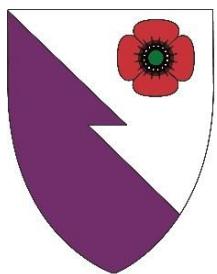


The Royal College of Emergency Medicine

Best Practice Guideline

**Pharmacological
Agents for Procedural
Sedation and Analgesia
in the Emergency
Department**



**October 2020
(Revised)**

Summary of recommendations

- 1.** Every Emergency Department should have a procedural sedation policy that is regularly reviewed, and audited
- 2.** The procedural sedation policy should provide guidelines for pharmacological agents that are routinely used in the Department
- 3.** A pro-forma should be used for procedural sedation and analgesia (PSA) as a checklist and as an auditable record of the procedure
- 4.** Adverse events should be reported using the SIVA reporting tool

Scope

This document provides guidelines for the use of common pharmacological agents in sedation in the Emergency Department in the UK. It provides guidelines for the use of these agents for adults and children (excluding neonates).

This guideline only applies to pharmacological agents used in sedation in the Emergency Department and is not intended to be used in any other setting. It does not set standards for safe sedation, these are set out in the "Safe Sedation of Adults in the Emergency Department" guideline developed by the Royal College of Anaesthetists and the College of Emergency Medicine in November 2012 (1) and NICE Clinical Guideline 112 "Sedation in children and young people" (2). This guideline is intended to be used as an adjunct to these guidelines.

It does not provide guidance for anaesthesia.

A [pro-forma template](#) has been developed for use as a checklist, for formal recording, for monitoring and for the purposes of audit (Appendix 1).

Reason for development

There are currently no recommended guidelines for the use of pharmacological agents in procedural sedation and anecdotally, practice varies widely. There are many sedation techniques available; this document provides guidance on common combinations of pharmacological agents and suggested drug doses.

Introduction

The practice of Emergency Medicine (EM) includes the performance of procedures that will cause pain and anxiety. The EM practitioner should always seek to minimise discomfort and apprehension with appropriate non-pharmacological and pharmacological interventions or techniques. Procedural sedation and analgesia (PSA) is an appropriate technique in some circumstances, using short-acting analgesics and sedative medicines to facilitate the effective performance of a procedure.

Current guidelines focus on procedural standards for delivering safe sedation in the Emergency Department. Some Emergency Departments have developed guidelines for PSA that include the use of pharmacological agents and suggested drug doses.

Comprehensive guidelines for PSA in both adults and children that include detailed review of common pharmacological agent's properties, usage and side effects are available through the point of care resource, UpToDate® (3), (4). The American College of Emergency Physicians published an official Clinical Policy on procedural sedation and analgesia in 2014 (5). These guidelines have informed this document.

Audit of procedural sedation is on the Royal College of Emergency Medicine's national clinical audit cycle, first audit August 2015 – January 2016.

Preparing for Procedural Sedation and Analgesia

Effective procedural sedation requires:

Analgesia: Pain experienced by the patient should be treated with analgesia rather than sedation. Pain should be assessed and managed prior to starting sedation using the WHO pain ladder principles and the need for ongoing pain relief post-procedure considered. Where opiate pain relief has been given prior to PSA, doses of sedatives should be adjusted accordingly.

Anxiolysis: Non-pharmacological methods of reducing anxiety are often the most effective and include consideration of the environment and patient comfort. Environment is particularly important for children and patients with dementia or learning difficulties. Family members often provide invaluable support and distraction. Most painful procedures are best performed with the patient supine.

Sedation: The Joint Commission on Accreditation of Healthcare Organizations in the United States has attempted to define the levels of sedation (6) (see Appendix 2). For most procedures in the ED, the level of required sedation will be moderate to deep, this should be determined in advance.

Amnesia: A degree of amnesia will minimise unpleasant memories associated with the procedure.

In most circumstances a combination of short acting analgesics and sedatives are required as the only pharmacological agent that has the potential to provide analgesia, sedation, anxiolysis and amnesia is ketamine.

Choice of Sedation Agents

The appropriate choice of pharmacological agents for PSA depends on:

- 1) The nature of the procedure
- 2) The planned level of sedation
- 3) Training and familiarity of the sedating practitioner with potential pharmacological agents
- 4) Patient factors
- 5) The local environment

Appendix 6 discusses this in greater detail.

Pharmacological Agents for PSA

Commonly used pharmacological agents are described in Appendix 3. There are many variations in the combinations of pharmacological agents described in the literature and many different ways of delivering the pharmacological agents. The most common agents are described along with the routes and doses for which there is most supportive evidence.

Having selected the appropriate drugs for the needs of the patient, doses of the pharmacological agents need to be tailored to the individual patient to deliver the required effects. Great care should be used when administering sedatives because of:

- Slow and variable onset time
- Inter-patient variability in dose requirement
- Synergistic action between drugs

Table 1 provides information for common pharmacological agents for PSA for adult patients and Table 2 provides equivalent information for paediatric patients.

There is a paucity of evidence of superiority of any one agent over another, and similarly there is no evidence that any agent has greater safety. There is some limited evidence of higher procedural success rates for some procedures with easily titratable sedative drugs.

During Sedation

Adherence to the guidance “Safe Sedation of Adults in the Emergency Department” guideline developed by the Royal College of Anaesthetists and the College of Emergency Medicine in November 2012 is advocated (1).

Post-Procedure Monitoring

Full monitoring post-procedure should continue in the same clinical area (with the same facilities available including staffing) until the patient meets the criteria for safe discharge:

- vital signs returned to normal levels
- the patient is awake with intact protective reflexes and no longer at risk of reduced level of consciousness
- nausea, vomiting and pain have been adequately addressed.

Post-Sedation Advice

All patients being discharged home should be discharged home in the care of a competent adult with verbal and written advice. Information and discharge advice for children who have had ketamine PSA is available on the College website at www.rcem.ac.uk/Shop-Floor/Clinical%20Guidelines/Local%20Guidelines/Default.asp?f=edit#Discharge. There is also generic post-discharge advice available at the same site.

Evidence

There is currently limited evidence on the safety of PSA in the UK both in adults and children (2). The American College of Emergency Physicians Clinical Policies Subcommittee on Procedural Sedation and Analgesia evaluated a number of clinical questions related to PSA including the safety of some common pharmacological agents and combinations of agents (6). The data from that clinical policy is included in this guideline (Appendix 4).

Adverse Event Reporting and Data Collection

A large prospective database of sedation cases including data on drugs, procedures, the depth of sedation and complications would help to clarify the safety of sedation and improve clinical practice. Recognising the wide variation in sedation practices and the importance of recording and tracking adverse events, The World Society for Intravenous Anaesthesia (World SIVA) International Sedation Task force has developed an event reporting tool for sedation-related adverse events. This tool is available as a web page through several organisations(e.g. <http://aesedationreporting.com/site/Register.aspx>), as a tick box form that could be incorporated into electronic patient record, or as a paper-based tool (8). This tool (Appendix 5) would complement the information collected on the PSA Checklist and Monitoring proforma (Appendix 2).

Table 1: Adult: Pharmacological Agents for Procedural Sedation and Analgesia

Click [here](#) to download Table 1 and Table 2

Agent (references)	Role	Route	Initial dose - Adult	Repeat dose	Initial dose - Adult	Repeat dose	Initial onset time (min)	Peak effect time (min)
			Elderly	Elderly	Adult	Adult		
Propofol (4)	Sedation/Amnesia	IV	10 - 20 mg (given slowly)	10 - 20 mg (given slowly)	0.5– 1.0 mg/kg	0.5mg/kg every 3-5mins	½ - 1	1 - 2
Midazolam (4)	Sedation/Amnesia	IV (over 1-2mins)	0.5mg	0.5mg	1 - 2 mg (max single dose 2.5mg)	After 2-5mins	1 - 2	3 - 4
Ketamine (9)	Sedation/Amnesia/Analgesia	IV (give over 30-60secs)	10 - 30 mg		1 mg/kg	0.25-0.5mg/kg every 5-10 mins	½ - 1	1 - 2
Ketamine (9)	Sedation/Amnesia/Analgesia	IM			4-5 mg/kg	2 – 2.5 mg/kg every 5-10 mins	½ - 1	1 - 2
Ketamine (4, 9)	Analgesia (sub-dissociative)	IV			0.3mg/kg		½ - 1	1 - 2
Fentanyl (4)	Analgesia with other sedatives	IV			Up to 0.5µg/kg	Up to 0.5µg/kg every 2 mins	1 - 2	3 - 5
Fentanyl (4)	Sedation/Analgesia	IV			Up to 0.5 to 1µg/kg	0.5 - 1.0µg/kg every 2 mins	1 - 2	3 - 5
Ketofol (ketamine and propofol) (6)	Sedation/Amnesia/Analgesia	IV			0.5mg/kg-0.75mg/kg of both agents		½ - 1	1-2

Table 2: Paediatric: Pharmacological Agents for Procedural Sedation and Analgesia

Agent (reference)	Role	Route	Age (see Appendix 6)	Initial dose - Paediatric	Maximum dose	Repeat dose	Maximum dose	Initial onset time (min)	Peak effect time (min)
Propofol (5)	Sedation/ Amnesia	IV	6months – 2 years	1mg/kg-2mg/kg*		0.5mg/kg every 3-5mins*	3mg/kg*	½ - 1	1 - 2
			> 2 years	0.5 – 1.0 mg/kg*					
Midazolam (5)	Sedation/ Amnesia	IV	6months - 5yrs	0.025-0.05mg/kg **	2mg (single dose)	Up to 0.2mg/kg after 2-5 mins	Total 6mg	1 - 2	3 - 4
			6 – 12 yrs	0.025-0.05mg/kg	2mg (single dose)	0.1mg/kg	Total 10mg		
IM Ketamine (9)	Sedation/ Amnesia/ Analgesia	IM	> 3months only	4-5mg/kg (many authorities/guidelines suggest 2-2.5mg/kg)		Half of first dose: 2-2.5mg/kg (1mg/kg if using lower dose). IM after 5-10mins		½ - 1	1 - 2
IV Ketamine (9)	Sedation/ Amnesia/ Analgesia	IV (over 30-60secs)	> 3months only	1.0mg/kg***		0.5 mg/kg IV after 5-10mins		½ - 1	1 - 2
Ketofol (ketamine and propofol) (5, 6)	Sedation/ Amnesia/ Analgesia	IV	> 6months only	0.5mg/kg propofol and ketamine				½ - 1	1-2

*reduce dose significantly in patients who are debilitated or have decreased cardiac function

** This is BNF dosing regime, many sources suggest higher dose range of 0.05-0.1mg/kg required

*** See RCEM [Ketamine Procedural Sedation for Children in EDs](#) (Feb 2020)

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Review

Usually within three years or sooner if important information becomes available.

Conflicts of Interest

None

Disclaimers

The College recognises that patients, their situations, Emergency Departments and staff all vary. This guideline cannot cover all possible scenarios. The ultimate responsibility for the interpretation and application of this guideline, the use of current information and a patient's overall care and wellbeing resides with the treating clinician.

Research Recommendations

There is a need for research into the safety of combinations of drugs, for example sub-dissociative doses of ketamine and propofol(4). There is also a need to develop a comprehensive database of PSA methodology and outcomes.

Audit standards

There should be documentation that allows the regular audit of the use of pharmacological agents in Procedural Sedation and Analgesia and the outcomes of their use in PSA.

Key words for search

Medline search performed for all publications in English after 2010 including the terms "procedural sedation" and "emergency department".

Appendix 1: Procedural Sedation and Analgesia (PSA) Checklist and Monitoring Proforma

Click [here](#) to download the checklist and proforma

Date Time	Patient name Date of birth affix label Hospital number		
Planned procedure:			
Planned sedation level:		minimal	
		moderate sedation	
		deep sedation	
		dissociative sedation	
Patient factors:			
Age:	<input type="text"/>	yrs	Weight <input type="text"/> Kg
Pregnant:	Yes	No <input type="text"/>	
Relevant co-morbidities	IHD	COPD/asthma	Obese
	Schizophrenia	other: <input type="text"/>	
Allergies			
Normal Medications			
Acute Medications			
Recreational drugs or alcohol			
Previous anaesthetic	Yes <input type="checkbox"/>	No <input type="checkbox"/>	
Anaesthetic complications			
Date and time of last food			
Date and time of last oral fluid intake			
ASA grade (please circle)	ASA I	A normal healthy patient	
	ASA II	A patient with mild systemic disease	
	ASA III	A patient with severe systemic disease	
	ASA IV	A patient with severe systemic disease that is a constant threat to life	
	ASA V	A moribund patient who is not expected to survive without the operation	
Difficult Airway?	no concern/ mild concern/significant concern		
Features to consider:			
BMV ventilation:	beard, no teeth, obesity, trauma, cachexia		
LMA:	Look for characteristics of difficult intubation, Evaluate mouth opening and		
Laryngoscopy:	thyromental distance, assess Mallampati score, look for Obstruction, assess Neck		
Crithyroidotomy:	mobility. (LEMON) Check front of neck.		
Consent: sedation	verbal <input type="text"/>	written <input type="text"/>	lacks capacity <input type="text"/>
procedure	verbal <input type="text"/>	written <input type="text"/>	lacks capacity <input type="text"/>
Preprocedural ECG:	Y <input type="checkbox"/>	N <input type="checkbox"/>	
Pain before procedure	mild (0-3)	moderate (4-6)	severe (7-10)
Pain post-procedure	mild (0-3)	moderate (4-6)	severe (7-10)

		Name	Grade	Speciality		Patient Information	
Sedating Practitioner						Name:	
Procedural Assistant						Hosp No:	Affix patient label:
Nursing staff							
Location for procedure	Resus	Y	N		Other (details)		
Date:							
Time:							
Respiratory rate (bpm)							
SpO2 %							
Oxygen delivered (l/min or %)							
Et CO2							
Blood pressure: Systolic/ Diastolic (mmHg)	240						
	230						
	220						
	210						
	200						
	190						
	180						
	170						
	160						
	150						
	140						
	130						
Heart Rate (bpm)	120						
	110						
	100						
	90						
	80						
	70						
	60						
	50						
	40						
	30						
Drugs	Units						
GCS/ Sedation level							
Level of sedation achieved:	minimal sedation						
	moderate sedation						
	deep sedation						
dissociative sedation							
anaesthesia							
Interventions needed:	none						
	hypotension rx						
	BMV						
	LMA						
ETT							
reversal agent							
other							
Adverse events:	none						
	hypoxia						
	hypotension						
	adverse reaction						
vomiting							
cardiac arrest							
aspiration							
death							
Return to baseline	yes	<input type="checkbox"/>	no	<input type="checkbox"/>	Eating/ drinki	yes	<input type="checkbox"/>
Ambulant	yes	<input type="checkbox"/>	no	<input type="checkbox"/>			<input type="checkbox"/>
Procedure Successful:	yes	<input type="checkbox"/>	no	<input type="checkbox"/>			
Discharge Advice given:	verbal	<input type="checkbox"/>	written	<input type="checkbox"/>	Sedating Practitioner signature:		
Patient satisfaction with procedure:	/10						

Appendix 2: Definitions of Sedation Level

The Joint Commission on Accreditation of Healthcare Organizations in the United States has attempted to define the levels of sedation, which range from minimal sedation to general anaesthesia(3):

- Analgesia; Relief of pain without intentionally producing a sedated state. Altered mental status may occur as a secondary effect of medications administered for analgesia.
- Minimal sedation (anxiolysis); The patient responds normally to verbal commands. Cognitive function and coordination may be impaired, but ventilatory and cardiovascular functions are unaffected.
- Moderate sedation and analgesia; The patient responds purposefully to verbal commands alone or when accompanied by light touch. Protective airway reflexes and adequate ventilation are maintained without intervention. Cardiovascular function remains stable.
- Deep sedation and analgesia; The patient cannot be easily aroused, but responds purposefully to noxious stimulation. Assistance may be needed to ensure the airway is protected and adequate ventilation maintained. Cardiovascular function is usually stable.
- General anaesthesia; The patient cannot be aroused and often requires assistance to protect the airway and maintain ventilation. Cardiovascular function may be impaired.
- Dissociative sedation; Dissociative sedation is a trance-like cataleptic state in which the patient experiences profound analgesia and amnesia, but retains airway protective reflexes, spontaneous respirations, and cardiopulmonary stability. Ketamine is the pharmacologic agent used for procedural sedation that produces this state (see 'Ketamine' Appendix 2).

Sedation exists on a continuum, and is difficult to divide into discrete clinical stages. Many sedatives can cause rapid changes in the depth of sedation. Dissociative sedation is considered to be separate from the continuum of sedation.

Appendix 3: Pharmacological Agents for PSA

Entonox:

Properties: Entonox is a gaseous 1:1 mixture of nitrous oxide and oxygen (to prevent hypoxaemia) that is inhaled. It has a rapid onset (30secs) and short duration of action (1 min). Thought to act by binding to opiate receptors in the CNS.

Uses: Sedation, anxiolysis and minor analgesic properties.

Administration: Inhaled but requires patient to be able to hold the inhaler, does not require venous access.

Side effects: Requires a well ventilated room to prevent exposure of clinician, can cause vomiting, contraindicated in head injury, pneumothorax, bowel obstruction, sinus disease.

Propofol:

Properties: Phenol derivative, highly lipophilic, crosses blood brain barrier rapidly. Action thought to be through positive modulation of GABA inhibitory neurotransmission. Used widely for induction and maintenance of anaesthesia.

Uses: Sedation and amnesia.

Administration: Much smaller doses are required for sedation than for general anaesthesia, with initial doses as low as 10 mg in the elderly or those with significant co-morbidities although the initial action may be seen within 30 s, the peak effect may take 2 min or more, particularly in the elderly. An alternative technique is to use a computer-controlled infusion, which estimates the administration profile required for a target plasma concentration, normally 0.5-1.5 µg/ml for sedation.

Children: Propofol is not licensed for use in sedation in children, however the NICE Guideline Development Group recommend off-license use of propofol for sedation in children(2).

Side effects: Hypotension, respiratory depression, pain at site of injection

Allergens: soya and egg lecithin in formulation

Evidence: American College of Emergency Physicians Clinical Policy level of evidence of safety for use in PSA in the emergency department: Level A for paediatrics, Level A for adults.

Midazolam:

Properties: Short-acting benzodiazepine metabolised in the liver commonly used for sedation, it is also a powerful amnesic.

Uses: Sedation and amnesia

Administration: Can be administered orally, buccally or as an IV infusion.

Children: Midazolam is not licensed for use in children under 6 months or for sedation either via the oral or buccal route in children. There is no UK marketing authority currently for oral or intranasal midazolam use in sedation.

Side effects: respiratory depression, hypotension, paradoxical disinhibition and agitation at low doses in children. Accumulates in adipose tissue, which can significantly prolong sedation. The elderly, obese and patients with hepatic or renal disease are at risk of prolonged sedation.

Ketamine:

Properties: Phencyclidine derivative that produces a dissociative state and profound analgesia with superficial sleep. Ketamine does not display a dose-response continuum as seen with other analgesic and sedative agents. There is a threshold dose for dissociation after which additional doses are required only to maintain the dissociative state. Sub-dissociative doses provide analgesia with disorientation rather than dissociation.

Uses:

1. dissociative state, amnesia and analgesia
2. analgesia

Administration: Can be administered in IV boluses, as an IV infusion or intramuscularly.

Side effects: tachycardia, hypertension, laryngospasm, unpleasant hallucinations (reduced by pre-medication with a benzodiazepine), nausea and vomiting, hyper-salivation, increased intracranial and intraocular pressure.

Contraindications: Absolute contraindications: age less than 3 months, known or suspected schizophrenia. Relative contraindications: age less than 1 year, active pulmonary disease or infection, known or suspected cardiovascular disease (including angina, hypertension and heart failure), CNS masses, abnormalities or hydrocephalus, globe injury or glaucoma (9).

Coadministered pharmacological agents(9):

1. Prophylactic anticholinergics have been used in the past to reduce hyper-salivation; however, there is no support in the literature to any benefit of prophylactic co-administration.
2. Benzodiazepines (primarily midazolam) have been recommended in the past to minimize recovery reactions. The literature is not supportive of prophylactic use of midazolam in children but there may be benefit in adults (single controlled study). Benzodiazepines can be used, when required to treat unpleasant reactions if they occur.
3. Anti-emetics are not considered mandatory, a trial using ondansetron showed a significant decrease in emesis (8%) in children; however, this evidence was not considered strong enough to mandate the use of anti-emetics.

Evidence: American College of Emergency Physicians Clinical Policy level of evidence of safety for use in PSA in the emergency department: Level A for paediatrics, Level C for adults.

Ketofol:

Properties: Combination of ketamine and propofol used at a 1:1 ratio in the same syringe in the belief that the lower doses reduce side-effects (hypotension and vomiting and emergence phenomena respectively) of the agents and that the agents act synergistically.

Uses: sedation, amnesia and analgesia

Administration: 1:1 mixture in the same syringe.

Side effects: as for propofol and ketamine

Evidence: American College of Emergency Physicians Clinical Policy level of evidence of safety for use in PSA in the emergency department: Level B for paediatrics, Level B for adults.

Fentanyl:

Properties: A synthetic opioid with 72-125x potency of morphine. Rapid onset (2-3 minutes) duration of effect 30-60mins.

Uses: analgesia and sedation

Administration: Best given a few minutes before sedation to maximise analgesic effect. Doses of no more than 0.5 mcg/kg with other sedation agents.

Side effects: Respiratory depression potentiated by sedatives (e.g. propofol). Patients with renal or hepatic disease and the elderly may experience more profound or prolonged effects.

Appendix 4: Methodology

Where possible, appropriate evidence has been sought and appraised using standard appraisal methods. High quality evidence is not always available to inform recommendations. Best Practice Guidelines rely heavily on the consensus of senior emergency physicians and invited experts.

American College of Emergency Physicians Definitions of Evidence Levels

American College of Emergency Physicians Clinical Policy: level of evidence of safety for use in PSA in the emergency department (6)

Agent	Children	Adults
Ketamine	Level A	Level C
Propofol	Level A	Level A
Ketofol	Level B	Level B

A = Generally accepted principles of patient care based on evidence from 1 or more randomised control trials/ meta-analysis or multiple non-randomised control trials.

B = Recommendations for patient care based on 1 or more non-randomised control trials or multiple case series or reports.

C = Recommendations for patient care based on case series or reports and/ or expert consensus.

Appendix 5: World SIVA adverse sedation event-reporting tool

World SIVA adverse sedation event reporting tool			
<p>World SIVA adverse sedation event recording tool configured for a web page or paper form. Completion of this tool requires execution of all five steps. Responses to each step will often occupy different columns.</p>			
Step 1: Was there one or more adverse events associated with this sedation encounter?			
<input type="radio"/> No, this form is now complete. <input type="radio"/> Yes, fill out remainder of form below.			
Step 2: Please DESCRIBE the adverse events(s). Check all that apply.			
<i>Minimal risk descriptors</i>	<i>Minor risk descriptors</i>	<i>Sentinel risk descriptors</i>	
<input type="radio"/> Vomiting / Retching <input type="radio"/> Subclinical respiratory depression ^a <input type="radio"/> Muscle rigidity, myoclonus <input type="radio"/> Hypersalivation <input type="radio"/> Paradoxical response ^b <input type="radio"/> Recovery agitation ^c <input type="radio"/> Prolonged recovery ^d	<input type="radio"/> Oxygen desaturation (75–90%) for <60 s <input type="radio"/> Apnoea, not prolonged <input type="radio"/> Airway obstruction <input type="radio"/> Failed sedation ^e <input type="radio"/> Allergic reaction without anaphylaxis <input type="radio"/> Bradycardia ^f <input type="radio"/> Tachycardia ^f <input type="radio"/> Hypotension ^f <input type="radio"/> Hypertension ^f <input type="radio"/> Seizure	<input type="radio"/> Oxygen desaturation, severe (<75% at any time) or prolonged (<90% for >60 s) <input type="radio"/> Apnoea, prolonged (>60 s) <input type="radio"/> Cardiovascular collapse/shock ^g <input type="radio"/> Cardiac arrest/absent pulse	<input type="radio"/> Other, specify below
Step 3: Please note the INTERVENTIONS performed to treat the adverse events(s). Check all that apply.			
<i>Minimal risk</i>	<i>Minor risk</i>	<i>Moderate risk</i>	<i>Sentinel intervention</i>
<input type="radio"/> No intervention performed Administration of: <input type="radio"/> Additional sedative(s) <input type="radio"/> Antiemetic <input type="radio"/> Antihistamine	<input type="radio"/> Airway repositioning <input type="radio"/> Tactile stimulation or the administration of: <input type="radio"/> Supplemental oxygen, new or increased <input type="radio"/> Antisialogogue	<input type="radio"/> Bag valve mask-assisted ventilation <input type="radio"/> Laryngeal mask airway <input type="radio"/> Oral/nasal airway <input type="radio"/> CPAP or the administration of: <input type="radio"/> Reversal agents <input type="radio"/> Rapid i.v. fluids <input type="radio"/> Anticonvulsant i.v.	<input type="radio"/> Chest compressions <input type="radio"/> Tracheal intubation or the administration of: <input type="radio"/> Neuromuscular block <input type="radio"/> Pressor / epinephrine <input type="radio"/> Atropine to treat bradycardia
Step 4: Please note the OUTCOME of the adverse events(s). Check all that apply.			
<i>Minimal risk outcome</i>	<i>Moderate risk outcome</i>	<i>Sentinel outcome</i>	
<input type="radio"/> No adverse outcome	<input type="radio"/> Unplanned hospitalisation or escalation of care ^h	<input type="radio"/> Death <input type="radio"/> Permanent neurological deficit <input type="radio"/> Pulmonary aspiration syndrome ⁱ	<input type="radio"/> Other, specify below
Step 5: Assign a SEVERITY rating to the adverse event(s) associated with this sedation encounter.			
<input type="radio"/> If there are any options checked in the Sentinel columns above, then this is a Sentinel ^j adverse event. <input type="radio"/> If the most serious option(s) checked above are Moderate risk, then this is a Moderate ^k risk adverse event. <input type="radio"/> If the most serious option(s) checked above are Minor risk, then this is a Minor ^l risk adverse event. <input type="radio"/> If the most serious option(s) checked above are Minimal risk, then this is a Minimal ^m risk adverse event.			
Additional details (including 'other' entries):			
Footnotes: a. "Subclinical respiratory depression" is defined as capnographic abnormalities suggesting respiratory depression that do not manifest clinically. b. "Paradoxical response" is defined as unanticipated restlessness or agitation in response to sedatives. c. "Recovery agitation" is defined as abnormal patient affect or behaviors during the recovery phase that can include crying, agitation, delirium, dysphoria, hallucinations, or nightmares. d. "Prolonged recovery" is defined as failure to return to baseline clinical status within 2 hours. e. "Failed sedation" is defined as inability to attain suitable conditions to humanely perform the procedure. f. Alteration in vital signs (bradycardia, tachycardia, hypotension, hypertension) is defined as a change of >25% from baseline. g. "Cardiovascular collapse/shock" is defined as clinical evidence of inadequate perfusion. h. Examples of "escalation of care" include transfer from ward to intensive care, and prolonged hospitalisation. i. "Pulmonary aspiration syndrome" is defined as known or suspected inhalation of foreign material such as gastric contents into the respiratory tract associated with new or worsening respiratory signs. j. "Sentinel" adverse events are those critical enough to represent real or serious imminent risk of serious and major patient injury. Once recognized, they warrant immediate and aggressive rescue interventions. Once clinically concluded, they warrant immediate reporting within sedation care systems, and the highest level of peer scrutiny for continuous quality improvement. k. "Moderate" adverse events are those that, while not sentinel, are serious enough to quickly endanger the patient if not promptly managed. Once clinically concluded, they warrant timely reporting within sedation care systems, and periodic peer scrutiny for continuous quality improvement. l. "Minor" adverse events are those encountered periodically in most sedation settings, and that pose little threat given appropriate sedationist skills and monitoring. m. "Minimal" adverse events are those that alone present no danger of permanent harm to the patient.			

Appendix 6

Procedure: The degree of pain and anxiety caused by the procedure and the length of time for the procedure.

Planned level of sedation and analgesia: The planned level of sedation should be decided prior to the procedure and any deviation from the planned level, whether intentionally or unintentionally should be recorded.

Practitioner Training and Familiarity: The sedating practitioner should have familiarity with using the pharmacological agent(s) chosen for the procedure.

Patient factors/ pre-sedation assessment:

Paediatric patients: Both pharmacological and non-pharmacological approaches that are developmentally appropriate should be considered to address anxiety and pain to optimise the success of PSA. Relative contraindications to PSA in paediatric patients include difficult airway or significant co-morbidities, sleep apnoea, special needs, and decreased GI motility (5). **Given the increasing risk of airway complications in younger ages, extreme caution is advised under the age of 2 years, especially when using benzodiazepines, and many local polices and RCEM sedation guidance 2009 limit this.**

Pregnant patients: Additional considerations include positioning, oxygenation, foetal monitoring and pre-procedural medication. In late second and third trimester, left lateral displacement of the uterus will reduce risk of hypotension and foetal hypoxaemia. Oxygenation should be provided. Pre-procedural administration of antacid and anti-emetic such as metoclopramide should be considered.

Older patients: Older patients may be at increased risk of complications with PSA particularly when deeply sedated (7).

Patients with significant co-morbidities: Co-morbidities such as anaemia, heart failure, COPD, dehydration and neuromuscular disease make patients more susceptible to the cardiorespiratory side effects of sedatives.

For older patients and those with significant co-morbidities, it is recommended that lower starting doses with slower rates of administration and repeated doses at less frequent intervals are employed (4).

Other factors: anaesthetic complications, allergies, weight, normal medications, acute medications, recreational drugs including alcohol, anxiety should be assessed.

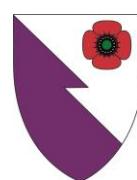
Fasting: The current recommendation is that PSA in the ED should not be delayed in adults or in paediatrics based on fasting time as there is no demonstrated reduction in risk of vomiting or aspiration (Level B evidence) (6).

Weight: Accurate weight should be recorded and used to calculate appropriate doses.

Environment: Time of day, availability of appropriate staff and facilities, availability of alternative approaches (theatre) should be a strong consideration in deciding what level of PSA should be performed.

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