b> : Facilitate hand-offs and monitor the perioperative intervention across department providers

2.1. Medications administered durin	ng pre-operation, induction.	Dexmed	etomidine	Propofol
and/or maintenance of anesthesia ph	ases		l	Remifentanil
			amhana	
			orphone	
			ie	□ Volatile agent
		□ Acetami	nophen	□ Oral opiates
2.2. Preemergence		□ NONE		Hydromorphone
		□ Dexmed	etomidine	□ Propofol
2.3. If applicable, in response to eme	ergence delirium	□ NONE		Propofol
		□ Dexmed	etomidine	□ Midazolam
	Staff who complete part 3: Anesth	nesia provide	er or PACU nurse	3
Part 3: Outcome	Purpose: Monitor effectiveness of	the intervent	ion across depar	tment providers and facilitate
	collecting information for patient re	ecord	-	-
3.1. Was there an episode of emerge	ence delirium? E.g., violence,			
thrashing, aggression, combativeness,	screaming, hallucinating,		- Var	
attempting to exit bed/table, and/or att	tempting to remove medical			
apparatuses				
If yes emergence delirium (3.1):		$\Box No$	□ Yes. descri	be:
<b>3.2A.</b> Was the behavior dangerou	us?		,	
3 2B Was anyone injured?		$\Box No$	<b><i>□</i>Ves</b> descri	he
5.2D. Was anyone injured.				
<b>3.2C. What may have triggered t</b>	he episode? E.g., loud sound,	Describe:		
bright light, or tactile stimulus				
Anesthesia Provider:	Date:			
PACU Nurse:				

Instructions for U Preoperative Anesthesia Asses	Using PAASO Form (v 6/17/2022) sment, Medication Strategy, and Outcome
Part 1: Preoperat Purpose: Identify patients who are at elevated risk for emerge to orient the patient upon emergence, and facilitate hand-offs Who completes part 1: Anesthesia provider	ive Anesthesia Assessment nce delirium, further refine the medication strategy, gather information
Questions from PAASO Form	How information will be utilized
1.1. Based on chart review and patient interview, is there a history of the following symptoms, conditions, or experiences (potential risk factors for emergence delirium)?	Identify factors that could be associated with an increased likelihood of emergence delirium
1.2. Based on chart review and patient interview, what is <i>your estimate</i> of the patient's risk for emergence delirium?	<ul> <li>Develop and prepare a team response, if warranted</li> <li>Improve the continuity of care between providers and/or staff</li> <li>Track and trend the intervention plan across department providers</li> </ul>
1.3. Based on chart review and patient interview, are there any current medications and/or substances used (drugs and alcohol) that may impact the effectiveness of analgesic or sedative agents?	This information will be used to guide the <i>medication strategy and the</i> <i>amount of medication administered</i>
<ul> <li>Questions to Facilitate Wake-Up</li> <li>1.4A. At home, do you often wake up confused, angry or violent?</li> <li>1.4B. Is there anything specific I should do or not do to wake you up at the end of your procedure?</li> <li>1.4C. Are there any names that I can use to help orient you when you wake up?</li> </ul>	<ul> <li>(Q 1.4A) Guide the preemergence medication strategy and the amount of medication administered. I.e., challenging behaviors may indicate a need for additional medications and/or a greater dose</li> <li>(Qs 1.4A, 1.4B) Identify stimuli (triggers) that could be associated with an increased likelihood of emergence delirium</li> <li>(Qs 1.4A, 1.4B, 1.4C) Develop and prepare a team response, in anticipation of emergence delirium</li> </ul>
Part 2: M Purpose: Facilitate hand-offs and monitor the perioperative in Who completes part 2: Anesthesia provider	edication Strategy tervention across department providers
Questions from PAASO Form	How information will be utilized
<ul> <li>2.1. Medications administered during <i>preoperation</i>, <i>induction</i>, <i>and/or maintenance of anesthesia phases</i></li> <li>2.2. <i>Preemergence</i></li> <li>2.3. If applicable, in <i>response to emergence delirium</i></li> </ul>	<ul> <li>Improve the continuity of care between providers and/or staff</li> <li>Track and trend the intervention plan across department providers</li> </ul>
Part	t 3: Outcome
<b>Purpose</b> : Monitor effectiveness of the intervention and facilita <b>Who completes part 3</b> : Anesthesia provider or PACU nurse	ate collecting information for patient record
Questions from PAASO Form	How information will be utilized
3.1. Was there an episode of emergence delirium?	<ul> <li>Collect information that will be entered into patient record and will guide provider practice during future procedures with the same patient</li> <li>Track and trend the intervention plan across department providers</li> </ul>
It yes emergence delirium: 3.2A. Was the behavior dangerous? 3.2B. Was anyone injured? 3.2C. What may have triggered the episode?	• If emergence delifium is observed, this information will facilitate a debrief meeting among staff and could be used to develop a patient safety event report

### **Supplemental Material S4** Timeline of Anesthesia Activities and Perioperative Care

### How material is used: Clinical reference

**Purpose**: This information is intended to help people better understand patient status and staff responsibilities across the perioperative phases. In particular, we anticipate that readers will find the following information useful: stages of anesthesia, risk of emergence delirium (ED), and medication strategies.

Phases	Patient Status and Staff Responsibilities for Intravenous (IV)/Monitored Sedation or General Anesthesia With or Without Regional Anesthesia
<b>Preoperation</b> <b>Preparation</b> Immediately prior to transfer to operating room	<ul> <li>Stage of anesthesia: None</li> <li>Duration: 5–15 min</li> <li>Risk of emergence delirium: None to low</li> <li>Commonly administered medications: Midazolam (Versed), dexmedetomidine, and/or fentanyl</li> <li>Our medication strategy: None or acetaminophen, dexmedetomidine, and/or an opioid (oral or IV), to reduce anxiety as needed. For more information about these medications, see supplemental materials S5 and S6.</li> <li>Desired patient status: Nonsedated or transitioning from conscious and communicative to lightly sedated, but remains voluntarily participative</li> <li>Staff responsibilities: Transition patient from preoperative area to surgical or procedural suite. Move patient from stretcher to operating room bed. Alarms and monitors are applied and activated. Supplemental oxygen is administered, as required.</li> </ul>
Induction of Anesthesia Administration of an agent to provide sedation or general anesthesia as necessary	<ul> <li>Stage of anesthesia: Transition from stage 1 to stage 2</li> <li>Duration: 2–5 min</li> <li>Risk of emergence delirium: Low to moderate during transition through stage 2, which is the stage with greatest risk of ED. Duration of stage 2 is typically greater with IV/monitored sedation than general anesthesia.</li> <li>Commonly administered medications: Midazolam, fentanyl, lidocaine, and propofol</li> <li>Our medication strategy: Lidocaine, propofol, ketamine, and hydromorphone. For more information about these medications, see supplemental materials S5 and S6.</li> <li>Desired patient status: Transitioning to a state of amnesia and analgesia</li> <li>Staff responsibilities: Support and maintain adequate vital signs; ventilatory support; and administer adjunct medications, as required, in preparation to achieve a surgical plane of anesthesia</li> </ul>
Maintenance of Anesthesia Maintain desired level of anesthesia required for the procedure	<ul> <li>Stage of anesthesia: 3</li> <li>Duration: Dependent on procedure</li> <li>Risk of emergence delirium: Low, depending on integrity of anesthesia delivery to maintain adequate depth of anesthesia</li> <li>Commonly administered medications: Propofol infusion (total intravenous anesthesia) and/or volatile anesthetic (inhalational), depending on type of anesthesia</li> <li>Our medication strategy: Propofol infusion with or without adjunct agents (e.g., dexmedetomidine, ketamine, hydromorphone), depending on type of anesthesia. For more information about these medications, see supplemental materials S5 and S6.</li> <li>Desired patient status: Unconscious; however, spontaneous respirations are preserved during IV/monitored sedation and respirations are often controlled during general anesthesia</li> <li>Staff responsibilities: Continuously reassess vital signs and administer maintenance and adjunct medications to preserve the appropriate level of anesthesia</li> </ul>

<b>Emergence</b> Patient awakens from anesthesia leading to the greatest potential for delirium or excitement	<ul> <li>Stage of Anesthesia: Transition from stage 3 to stage 2 to stage 1</li> <li>Duration: 5–20 min</li> <li>Risk of emergence delirium: Moderate to high, relative to other perioperative phases</li> <li>Commonly administered medications: Administration of midazolam and/or propofol, if a patient demonstrates emergence delirium</li> <li>Our medication strategy: Preemergence administration of dexmedetomidine and/or hydromorphone, as a preemptive treatment for a patient who is at elevated risk for emergence delirium. If a patient demonstrates emergence delirium, then administer propofol. For more information administer propofol. For more</li> </ul>
	<ul> <li>delirium. If a patient demonstrates emergence delirium, then administer propotol. For more information about these medications, see supplemental materials S5 and S6.</li> <li>Desired patient status: Transitioning from semiconscious or unconscious to conscious and cooperative</li> <li>Staff responsibilities: Maintain monitoring of vital signs; maintain/secure airway and IV; and assess/treat emergence delirium, as needed</li> </ul>

**Note**: Stage 1 of anesthesia is commonly referred to as the induction stage. Patients are sedated but remain conscious and perhaps even conversational. Breathing is slow and regular. This stage progresses from analgesia with no amnesia, to analgesia with amnesia, and finally ends with the loss of consciousness.<sup>18</sup>

Stage 2 of anesthesia is referred to as the delirium or excitement stage. This stage is distinguished by disinhibition, delirium, uncontrolled movements, loss of eyelash reflex, hypertension, and tachycardia.<sup>18</sup> Airway reflexes remain intact during this phase and are often hypersensitive to stimulation and thus airway manipulation should be avoided. There is a higher risk of laryngospasm (involuntary closure of vocal cords) at this stage, which may be precipitated by any airway manipulation or another stimulus such as loud noises and conversation in the OR.<sup>19</sup> This stage is often where airway compromise occurs. Fast-acting agents help reduce the time spent in stage 2 as much as possible and facilitate entry to stage 3.

Stage 3 is a surgical plane of general anesthesia. Eye movements cease and respiratory depression occurs to various degrees.<sup>18</sup> Airway manipulation is safe at this stage.

Stage 4 is rarely seen and is an overdose state of anesthesia. This stage occurs when too much anesthetic agent is given relative to the amount of surgical stimulation, which may result in central respiratory paralysis, cardiac failure, and then death.<sup>18</sup>

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### Supplemental Material S5 Medication Strategy for Patients at Elevated Risk for Emergence Delirium

How material is used: Clinical reference

**Purpose**: This medication strategy (individualized anesthesia care plan) was designed to improve the prevention and treatment of emergence delirium

Phases		Types of Anesthesia and	Medications
Preoperation Preparation	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia
Immediately prior	1	Acetaminophen: 1000–1300 mg, oral	
operating room	2	<i>Optional</i> Dexmedetomidine: 8–20 mcg (0.25–effect), IV	-0.5 mcg/kg ideal body weight, titration to
	3	<ul> <li>Optional, one of the following:</li> <li>Fentanyl: 50–100 mcg, IV</li> <li>Hydromorphone: 0.2–1.0 mg, IV (Note: no anesthesia)</li> <li>Oral Opioid</li> </ul>	t administered as a premedication with regional
	Note: With dez administration.	xmedetomidine, the clinically effective onset of	sedation occurs 5–10 minutes after bolus
Induction of Anesthesia	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia
Administration of	1	Lidocaine: 5	50–100 mg, IV
an agent to provide sedation or general anesthesia as necessary	2	<ul> <li>Propofol, <i>one or both</i> of the following:</li> <li>Bolus: Incremental doses, 25–100 mg, IV</li> <li>Infusion: 25–150 mcg/kg/min, IV</li> </ul>	<ul> <li>Propofol, <i>both</i> of the following:</li> <li>GETA: 1.0–2.0 mg/kg, IV</li> <li>Infusion: 50–150 mcg/kg/min, IV</li> </ul>
	3	* <b>Ketamine</b> : <0.5 mg/kg (low dose	ketamine, maximum of 50 mg), IV
	4	*Hydromorphor	<b>ne</b> : 0.2–1.0 mg, IV
	<b>Note:</b> *Durin analgesic.	g most procedures we (VAPHS) recommend use	e of Ketamine and/or Hydromorphone as an
Maintenance of Anesthesia	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia
Maintain desired	1	Propofol infusion: 2:	5–150 mcg/kg/min, IV
required for the procedure	2	<ul> <li>Dexmedetomidine:</li> <li>Optional loading dose: 0.5–1.0 mcg/kg, IV</li> <li>Infusion: 0.2–1.0 mcg/kg/hour, IV</li> </ul>	, over 10 minutes
	3	Not applicable	<i>Optional</i> Remifentanil infusion: 0.04-0.1mcg/kg/min, IV
	4	*Ketamine: <0.5 mg/kg (low dose	e ketamine, maximum of 50 mg), IV
	5	*Hydromorphor	ne: 0.2–2.0 mg, IV
	awake fiberop use of ketamin	tic intubation) as an IV/monitored sedation. *Dene and/or hydromorphone as an analgesic.	uring most procedures we (VAPHS) recommend

Emergence-	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia
from anesthesia	of wieus	Preemergence, <i>bolus</i> or <i>infusion</i> of deymede	tomidine:
leading to the		<ul> <li>Bolus: 8–20 mcg (0.25–0.5 mcg/kg ideal bolt</li> </ul>	dy weight, titration to effect). IV, repeat as
greatest potential	1	needed	
for emergence		• Infusion: 0.2–1.0 mcg/kg/hour, IV (loading	g dose: 0.5–1.0 mcg/kg, IV, over 10 minutes)
delirium	2	Preemergence, hydromorphone bolus: 0.2-1.	0mg, IV, repeat as needed
	3	Preemergence, if not previously administere	d, acetaminophen infusion: 1000 mg, IV,
		Post-emergence, if patient demonstrates low	-to-moderate-risk behavior, <i>bolus</i> or
		<i>infusion</i> of dexmedetomidine:	
	4	• Bolus: 8–20 mcg (0.25–0.5 mcg/kg ideal bo	dy weight, titration to effect), IV, repeat as
		needed	
		• Infusion: 0.2–1.0 mcg/kg/hour, IV (loading	dose: 0.5–1.0 mcg/kg, IV, over 10 minutes)
		Post-emergence, if patient demonstrates mod	derate-to-high-risk behavior, <i>bolus</i> or
	5	Bolus: 50–200 mg IV repeat as needed	
		<ul> <li>Infusion: 50–150 mg/kg/min, IV</li> </ul>	
	Note: We (V.	APHS) do not recommend administering ketamin	he pre-emergence (late in the case), as this may
	increase the l	ikelihood of emergence delirium.	
Warnings,	General infor	mation	
Precautions, and	1 Condmini	stration of anosthatic agants and/or adjuncts can	anhanaa nhamaaa dumamiasu tharafara agutian
Strategies	n. Coadmin	ken to account for the additive impact on the nat	ient's vital signs <sup>20</sup>
	2. Note that	this medication strategy is only designed to prev	ent and/or treat emergence delirium.
	Dexmedetomi	idine	U
			1 1 1 2 20
	1. Clinically 2 Not all pa	significant risks of using dexmedetomidine: sev	who have a history of congestive heart failure
	advanced	heart blocks, bradvarrhythmias, and/or severe ve	entricular dysfunction should not receive
	dexmedet	omidine. <sup>20</sup> Alternatively, a reduced administratio	n plan for dexmedetomidine should also be
	anticipate	d for patients with one or more chronic and/or ag	e-related medical conditions. <sup>20</sup> For example,
	elderly pa	tients, diabetics, and those with expected or prov	en renal and/or hepatic impairment may not be
	ideal dexn	nedetomidine candidates.	
	3. Caution sl	hould be taken with patients who have a low rest	ing heart rate and/or low blood pressure (e.g.,
	young/nea	attny, athletes, soldiers, high vagal tone, and spin strategies should be applied to prevent clinically	al cord injury patients). <sup>2029</sup> with these patients,
	4 Prior to ac	iministration of dexmedetomidine staff should c	onsider the following two proactive strategies to
	offset anti	cipated systemic effects on vital signs:	
	a. Prehy	drate, increase IV fluid administration (e.g., crys	stalloid) to compensate for an anticipated drop in
	systol	lic blood pressure. <sup>24</sup>	
	b. The d	lrug may require preemptive glycopyrrolate IV to	elevate heart rate to a safe level.
	5. Post-admi	nistration of dexmedetomidine, staff should cons	sider the following four reactive strategies to
	offset syst	temic effects on vital signs:	
	a. Decre	ase IV fluid administration <sup>20</sup>	
	c. Eleva	te lower extremities. <sup>20</sup>	
	d. Use o	of pressor agents (e.g., atropine, glycopyrrolate, e	phedrine, and/or phenylephrine). <sup>20</sup>
	6. Clinically	effective onset of sedation occurs 5-10 minutes	after IV bolus administration, <sup>25</sup> 10–15 minutes
	after IV in	nfusion, <sup>20</sup> and the duration of action is approxima	tely 2 hours.

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### **Supplemental Material S6** Literature Review in Support of the Medication Strategy

### How material is used: Clinical reference

**Purpose**: This information represents a literature review of anesthetics and adjunct agents that are associated with an increased or decreased likelihood of emergence delirium. This information was used to shape the medication strategy (individualized anesthesia care plan) described in supplemental materials **S4** and **S5**.

**Midazolam (Benzodiazepine)**. Midazolam, an anxiolytic, amnestic benzodiazepine,<sup>25</sup> is perhaps the most universally administered sedative.<sup>26,27</sup> The clinical effects of midazolam occur as a result of agonist action on the binding sites on GABA<sub>A</sub> neurotransmitter receptors throughout the central nervous system.<sup>28</sup> Midazolam is often administered preoperatively to treat anxiety and is widely used due to its rapid onset of 2–5 minutes, 20–30 minute duration, and ability to be reversed.<sup>25,29</sup>

Despite widespread use of midazolam/benzodiazepines, several studies reported an association between administration of the medication and an increased likelihood of emergence delirium.<sup>3,30-32</sup> For example, one study found that preoperative use of benzodiazepines with patients >15 years of age was associated with nearly twice the risk of emergence delirium.<sup>3</sup> Similarly, in a prospective observational study of 1,868 adult patients, they reported that patients who were administered 0.1 mg/kg of midazolam as a premedication had a significantly greater incidence of emergence delirium than patients who did not receive midazolam.<sup>30</sup> Previous literature also suggests that patients with a history of post-traumatic stress disorder (PTSD) or trauma and who are administered midazolam are at a higher risk for emergence delirium.<sup>6,33-35</sup> There is some debate among providers about the relation between midazolam/benzodiazepines and emergence delirium;<sup>13</sup> nevertheless, in our opinion, the previous studies and other literature collectively indicate that anesthesia providers may reduce the likelihood of emergence delirium by avoiding administration of midazolam/benzodiazepines.

**Dexmedetomidine**. Dexmedetomidine, an alternative to midazolam,<sup>25,36-41</sup> is a sedative-analgesic (opioid-sparing) and a potent anxiolytic.<sup>6,20,33,42-45</sup> Dexmedetomidine is an agonist of the alpha-2 adrenergic receptor and acts by reducing the sympathetic noradrenergic outflow and inhibits the central release of norepinephrine, thereby attenuates the hyperarousal state and fight-or-flight response.<sup>33,43-49</sup> Dexmedetomidine, when administered prior to emergence (preoperatively, intraoperatively, and/or at the end of surgery), will facilitate cooperative and tranquil patient behavior.<sup>33,36,38-41,43-47,49-56</sup> Dexmedetomidine has an onset of 5–10 minutes and duration of 2–3 hours.<sup>20,25</sup>

Numerous studies have shown that dexmedetomidine is effective in preventing emergence delirium,<sup>31,38-41,46,48-57</sup> while few studies indicate that it may be used to treat an episode of emergence delirium.<sup>35,58,59</sup> One study showed that preoperative and intraoperative administration of dexmedetomidine significantly reduced the incidence and severity of emergence delirium among anxiety afflicted combat veterans.<sup>46</sup> In a randomized study of children undergoing surgery, administering 0.3 mcg/kg of dexmedetomidine 10 minutes after induction was associated with 27% less incidence of emergence delirium than the control group.<sup>60</sup> Overall, administration of dexmedetomidine is associated with a more cooperative patient and safer behavior throughout emergence.

**Ketamine**. Ketamine is a dissociative sedative-analgesic with opioid-sparing properties.<sup>25,61,62</sup> Ketamine is an antagonist of the N-methyl-D-aspartate (NDMA) receptor and modulates the presynaptic release of glutamate, which is the most abundant excitatory neurotransmitter in the central nervous system.<sup>29,63,64</sup> Inhibition of and reduction in available glutamate influences a decrease in the pathophysiologic response to stress, which is associated with a prevention of emergence delirium.<sup>29,34,64</sup> Timing of ketamine administration has varied widely across previous studies and included the following phases: preoperative, induction, and/or at the end of surgery (pre-emergence).<sup>32,33,37,50,64-78</sup> Ketamine has an onset of less than 1 minute and has a dose-dependent duration of 5–20 minutes.<sup>25,29,79</sup>

Previous research on ketamine has reported conflicting findings,<sup>66</sup> with some indicating that it increases the likelihood of emergence delirium<sup>67,68,77</sup> and others suggesting that it can reduce the likelihood.<sup>31,32,37,50,69,73-75,78,80,81</sup> Among the studies reporting an association between ketamine and an increased likelihood of emergence delirium, they administered ketamine under both of the following conditions: 1) administered a relatively higher dose (e.g., >0.5 mg/kg);<sup>67,77</sup> and 2) without an additional active anesthetic agent (e.g., propofol, volatile agent) or an active sedative adjunct (e.g., dexmedetomidine).<sup>67,77</sup> In contrast, in the studies that reported an association between use of ketamine and a decreased likelihood of emergence delirium, they administered ketamine under one or more of the following conditions: 1) administered a relatively low dose (e.g., <0.5 mg/kg);<sup>74,75,78,81</sup> and/or 2) paired with an active anesthetic agent (e.g., propofol, volatile agent) and/or an active sedative adjunct (e.g., dexmedetomidine).<sup>37,69,73-75,78,80,81</sup> For example, a double-blind study of pediatric cases with sevoflurane-based anesthesia found that patients who were administered 0.15 mg/kg of ketamine and 0.3 mcg/kg of dexmedetomidine, 10 minutes before the end of surgery, resulted in a significantly lower rate and severity of emergence delirium, relative to administration of saline (control condition).<sup>81</sup> Our review of literature indicates that when ketamine is administered as a low dose and/or when paired with an active anesthetic agent or sedative adjunct, there is a lower likelihood and severity of emergence delirium.

**Volatile Anesthetics (Inhalational)**. Volatile anesthetics (e.g., desflurane, halothane, isoflurane, and sevoflurane) are inhaled and produce titration-dependent amnesia, unconsciousness, and immobility, but not analgesia.<sup>28,29,82</sup> Volatile anesthetics are used for the rapid induction and maintenance of general anesthesia. This type of drug acts on the GABA<sub>A</sub> receptor by suppressing the excitatory channels and augmenting the inhibitory channels.<sup>28,29,82-84</sup> Volatile anesthetics, depending on the type and concentration, have an uptake and onset of 1–4 minutes and the patient will emerge (eye opening) 10–15 minutes following discontinuation.<sup>29,82</sup>

Much of the previous research reported that the administration of volatile anesthetics is associated with a higher rate and/or severity of emergence delirium.<sup>3,50,85-96</sup> For example, a randomized clinical trial of 80 adults undergoing invasive nasal surgery compared the rates of emergence delirium between a patient group with volatile anesthetic induction/maintenance and another group with propofol/total intravenous anesthesia (TIVA).<sup>92</sup> They used two different agitation assessment scales (Richmond Agitation-Sedation Scale and Riker Sedation-Agitation Scale) and, across both scales, they found 20%–25% rates of agitation in the volatile anesthetic group and a rate of 2.5% in the propofol/TIVA group. Also, several studies of the effects of volatile anesthetics found that patients who had noninvasive procedures (e.g., MRI) and were without pain also experienced a high rate of emergence delirium.<sup>86,97,98</sup> As a whole, a large portion of the previous literature indicates that volatile anesthetics are associated with an increased risk and severity of emergence delirium.

**Propofol/Total Intravenous Anesthesia (TIVA).** Propofol is a sedative-hypnotic that is administered for the rapid induction and maintenance of anesthesia,<sup>25</sup> and can also be administered to rapidly treat an episode of emergence delirium.<sup>99,100</sup> Propofol, like volatile anesthetics, augments and extends the duration of GABA<sub>A</sub>-mediated synaptic inhibition.<sup>83,84</sup> Propofol has an onset of less than 1 minute and a dose-dependent duration of 3–10 minutes.<sup>25,28</sup>

Several studies have shown that the administration of propofol, as a maintenance anesthetic and/or pre-emergence bolus, was associated with a lower rate and/or severity of emergence delirium.<sup>3,31,50,85-92,101-105</sup> For example, in two studies, each with >790 adult patients, the authors found that administration of propofol (i.e., TIVA) as a maintenance anesthetic was associated with a significantly reduced incidence of emergence delirium, when compared with patients who were administered a volatile anesthetic.<sup>89,90</sup> Unfortunately, few previous studies have systematically evaluated the use of propofol as an acute treatment during an episode of emergence delirium; nevertheless, providers have reported that propofol is an effective treatment for an episode of emergence delirium.<sup>99,100</sup>

**Summary of Relation Between Medications and Emergence Delirium**. Previous literature indicates that use of midazolam and/or volatile anesthetics are associated with an increase of the likelihood and severity of emergence delirium when compared with administration of alternative medications. As a result, we recommend and use a medication strategy that includes dexmedetomidine, ketamine, and propofol (TIVA), in an effort to reduce the likelihood and severity of emergence delirium among our high-risk patients.

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### Supplemental Material S7 General Information for the Lecture and Hands-On Training

### How material is used: Training

**Purpose:** Use during development and rollout of the training program. The document identifies the learning objectives, participants, time requirements, when and where the training should occur, and required material/equipment.

### **Learning Objectives**

### Lecture

- 1. Restate risk factors that are associated with emergence delirium.
- 2. Identify team strategies to prevent, treat, and de-escalate episodes of emergence delirium.
- 3. Describe a medication strategy related to emergence delirium.
- 4. Identify medically appropriate manual restraints for patients exhibiting dangerous behavior related to emergence delirium.

### Hands-On Training

- 1. Employ strategies to prevent, treat, and de-escalate dangerous and disruptive behavior associated with emergence delirium.
- 2. Utilize a team response to the patient actor exhibiting emergence delirium.
- 3. Employ safe and medically appropriate manual restraints with the patient actor exhibiting emergence delirium.

### Who is Involved in the Training

- **Instructors**: They should collectively have expertise in perioperative clinical management of a patient and hands-on restraint of patients. The instructor teaching restraint should have formal training in how to safely conduct hands-on restraint.
- Learners: Staff who work in a perioperative environment, particularly staff who administer anesthetics and adjunct agents, staff who work in a post-anesthesia care unit, and other staff who could be called to assist with patient restraint.
- **Patient actor**: During simulation, this role will be performed by a person who has been thoroughly trained to resist in a way that does not cause harm to the learners or themselves. Ideally, the role will be performed by a standardized patient who is a medically trained educator or an instructor. In the event that no patient actors are available, then the instructor may consider using a mannequin in place of the patient actor.

### When Should the Training Occur

- This training should be administered during rollout of the intervention and staff orientation, and as a refresher training as needed.
- Depending on the number of learners, we recommend that the instructor spend 30–60 minutes on the lecture and 30–60 minutes on the hands-on training (total of 60–120 minutes for both trainings).

### Location and Required Materials/Equipment

### Lecture

- The lecture should be given in a private room.
- Ideally, the instructor will use a projector or screen to show the PowerPoint file.
- All lecture materials are included in the packet of supplemental material.

### Hands-On

- The training should be performed in a private room or a room specifically designed for operating room simulation.
- The simulation room should have at minimum a stretcher, gastrointestinal bed, or table. Also, we recommend using simulated intravenous fluids and oxygen.
- All other materials will come from this packet.

### Supplemental Material S8 Agenda for the Lecture and Hands-On Training

How material is used: Training

Purpose: Provide the instructor with a concise list of training topics and corresponding durations

Recommended Time (Total 60–120 min)		Type of Training and Topics	Instructor(s)
2–4 min		Introduction	[enter name]
5–10 min		<b>PAASO Form:</b> Preoperative Anesthesia Assessment, Medication Strategy, and Outcome	[enter name]
3–6 min	nre	Communication Among Staff	[enter name]
2–4 min	ect	Adjust Environment	[enter name]
5–10 min	Ĺ	Medication Strategy	[enter name]
5–10 min		Patient Orientation Upon Emergence	[enter name]
5–10 min		<b>Response to Emergence Delirium</b>	[enter name]
3–6 min		Post-Emergence Delirium	[enter name]
		Transition	
5–10 min	_	Prebrief	[enter name]
10–18 min	n)	Guided Practice	[enter name]
2–4 min	on acti	Pre-Scenario 1 Talking Points	[enter name]
3–5 min	ds e Pr: ula	Simulation Scenario 1	[enter name]
3–5 min	andled	Post-Scenario 1 Talking Points	[enter name]
1–2 min	H uic S pi	Pre-Scenario 2 Talking Points	[enter name]
3–6 min	an an	Simulation Scenario 2	[enter name]
3–5 min		Post-Scenario 2 Talking Points	[enter name]
		Transition	
0–5 min		Highlights From Training	[enter name]
Notes The time on and simula		1.4 1.1 1.1	4

**Note**: The times are simply a recommendation and should vary based on learner needs and time constraints at the healthcare facility

### **Supplemental Material S9**

Perioperative Intervention To Prevent and Treat Emergence Delirium (PowerPoint, 19 slides plus talking points)

### How material is used: Training

**Purpose**: Guide staff's skill development and facilitate rollout and continuity of the intervention. This content was designed to cover all components of the intervention.

For a copy of the **PowerPoint file with talking points (S9)**, go to the webpage for this article (Vol. 4, No. 4 at PatientSafetyJ.com).



# Learning Objectives

At the conclusion of this presentation, participants will be able to:

- Restate risk factors that are associated with emergence delirium
- Identify team strategies to prevent, treat, and de-escalate episodes of emergence delirium
- Describe a medication strategy related to emergence delirium
- Identify medically appropriate manual restraints for patients exhibiting dangerous behavior related to emergence delirium



<b>Emergence</b> De	lirium
---------------------	--------

During the condition, the patient may engage in a wide range of behavior:

- Violence (e.g., punching, kicking, hitting, biting)
- Thrashing movement
- Aggression
- Combativeness
- Screaming

- Disconnect with current time and place (e.g., flashbacks, hallucinating),
- Attempting to exit bed/table
- Attempting to remove medical apparatuses

The condition begins with awakening from the anesthetic and/or adjunct agent and is followed by a return to baseline behavior after a short time period (typically less than 30 minutes).

3

### **Training Components** Lecture Hands-On Learning Team response and manual restraint • PAASO Form: Preoperative Anesthesia Assessment, Medication Strategy, and Guided practice: instructor-modeled Outcome manual restraint, learner rehearsal, and feedback • Communication among staff • Simulation: learners, as a team, Adjust environment physically interact to mimic a real-life Medication strategy response to emergence delirium Patient orientation upon emergence Response to emergence delirium Post-emergence delirium 4



Part 2: Medication Strategy	Staff who complete part 2: Anesthesia provider Purpose: Facilitate hand-offs and monitor the perioperative intervention across department providers			
2.1. Medications administered and/or maintenance of anesthes	luring pre-operation, induction, ia phases	□ Dexmed □ Fentany □ Hydron □ Ketamin	detomidine yl norphone ne	□ Propofol □ Remifentanil □ Midazolam □ Volatile agent
2.2. Preemergence 2.3. If applicable, in response to	emergence delirium	Acetam     NONE     Dexmed     NONE     Decrmed	inophen detomidine	Oral opiates     Hydromorphone     Propofol     Propofol     Midazalam
Part 3: Outcome	Staff who complete part 3: Ane Purpose: Monitor effectiveness facilitate collecting information	sthesia pro of the inter for patient	vider or PACU vention acros record	I nurse is department providers and
<b>3.1. Was there an episode of em</b> thrashing, aggression, combativ attempting to exit bed/table, and medical apparatuses	ergence delirium? E.g., violence, eness, screaming, hallucinating, /or attempting to remove	□No	□Yes	
If yes emergence delirium (3.1) 3.2A. Was the behavior dang	: erous?	□No	□ <b>Yes</b> , desc	ribe:
3.2B. Was anyone injured?		□No	□ <b>Yes</b> , desc:	ribe:
3.2C. What may have trigger bright light, or tactile stimulu	<b>ed the episode?</b> E.g., loud sound, s	Describe:		

## **Communication Among Staff**

- Implement plan across perioperative phases to help staff recognize the high-risk patient (e.g., patient wears a unique colored cap or wristband)
- Transitions of care (e.g., ensure communication about patient's risk status of emergence delirium)
- During time-out discuss patient's risk status for emergence delirium, plan to treat, and staff roles
- Use overhead paging system to announce "Anesthesia stat" when staff support is required. Announce location and room number.





Preoperation	Somoneo	Intravanous (IV)/Manitored Sodation	Coneral Anasthasia With or Without			
Preparation	of Meds	With or Without Regional Anesthesia	Regional Anesthesia			
Immediately	1	Acetaminophen: 1000–1300 mg, oral				
to operating room	2	<i>Optional</i> Dexmedetomidine: 8–20 mcg (0.25–0.5 mcg/kg ideal body weight, titration to effect), 1V				
	3	<ul> <li>Fentanyl: 50–100 mcg, IV</li> <li>Hydromorphone: 0.2–1.0 mg, IV (Note:</li> </ul>	not administered as a premedication with			
	5	• Hydromorphone. 0.2-1.0 mg, 1v (Note.	not administered as a premedication with			
		regional anesthesia) • Oral Opioid				
	Note: With de bolus admini	regional anesthesia) • Oral Opioid exmedetomidine, the clinically effective on istration.	set of sedation occurs 5–10 minutes after			
Induction of	Note: With de bolus admini Sequence	regional anesthesia) • Oral Opioid exmedetomidine, the clinically effective on istration. Intravenous (IV)/Monitored Sedation	set of sedation occurs 5–10 minutes after General Anesthesia With or Without			
Induction of Anesthesia	Note: With de bolus admini Sequence of Meds	regional anesthesia) • Oral Opioid exmedetomidine, the clinically effective on istration. Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia			
Induction of Anesthesia Administration	Note: With de bolus admini Sequence of Meds 1	regional anesthesia) • Oral Opioid exmedetomidine, the clinically effective on istration. Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia Lidocaine:	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV			
Induction of Anesthesia Administration of an agent to	Note: With debolus admini	regional anesthesia) • Oral Opioid exmedetomidine, the clinically effective on istration. Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia Lidocaine: Propofol, one or both of the following:	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV Propofol, both of the following:			
Induction of Anesthesia Administration of an agent to provide sedation or concern	Note: With do bolus admini Sequence of Meds 1	<ul> <li>• Oral Opioid</li> <li>• Oral Opioid</li> <li>exmedetomidine, the clinically effective on istration.</li> <li>Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia</li> <li>Lidocaine:</li> <li>Propofol, one or both of the following:</li> <li>• Bolus: Incremental doses, 25–100 mg,</li> </ul>	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV Propofol, both of the following: • GETA: 1.0–2.0 mg/kg, IV			
Induction of Anesthesia Administration of an agent to provide sedation or general anethesia as	Note: With de bolus admini Sequence of Meds 1 2	<ul> <li>Oral Opioid</li> <li>Oral Opioid</li> <li>exmedetomidine, the clinically effective on istration.</li> <li>Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia</li> <li>Lidocaine:</li> <li>Propofol, one or both of the following:</li> <li>Bolus: Incremental doses, 25–100 mg, IV</li> </ul>	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV Propofol, both of the following: • GETA: 1.0–2.0 mg/kg, IV • Infusion: 50–150 mcg/kg/min, IV			
Induction of Anesthesia Administration of an agent to provide sedation or general anesthesia as necessary	Note: With de bolus admini Sequence of Meds 1 2	<ul> <li>Oral Opioid</li> <li>Oral Opioid</li> <li>exmedetomidine, the clinically effective on istration.</li> <li>Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia Lidocaine:</li> <li>Propofol, one or both of the following:</li> <li>Bolus: Incremental doses, 25–100 mg, IV</li> <li>Infusion: 25–150 mcg/kg/min, IV</li> </ul>	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV Propofol, both of the following: • GETA: 1.0–2.0 mg/kg, IV • Infusion: 50–150 mcg/kg/min, IV			
Induction of Anesthesia Administration of an agent to provide sedation or general anesthesia as necessary	Note: With do bolus admini	<ul> <li>Oral Opioid</li> <li>Oral Opioid</li> <li>exmedetomidine, the clinically effective on istration.</li> <li>Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia Lidocaine:</li> <li>Propofol, one or both of the following:</li> <li>Bolus: Incremental doses, 25–100 mg, IV</li> <li>Infusion: 25–150 mcg/kg/min, IV</li> <li>*Ketamine: &lt;0.5 mg/kg (low dose</li> </ul>	set of sedation occurs 5–10 minutes after General Anesthesia With or Without Regional Anesthesia 50–100 mg, IV Propofol, both of the following: • GETA: 1.0–2.0 mg/kg, IV • Infusion: 50–150 mcg/kg/min, IV ketamine, maximum of 50 mg), IV			

			-0,			
Phases		Types of Anesthesia and	Medications			
Maintenance of Anesthesia	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia			
Maintain desired	1	Propofol infusion: 25–150 mcg/kg/min, IV				
level of anesthesia required for the	2	Dexmedetomidine: 2 • Optional loading dose: 0.5–1.0 mcg/kg, IV, over 10 minutes • Infusion: 0.2–1.0 mcg/kg/hour, IV				
procedure	3	Not applicable	Optional Remifentanil infusion: 0.04-0.1mcg/kg/min, IV			
	4	*Ketamine: <0.5 mg/kg (low dose ketamine, maximum of 50 mg), IV				
	5	5 <b>*Hydromorphone:</b> 0.2–2.0 mg, IV				
	<b>Note:</b> A dexmedetomidine infusion alone can be used with a limited number of specific procedures (e.g., awake fiberoptic intubation) as an IV/monitored sedation. *During most procedures we (VAPHS) recommend use of ketamine and/or hydromorphone as an analgesic.					

Phases		Types of Anesthesia and	Cegy Medications
<b>Emergence</b> - Patient awakens	Sequence of Meds	Intravenous (IV)/Monitored Sedation With or Without Regional Anesthesia	General Anesthesia With or Without Regional Anesthesia
from anesthesia leading to the greatest potential for emergence dutinium	1	<ul> <li>Preemergence, <i>bolus</i> or <i>infusion</i> of dexm</li> <li>Bolus: 8-20 mcg (0.25-0.5 mcg/kg ideal as needed</li> <li>Infusion: 0.2-1.0 mcg/kg/hour, IV (load minutes)</li> </ul>	edetomidine: body weight, titration to effect), IV, repeat ling dose: 0.5–1.0 mcg/kg, IV, over 10
demrum	2	Preemergence, hydromorphone bolus: 0 Preemergence, if not previously adminis	.2-1.0mg, IV, repeat as needed tered, acetaminophen infusion: 1000 mg,
	4	<ul> <li>Post-emergence, if patient demonstrates infusion of dexmedetomidine:</li> <li>Bolus: 8-20 mcg (0.25-0.5 mcg/kg ideal as needed</li> <li>Infusion: 0.2-1.0 mcg/kg/hour, IV (load minutes)</li> </ul>	<b>low-to-moderate-risk behavior</b> , <i>bolus</i> or body weight, titration to effect), IV, repeat <b>ting dose:</b> 0.5–1.0 mcg/kg, IV, over 10
	5	Post-emergence, if patient demonstrates infusion of propofol: • Bolus: 50-200 mg, IV, repeat as needed • Infusion: 50-150 mg/kg/min, IV	moderate-to-high-risk behavior, <i>bolus</i> or

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# Patient Orientation Upon Emergence Mention familiar people and places (e.g., patient's name and nickname, names of immediate family members, pet's name, location) Use patient's preferred physical interactions (e.g., tap on right shoulder and do not touch legs) Use patient's gender preference for the nursing care team Present comforting visual cues (e.g., picture projected onto ceiling of the U.S. flag)







**Note:** This video is an example of staff who were <u>learning</u> manual restraint in a simulation with a patient actor





# **Risks with Manual Restraint**

### Positional asphyxiation

· Weight should never be applied to the patient's thoracic cavity

Aspiration

• Risk should be low due to patient's NPO status

**Orthopedic injuries** 

- Do not exceed patient's range of motion
- Do not use joint lock or pain compliance techniques

### Skin integrity

• Apply a measured amount of force during restraint and do not default to full force



# Image: Contract information Image: Contract information</td

### Supplemental Material S10 Outline of Phases, Roles, and Materials for the Hands-On Training

### How material is used: Training

Purpose: Provide the instructor with information about phases, roles, and materials used during the hands-on training

Dh	2000			Supplemental	
rı	lases	Patient Actor	Learner	Instructor	Material
Pre-Brief (5–10 min)		Instructor reads the pre- learning environment, c time-out, and psycholog	<b>∢</b> S11		
Guided Pract (10–20 min)	tice	Guided practice consist rehearsal, and immedia Guided practice involvi gradual fading of mode proficiency.	restraint, learner behavior skills training). -step instruction with her demonstrated	<b>∢</b> \$12	
N	Note: The follow	ving phases, roles, and m	aterials apply to both simula	tion <u>scenario 1</u> and <u>scenar</u>	<u>io 2</u>
Simulation P Talking Point (2–4 min)	re-Scenario ts	Instructor reads pre-sce includes a reminder of s answering questions.	<b>∢</b> S13		
	Starting Behavior	Begin lying on a bed or stretcher in the supine or side position	Stand to the side or at the foot of the bed	Present with learners and prepared to closely observe their behavior	<b>⊲</b> \$14, \$15
Simulation Scenario (3–5 min)	Intervening Behavior	Emerges from anesthesia by yelling, sitting up in the bed, pulling at their arm or face to simulate removal of IV or oxygen	<ul> <li>Call for assistance (e.g., "Anesthesia stat")</li> <li>Attempt to verbally orient the patient actor</li> <li>Apply manual restraint techniques to the patient actor (identical to training)</li> <li>Protect IV and airway, if necessary</li> <li>Administer medication to sedate the patient actor</li> <li>One team member acts as an observer to monitor patient's safety</li> </ul>	Continuously moves to a position within the room to observe all learners, monitoring safety and learner performance against the simulation checklist	-
	End Behavior	Once the patient actor determines that the learners have applied restraint correctly, then they will stop resisting	Restraint applied to the patient	Verifies that learners performed tasks correctly and then stops the scenario	
Simulation Point Talking Point (3–5 min)	ost-Scenario ts	Provides feedback to learners about their restraint application and any safety concerns	Receives feedback on their performance and asks questions	Reviews simulation checklist and provides feedback to learners and facilitates discussion with the post-scenario talking points	<b>∢</b> \$13, \$15

### Supplemental Material S11 Pre-Brief for the Hands-On Training

### How material is used: Training

**Purpose**: Provide personnel with fundamental information about the components of the hands-on training. Conveying this information is essential to ensure that everyone understands their roles, behavioral expectations, and the context of the hands-on training.

Components	Description
Supportive Learning Environment and Confidentiality	Hands-on training is intended to be a supportive learning environment. This means that it is a safe space to make mistakes and discuss questions, difficulties, or errors. The learner will be an active participant in the hands-on training as well as an observer of peers. All actions, discussions, and feedback that occur in this session should not be shared with others outside of the training. Likewise, the training team will not speak about learner performance or direct comments to supervisors, co-workers, etc.
Fiction Contract	The simulation environment and patient actor have certain limitations in their ability to exactly mirror real life. The learner is expected to suspend judgement of realism when the simulation does not precisely match real life. Learners should engage with the same amount of care, respect, and attention that would be given in a real-world event of this nature.
Learning Objectives	<ol> <li>Employ strategies to prevent, treat, and de-escalate dangerous and disruptive behavior associated with emergence delirium.</li> <li>Utilize a team response to the patient actor exhibiting emergence delirium.</li> <li>Employ safe and medically appropriate manual restraints with the patient actor exhibiting emergence delirium.</li> </ol>
Environment	The simulated operating room has been set up to appear as close to real as can be accomplished. The room will include simulated operating room equipment such as the anesthesia gas machine (AGM), intravenous (IV) infusion pump, vital signs monitors, and wall-mounted and supplied oxygen and suction.
Roles	<b>Patient actor</b> : This role should be fulfilled by an instructor or medically trained educator (e.g., standardized patient) that will portray the patient. They are permitted to provide resistant force against the restraints and have been thoroughly trained to resist in a way which does not cause harm to the learner or to themselves. They have been trained to maintain their balance and remain aware of their limbs and head. The actors may push against the learners' restraint hold but will <u>not strike or attack the learners</u> in any way. The actors are permitted to resist, sit up, and attempt to remove simulated medical interventions (such as the IV). <b>Learners</b> : They are permitted to touch and manually restrain the patient actor. They are <u>not permitted</u> to strike, push, trip, lift, takedown to the ground, press against a wall, choke, or place any pressure on the thoracic cavity of the actor. They must not use any technique that they may have learned outside of this training. Any restraints used should follow the proper physical intervention as instructed in the emergence delirium training courses regarding arm control, standing escort, and supine or side manual restraints. <b>Instructors</b> : They may act as other responding team members but will not initiate or take the lead with any verbal intervention or restraints; they will only take cues from learners and provide support after all participating learners have attempted to apply the manual restraint. Instructors will always acknowledge before the start of a simulation if they are participating or not.
Time-Out	At any time, the instructors, learners, or patient actor may call a "time-out" and stop the hands-on training, in the event that something is unsafe. Instructors may also call a time-out to provide further instruction or guidance. Learners may also call a time-out if they become stuck and would like help formulating a new strategy.
Simulation Pre- and Post-Scenario Talking Points	Immediately before and after each of the two simulation scenarios, the instructor will address a few talking points. In the pre-scenario talking points, the instructor will communicate what will happen during the scenario and what they expect the learner to demonstrate. In the post-scenario talking points, the instructor will discuss what went well and what didn't go well.
Psychological Safety	The simulation scenarios may be a trigger or hit close to home for some of the learners. If that is the case, they are not obligated to participate and can feel free to step out or take a break at any time. If staff do not complete the training, they are instructed not to use manual restraint.

### Supplemental Material S12 Guided Practice for the Hands-On Training

### How material is used: Training

**Purpose**: Guided practice consists of instructor-modeled manual restraint (i.e., hands-on) followed by learner rehearsal with immediate instructor feedback (i.e., behavior skills training).

### **Overview of Guided Practice**

Guided practice consists of instructor-modeled manual restraint (i.e., hands on) followed by learner rehearsal with immediate instructor feedback (i.e., behavior skills training). This process will start slowly and include step-by-step instruction with gradual fading of modeling and prompts until the learners demonstrate proficiency. In the first round of practice, the patient actor (e.g., standardized patient or instructor) will offer light resistance against the restraint to help the learner safely measure the effectiveness of their hand placement and force. After the first round of restraint practice, the process will restart with the patient actor engaging in increased resistance against the restraint and using greater variability of movement. This process will repeat until the learners demonstrate proficiency (typically two to three rounds of practice are required).

### **Manual Restraint Description**

- 1. This section is not intended to be a "how-to" restraint application guide. Please consult your local restraint expert for additional information about how to safely apply manual restraint.
- 2. Arms Pressure will be applied to the shoulder area and below the elbow (forearm) of both arms. At no time should pressure be applied to the elbow joint. The arms will remain in line with the body and will not be manipulated in a way that produces pain to force compliance. The arms will not be maneuvered away from the body or exceed the patient's natural range of motion.
- 3. Legs Pressure will be applied to the thighs and shin areas. The legs can typically support pressure that is placed above or below the knees, but not directly on the knees.
- 4. Head It may be necessary to support the head of the patient in a way that will prevent the patient from removing or dislodging the oxygen apparatus. Pressure may be placed on the patient's forehead and under the chin, but at no time should weight be applied to the neck (i.e., do not use a choke hold).
- 5. Intravenous (IV) site The area around the IV site will need to be secured to prevent IV removal. This will be accomplished by cupping a hand over the IV site to prevent dislodgment.
- 6. General guidelines Do not apply weight to the thoracic cavity of the patient and do not lie across the patient's upper body, which may restrict the patient's respiration. Please note that learners may have difficulty gauging the difference between adequate and excessive force when restraining a patient; therefore, this topic should be emphasized during training.

### Supplemental Material S13 Simulation Pre- and Post-Scenario Talking Points for the Hands-On Training

### How material is used: Training

**Purpose**: The pre-scenario talking points will convey fundamental information to the learners and patient actors about the expected behavior. The post-scenario talking points will facilitate learning and evaluation.

### **Pre-Scenario Talking Points**

**Context for the scenario**: It will begin with the surgical team having just completed surgery and preparing for the patient to emerge from sedation and orient to the environment. As a reminder, the actor will engage in behavior associated with emergence delirium.

**Sequence of events**: The instructor will announce "Begin" to start the simulation. The simulation is expected to last approximately 3–5 minutes. Once the scenario begins, the learners are expected to assess and react to the situation as trained, except for making real calls for assistance. Instead, the learners may simulate a call by picking up a phone and announcing who they would call and what they would say.

We are about to begin the scenario. Please keep the following in mind during the scenario:

- Patient actor will be restrained in a [side or supine] position.
- Patient actor will not strike anyone.
- Call a time-out if you need help or see a safety concern.
- Learner should call for assistance (e.g., "Anesthesia stat").
- Learners should attempt to verbally orient the patient. For example, the learner will state the patient's name (e.g., Rebecca Smith), her spouse's name (e.g., John Smith) and where the procedure is being performed (e.g., Pittsburgh, Pennsylvania).
- Learners should only use restraint techniques taught in this course.
- If appropriate, a learner should protect the airway and IV.
- If appropriate, a learner should simulate re-sedation of the patient actor.
- A learner should act as an observer to monitor the patient actor's safety.

Any questions before we begin the scenario?

### **Post-Scenario Talking Points**

- Overall, how did the scenario go? Purpose of question: Will get the learners to begin talking about their experience.
- What was your role in the scenario? *Purpose of question*: Instructor will use the learners' responses to determine if all roles were performed as trained.
- What went well? *Purpose of question*: Instructor will use the learners' responses to evaluate whether they recognize what tasks were performed correctly. Instructor will also use this as an opportunity to praise their performance.
- What didn't go well? *Purpose of question*: Instructor will use the learners' responses to determine if they recognize what tasks were performed incorrectly and better understand what challenges they encountered. This is an opportunity for the patient actor to provide feedback to the learner. Depending on the learners' performance, the instructor may need to rerun the scenario.
- Other questions the instructor may want to pose
  - Did you perform your technique(s) as trained? If you varied from the technique, do you know why?
  - Did the team clearly communicate the safety needs of the patient?
  - What was most helpful about this scenario?
  - What would you do differently next time?
- Do you have any questions or concerns about the training or simulation?

### **Supplemental Material S14** Simulation Scenarios 1 and 2 for the Hands-On Training

### How material is used: Training

**Purpose:** Identifies learning objectives, context, setting information, and expected behavior during the scenario. This will be used by the patient actor and instructor during planning and implementation of the simulation training.

Components	Description
Learning objectives	<ol> <li>Learner will employ strategies to prevent, treat, and de-escalate dangerous and disruptive behavior associated with emergence delirium.</li> <li>Learners will utilize a team response to the patient actor exhibiting emergence delirium.</li> <li>Learners will employ safe and medically appropriate manual restraints with the patient actor exhibiting emergence delirium.</li> </ol>
Information for both scenarios	<ol> <li>Background on emergence delirium: The condition often begins with awakening or emergence from the anesthetic and/or adjunct agent and is followed by a return to baseline behavior after a short time period (typically less than 30 minutes). During the bout of emergence delirium, the patient may engage in a wide range of behavior that includes one or more of the following: violence, thrashing movement, aggression, combativeness, screaming, disconnect with current time and place, attempting to exit bed/table, and/or attempting to remove medical apparatuses.</li> <li>Patient actor behavior: While waking from anesthesia, the actor demonstrates emergence delirium by yelling; attempting to get up from the table/stretcher; attempting to remove an intravenous (IV) line or oxygen; and acting as if they are confused, agitated, and hallucinating (flashbacks of a traumatic experience). The actor should not attempt to strike the learner and must avoid causing harm.</li> <li>Learner behavior: The learners will use specific techniques to manually restrain the actor.</li> <li>Required equipment: At minimum a stretcher, gastrointestinal bed, or table. Also, we recommend using a simulated IV and oxygen.</li> <li>Location: A private room or a room specifically designed for operating room (OR) simulation</li> <li>Information for verbal orientation of patient: The learner will use names and locations to verbally orient the patient actor during their simulated bout of emergence delirium. For example, the learner will state the patient's name (e.g., Rebecca Smith), her spouse's name (e.g., John Smith), her dog's name (e.g., Jaffy), and where the procedure is being performed (e.g., Pittsburgh, Pennsylvania).</li> <li>For additional information to prepare the patient actor, see S10, "Outline of Phases, Roles, and Materials for the Hands-On Training."</li> </ol>
	<ol> <li>Patient actor had complaints of abdominal pain and is waking from an appendectomy procedure.</li> </ol>
Information for scenario 2	<ol> <li>Patient actor's position during simulation: The actor will start the scenario lying in a <i>side</i> position and will end by being restrained in a <i>side</i> position.</li> <li>Patient actor had complaints of gastrointestinal pain and is waking from a colonoscopy procedure.</li> </ol>

# Supplemental Material S15 Simulation Checklist for the Hands-On Training

How material is used: Training Purpose: Instructor uses it to facilitate staff learning and evaluation

Scenario 1: Manual Restraint of Supine Position	Yes	No	Comments
1. Did a team member call for assistance (e.g., "Anesthesia stat")?			
2. Did a team member attempt to verbally orient the patient by saying familiar			
names and locations?			
3. Did a team member manually restrain the patient's <b>right arm</b> correctly?			
4. Did a team member manually restrain the patient's left arm correctly?			
5. Did a team member manually restrain the patient's legs correctly?			
6. Did a team member manually restrain the patient's head correctly?			
7. If necessary, did the team protect the <b>airway and the IV</b> ?			
8. If necessary, did a team member administer medication to resedate the patient?			
9. Did a team member act as an observer to monitor patient's safety?			
Scenario 2: Manual Restraint of Side Position	Yes	No	Comments
1. Did a team member call for assistance (e.g., "Anesthesia stat")?			
2 Did a team member attempt to verbally orient the nation by saving familiar			
2. Did a team member attempt to verbany orient the patient by saying fammar			
names and locations?			
<ul><li>a. Did a team member attempt to versary orient the patient by saying familiar names and locations?</li><li>3. Did a team member manually restrain the patient's right arm correctly?</li></ul>			
<ul> <li>a. Did a team member manually restrain the patient's right arm correctly?</li> <li>4. Did a team member manually restrain the patient's left arm correctly?</li> </ul>			
<ul> <li>a. Did a team member antempt to versary orient the patient by saying familiar names and locations?</li> <li>3. Did a team member manually restrain the patient's right arm correctly?</li> <li>4. Did a team member manually restrain the patient's left arm correctly?</li> <li>5. Did a team member manually restrain the patient's legs correctly?</li> </ul>			
<ul> <li>a. Did a team member annually restrain the patient's right arm correctly?</li> <li>4. Did a team member manually restrain the patient's left arm correctly?</li> <li>5. Did a team member manually restrain the patient's legs correctly?</li> <li>6. Did a team member manually restrain the patient's head correctly?</li> </ul>			
<ul> <li>a. Did a team member attempt to versary orient the patient by saying familiar names and locations?</li> <li>3. Did a team member manually restrain the patient's right arm correctly?</li> <li>4. Did a team member manually restrain the patient's left arm correctly?</li> <li>5. Did a team member manually restrain the patient's legs correctly?</li> <li>6. Did a team member manually restrain the patient's head correctly?</li> <li>7. If necessary, did the team protect the airway and the IV?</li> </ul>			
<ul> <li>a. Did a team member annually restrain the patient's right arm correctly?</li> <li>3. Did a team member manually restrain the patient's left arm correctly?</li> <li>4. Did a team member manually restrain the patient's left arm correctly?</li> <li>5. Did a team member manually restrain the patient's legs correctly?</li> <li>6. Did a team member manually restrain the patient's head correctly?</li> <li>7. If necessary, did the team protect the airway and the IV?</li> <li>8. If necessary, did a team member administer medication to re-sedate the patient?</li> </ul>			