

Review Article

Paediatric procedural sedation for radiological imaging

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Abstract

Sedation is frequently undertaken for radiological imaging procedures in paediatric patients. Movement during procedure degrades all images of a particular sequence. A deeper level of sedation is needed. The sedation of children is different from the sedation of adult. The safe sedation of children for imaging procedure requires a systematic approach that includes the followings. Careful presedation health evaluation of the child with ASA classification. Appropriate fasting guidelines for sedation procedure. Detailed airway examination for any airway abnormalities that might increase the potential for airway obstruction. Adequate training and skills of sedating personnel in paediatric airway management. Age and size appropriate equipment for airway management and venous access. Adequate medications to combat adverse events. Monitoring of vital parameters during and after the procedure. A properly equipped and staffed recovery area. Recovery to presedation level of consciousness of patient before discharge from medical supervision and appropriate discharge instructions. The whole procedure should be well documented. Children who has contraindications to sedation should be selected for general Anaesthesia.

This review article has been made to discuss the need for sedation of children during radiological imaging, currently practiced different regimens of sedation, safe guidelines for sedation and also covers the debate between need for GA versus sedation.

Keywords : Procedural sedation, Paediatric, Radiological Imaging.

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Introduction

Sedation is usually a need for radiological imaging procedures in paediatric populations. Among them magnetic resonance imaging (MRI) takes a longer duration than other radiological imaging procedures and any movement usually degrades all images of a particular sequence. Mild and moderate sedation is unable to guarantee patients compliance and therefore a deeper level of sedation is required. Sedation in children is often administered to control behavior to allow safe completion of procedure. Children younger than 7 Years and those with developmental delay often require deep level of sedation to gain control their behavior.¹ Children in this group are particularly

vulnerable to sedating medications due to effect on respiration, and protective reflexes.² However, general anaesthesia cannot be organized routinely due to need for special, costly equipments, monitors and personnel as well as general anaesthesia is not without risk.

In practice, anaesthesiologists have to deal with these patients with request from radiologist, paediatricians and other clinical staff. There are various general protocols and standard operating procedures,³ made by medical and nursing organizations and societies. They are often general in nature, and thus the anaesthesiologists should design specific protocols to use in their own hospitals for their personnel.

The Need for Deep Sedation

As movement interferes with effective MRI, patients unable to lie still, provide a challenging problem. Moderate sedation is unable to guarantee patient compliance and therefore a deeper level of sedation is required. Infants may go to sleep with a feed and children older than 7 years can comply with instructions to remain still.^{4,5-14} Intravenous deep sedation is thus required for many of those between one and 7 years, and some older children with learning difficulties or claustrophobia.¹⁵

Intravenous sedation is more predictable in this group as it has an immediate effect and is much less reliant on other factors such as absorption. However, it has also been suggested that there are varying levels of deep sedation at the end, which is an overlap with general anaesthesia.¹⁶⁻²⁰

Sedation Regimens for Children

There has been debate over appropriate drugs and their dosage, and those who sedate children have their favorite regimens. It is important that persons administering the drug are familiar with them and cocktails of more than two drugs are to be avoided because of unpredictability of drug interactions and the increased incidence of important side effects.^{3,21,22}

Table I : *Different sedation regimens with dose and route of administration*

Drug Regimen	Dose/route of administration
1. Chloral hydrate	50-100 mg/kg PO
2. Pentobarbital	4-6 mg/kg IV or PO
3. Midazolam	0.5-0.75 mg/kg PO 0.025-0.5 mg/kg IV
4. Propofol	0.2 mg/kg intranasal
5. Methohexital	100-200 µg/kg/min IV 0.25-0.50 mg/kg IV
6. Ketamine	20-25 mg/kg rectal 3-4 mg/kg IM 1-2 mg/kg IV
7. Propofol with fentanyl	Propofol 50-150 µg/kg IV Fentanyl 1- 2 µg/kg IV
8. Midazolam with fentanyl	Midazolam 0.02 mg/kg IV Fentanyl 1-2 µg/kg IV

PO - Per Os (Oral)
IV - Intravenous
IM - Intramuscular

Chloral hydrate is an extremely useful and safe and can be used with good effect in children upto 10 kg.²⁴ Pentobarbital has a long history of effective use but emergence can be prolonged. Midazolam has track record of safe use both oral and intravenous, paradoxical reactions are not frequent. Intranasal route should not be recommended due to irritation.²⁵ Propofol is an ideal agent for nonpainful diagnostic procedures but only for use by expert airway managers with good backup systems.²³ Methohexital gives effective sedation in intravenous route and rectal route is not recommended because of high frequency of apnoea and desaturation events.²⁶ Ketamine is a very popular drug for effective sedation and analgesia for painful procedures, nausea and vomiting is relatively common after procedure and there are reports of laryngospasm.²⁸ Propofol combined with fentanyl is best for deep sedation to anaesthesia, but risk of requiring advanced airway management is high.²⁷ Midazolam with fentanyl is another common combination for painful procedures but risk of apnoea and hypoxia is significant.²⁹

Sedating Personnel

Present American Pediatric Guidelines¹⁶ augmented by a literature suggests that if deep sedation is required then it should be performed by someone.^{30,31}

- Who is working to an acceptable guideline.
- With sole responsibility for the sedation.
- Who has been trained to an acceptable level (Such as, advanced paediatric life support provider status).
- Who is familiar with the drugs, dosages, monitoring equipment, and requirement of the procedure.
- Who is supported by other skilled staff such as a children's nurse.

Preparation of Patient

Children should be prepared in a similar way to child undergoing general anaesthesia.

- Informed consent for sedation taken.
- Children fasted by withholding milk and solids as per direction.
- Reliable intravenous access essential before, during, after the procedure if intravenous drugs are used, in addition, for administration of resuscitation drugs if required.

d. The paediatric patient should be accompanied to and from the scanning department by a parent, legal guardian or other responsible person.

Monitoring of Patient

Monitoring of the patient during and following procedure is the cornerstone of safe practice.^{16,21} There must be one person available whose only responsibility is to constantly observe the patient's vital signs, airway patency, and adequacy of ventilation and administration of drugs. Pulse oximetry is essential during sedation and recovery; oxygen saturation should stay above 93%. Vital signs such as level of consciousness, pulse, respiratory rate and oxygen saturation readings should be taken at given minute's intervals during the procedure and any adverse events must be recorded. Non-ferromagnetic monitoring systems are available and therefore compatible with use in the MRI.³²

Post Sedation Care

After completion of imaging procedure, monitoring and observations should be continued and recorded until recovery of consciousness. All the resuscitation equipment should be readily available in the recovery area. Post anaesthesia recovery nurses with paediatric education and experience are required. Nurses providing such care should be capable managing the paediatric airway and skilled in basic resuscitation techniques.

Recommended Discharge Criteria³³

- a. Cardiovascular function and airway patency are satisfactory and stable.
- b. The patient is easily arousable, and protective reflexes are intact.
- c. The patient can talk (if age appropriate).
- d. The patient can sit up unaided (if age appropriate).
- e. For a very young or handicapped child incapable of the usually expected responses, the pre-sedation level of responsiveness or a level as close as possible to the normal level for that child should be achieved.
- f. The state of hydration is adequate.

Documentation

As like any anaesthetic proper documentation should be done throughout the whole procedure of sedation.³⁴⁻³⁷

- a. Documentation before sedation
 - (i) Informed consent for sedation.
 - (ii) Patients detail particulars.
 - (iii) A complete health evaluation with ASA classification as discussed.
 - (iv) Instructions and information's about fasting and sedation provided to the responsible person.
 - (v) Any special instructions for individual case.
- b. Documentation at the time of sedation.
 - (i) Previous health evaluation will be thoroughly reviewed by sedation team.
 - (ii) Baseline vital parameters of child including patients level of consciousness and responsiveness, heart rate, blood pressure, respiratory rate and oxygen saturation.
- c. Documentation during sedation.
 - (i) The patients chart shall contain a time based record that includes the name, route, site, time, dosage and patient effect of administered drugs.
 - (ii) The patients chart shall contain documentation during sedation the patients level of consciousness, heart rate, blood pressure, respiratory rate, and oxygen saturation at regular interval. This documentation will be continued until the patient attained predetermined discharge criteria.
 - (iii) Any adverse event and treatment of that shall be documented.
- d. Documentation after sedation. The time and condition of the child at discharge from the sedation area or facility shall be documented; this should include documentation that the child's level of consciousness and oxygen saturation in room air have returned to a state that is safe for discharge by recognized criteria.

Resuscitation Equipment

Following resuscitation equipment should be available during procedure and recovery and should include.^{16,21}

- a. Suction apparatus - size appropriate suction catheters and a functioning suction apparatus.
- b. Oxygen - adequate oxygen supply and functioning flow meters/other devices to allow its delivery.
- c. Airway – size appropriate airway equipment, nasopharyngeal and oropharyngeal airways, laryngoscope blades, endotracheal tubes, stylets, facemask, bag-valve-mask or equivalent device (functioning).
- d. Monitors – functioning pulse oximeter with size appropriate oximeter probes and other monitors as appropriate for the procedure (e.g. non-invasive blood pressure, end tidal carbon dioxide, ECG and stethoscope).
- e. Special equipment or drugs for a particular case (e.g. defibrillator).

Resuscitation Drugs^{16,21}

Sulbutamol for nebulization and inhalation, Suxamethonium, Atropine, Diazepam, Adrenaline (1:1000, 1:10,000), Flumazenil, Glucose (25% or 50%), Lignocaine (cardiac lignocaine, local infiltration), Midazolam, Hydrocortisone, Methylprednisolone, Naloxone, Oxygen, Vecuronium, Sodium bicarbonate,

Contraindications of Sedation

- a. Potential airway obstructions for example, sleep apnoea, anatomic airway abnormalities or extreme tonsillar hypertrophy.
- b. Respiratory centre abnormalities for example, brain stem tumours.
- c. Respiratory centre desensitized to carbon dioxide for example, conditions with chronically raised PaCO₂.
- d. Renal or hepatic dysfunction leading to altered drug kinetics.
- e. Conditions in which a rise in PaCO₂ would be detrimental for example raised intracranial pressure.
- f. Conditions with high risk of pulmonary aspiration of gastric content.

These candidates should be exclusively selected for general anaesthesia.

Adverse Events with Sedation in Children³⁸⁻⁵³

Sedation of pediatric patients has serious associated risks, such as

- a. Hypoventilation.
- b. Apnoea.
- c. Airway obstruction.
- d. Hypothermia.
- e. Reaction to contrast agent.
- f. Cardiopulmonary impairment.

Common Causes of Adverse Events

- a. Drug overdose.
- b. Inadequate monitoring.
- c. Premature discharge.
- d. Inadequate help.
- e. Drug interaction.
- f. Drug error.

These adverse responses during and after sedation can be minimized. Careful pre-procedure assessment, appropriate drug selection, appropriate monitoring, as well as the presence of a skilled individual needed to rescue a patient from an adverse responses are essential.^{2,43,44,54}

Ketamine-The Most Common and Effective Paediatric Sedation Agent for Radiological Imaging

The phencyclidine derivative ketamine has been described as a safe and effective paediatric sedation agent in the developed as well as developing world.⁵⁵ Ketamine produces a dissociative state, combination of analgesia, amnesia and sedation at subanaesthetic doses, with minimal effects on the airway and vital reflexes. It is best if combined with an anticholinergic to control secretion and with a benzodiazepine like diazepam to prevent agitation and nightmares.⁵⁶ Ketamine is a safe, useful procedure sedation agent but it delays recovery when used with long acting benzodiazepine like diazepam.⁵⁷⁻⁵⁹ Ketamine does not fit easily into standard drug classification. At low doses full general anaesthesia is not achieved rather a dissociative state in which airway and respiratory tone are maintained. The specific dangers of airway compromise and cardio respiratory instability are suggested to be less with ketamine.

In developing country like Bangladesh, cost of drug is a matter of consideration during sedation procedure. Ketamine is easily available in Bangladesh and more cost effective than other sedation agents and can safely administered for paediatric sedation during radiological imaging.

Sedation versus General Anaesthesia

MRI investigations in children can be done by safe sedation providing previously stated guidelines were followed. Newer short acting drugs like propofol, midazolam, fentanyl etc can be safely employed to provide sedation. Deep sedation has been found to cause respiratory adverse events in ASA III or IV paediatric patients.⁶⁰ Early identification of patients who are at risk of failing sedation or experiencing adverse events may help in choosing patients for whom general anaesthesia would be a safer or successful alternative.⁶¹ Advantages may be achieved by using general anaesthesia instead of deep sedation. There should be less failure and there may be faster turn around. The disadvantages of general anaesthesia include need for costly dedicated equipment, and a greater availability of paediatric anaesthesiologists.⁶²

Conclusion

Providing sedation to children during radiological imaging procedures is an area of rapid change marked by evolving standards. It is possible to use deep sedation to produce satisfactory conditions for children having scans unless there is any contraindication for sedation. Anaesthesiologists have played a critical role in establishing guidelines for safe sedation, considerable work remains in defining what represents effective and safe practice. It is the time that anaesthesiologists should establish the identity as the ultimate experts in this field with a proven track record of practice improvement to assure that clinical practice, training, and safety in this field.

References

1. Maxwell LG, Yester M. The myth of conscious sedation. *Arch pediatr Adolesc Med* 1996; 150:665-667.
2. Cote CJ, Notterman DA, Karl HW, Weinberg JA, McClosky C. Adverse sedation events in pediatrics: a critical incident analysis of contributory factors. *Pediatrics* 2000; 105:805-814.
3. Royal Colleges of Anaesthetists and Radiologists. Report of a joint working party. Sedation and anaesthesia in radiology. London: Royal colleges of Anaesthetists and Radiologists, 1992.
4. Kennedy RM, Luhmann JD. The "ouchless emergency department." Getting closer: Advances in decreasing distress during painful procedures in the emergency department. *Pediatr Clin North Am* 1999;46:1215-47.
5. Newton JT, Shah S, Patel H, Sturme P. Non-pharmacological approaches to behaviour management in children. *Dent Update* 2003;30:194-9.
6. Peretz B, Bimstein E. The use of imagery suggestions during administration of local anesthetic in pediatric dental patients. *ASDC J Dent Child* 2000;67:263-7.
7. Iserson KV. Hypnosis for pediatric fracture reduction. *J Emerg Med* 1999;17:53-6.
8. Rusy LM, Weisman SJ. Complementary therapies for acute pediatric pain management. *Pediatr Clin North Am* 2000;47:589-99.
9. Ott MJ. Imagine the possibilities! Guided imagery with toddlers and pre schoolers. *Pediatr Nurs* 1996;22:34-8.
10. Singer AJ, Stark MJ. LET versus EMLA for pretreating lacerations: A randomized trial. *Acad Emerg Med* 2001; 8:223-30.
11. Taddio A, Gurguis MG, Koren G. Lidocaine-prilocaine cream versus tetracaine gel for procedural pain in children. *Ann Pharmacother* 2002;36:687-92.
12. Eichenfield LF, Funk A, Fallon-Friedlander S, Cunningham BB. A clinical study to evaluate the efficacy of ELA-Max (4% liposomal lidocaine) as compared with eutectic mixture of local anesthetics cream for pain reduction of venipuncture in children. *Pediatrics* 2002;109:1093-9.
13. Shaw AJ, Welbury RR. The use of hypnosis in a sedation clinic for dental extractions in children: Report of 20 cases. *ASDC J Dent Child* 1996;63:418-20.
14. Aitken JC, Wilson S, Coury D, Moursi AM. The effect of music distraction on pain, anxiety and behavior in pediatric dental patients. *Pediatr Dent* 2002;24:114-8.
15. Sury MRJ, Hatch DJ, Deeley T, Dicks-Mireaux C, Chong WK. Development of a nurse-led sedation service for paediatric magnetic resonance imaging. *Lancet* 1999; 353:1667-71.

16. American Academy of Pediatrics Committee on Drugs. Guidelines for monitoring and management of pediatric patients during and after sedation for diagnostic and therapeutic procedures, *Pediatrics* 1992; 89: 1110-15.
17. Dial S, Silver P, Bock K, Sagy M. Pediatric sedation for procedures titrated to a desired degree of immobility results in unpredictable depth of sedation. *Pediatr Emerg Care* 2001;17:414-20.
18. Maxwell LG, Yaster M. The myth of conscious sedation. *Arch Pediatr Adolesc Med* 1996;150:665-7.
19. Motas D, McDermott NB, VanSickle T, Friesen RH. Depth of consciousness and deep sedation attained in children as administered by nonanaesthesiologists in a children's hospital. *Pediatr Anaesth* 2004;14:256-60.
20. Malviya S, Voepel-Lewis T, Tait AR, Merkel S, Tremper K, Naughton N. Depth of sedation in children undergoing computed tomography: Validity and reliability of the University of Michigan Sedation Scale (UMSS). *Br J Anaesth* 2002;88:241-5.
21. American Academy of Pediatrics, American Academy of Pediatric Dentistry, Cote CJ, Wilson S, Work group on sedation. Guidelines for monitoring and management of pediatric patients during and after sedation for diagnostic and therapeutic procedures; an update. *Pediatrics* 2006; 118(6):2587-602.
22. Goodson JM, Moore PA. Life-threatening reactions after ped-odontic sedation: an assessment of narcotic, local anaesthetic, and antiemetic drug interaction. *J Am Dent Assoc* 1983; 107: 239-45.
23. Scheiber G, Ribeiro FC, Karpinski H, Strehl K. Deep Sedation with propofol in preschool children undergoing radiation therapy. *Pediatr Anaesth* 1996; 6:209-13.
24. Rooks VJ, Chung T, Connor L, et al. Comparison of oral pentobarbital sodium (Nembutal) and oral chloral hydrate for sedation of infants during radiologic imaging: preliminary results. *Am J Roentgenol* 2003; 180:1125-1128.
25. Harcke HT, Grisson LE, Meister MA. Sedation in pediatric imaging using intranasal midazolam. *Pediatr Radiol* 1995; 25: 341-343.
26. Pomeranz ES, Chudnofsky CR, Deegan TJ, et al. Rectal methohexital sedation for computed tomography imaging of stable pediatric emergency patients. *Pediatrics* 2000; 105: 1110-4.
27. Bauman LA, Kish I, Baumann RC, Politis GD. Pediatric sedation with analgesia. *Am J Emerg Med* 1999; 17:1-3.
28. Green SM, Denmark TK, Cline J, et al. Ketamine sedation for pediatric critical care procedures. *Pediatr Emerg Care* 2001; 17 : 244 – 248.
29. Pitetti RD, Singh S, Pierce MC. Safe and efficacious use of procedural sedation and analgesia by non-anaesthesiologists in a pediatric emergency department. *Arch Pediatr Adolesc Med* 2003; 157: 1090-6.
30. American Heart Association. Pediatric Advanced Life Support Provider Manual. Dallas, Tx: American Heart Association; 2002
31. American Academy of Pediatrics, American College of Emergency Physicians. Advanced Pediatric Life Support. 4th ed. Fuchs S, Gausche-Hill M, Yamoto L, eds. Boston, Ma: Jones and Bartlett Publishers; 2004..
32. Peden CJ, Menon DK, Hall AS, et al. Magnetic resonance for anaesthetist. Part II. Anaesthesia and monitoring in MR units. *Anaesthesia* 1992; 47: 508–17.
33. Coté CJ. Discharge criteria for children sedated by nonanesthesiologists: Is “safe” really safe enough?” *Anesthesiology* 2004;100:207-9.
34. Malviya S, Voepel-Lewis T, Ludomirsky A, Marshall J, Tait AR. Can we improve the assessment of discharge readiness? A comparative study of observational and objective measures of depth of sedation in children. *Anesthesiology* 2004;100:218-24.
35. Malviya S, Voepel-Lewis T, Prochaska G, Tait AR. Prolonged recovery and delayed side effects of sedation for diagnostic imaging studies in children. *Pediatrics* 2000;105(3):e42.
36. Mayers DJ, Hindmarsh KW, Sankaran K, Gorecki DK, Kasian GF. Chloral hydrate disposition following single dose administration to critically ill neonates and children. *Dev Pharm & Ther* 1991;16:71-7.
37. Terndrup TE, Dire DJ, Madden CM, Davis H, Cantor RM, Gavula DP. A prospective analysis of intramuscular meperidine, promethazine, and chlorpromazine in pediatric emergency

- department patients. *Ann Emerg Med* 1991;20:31-5.
38. Law AK, Ng DK, Chan KK. Use of intramuscular ketamine for endoscopy sedation in children. *Pediatr Int* 2003;45:180-5.
39. De Blic J, Marchac V, Scheinmann P. Complications of flexible bronchoscopy in children: Prospective study of 1,328 procedures. *Eur Respir J* 2002;20:1271-6.
40. Pena BM, Krauss B. Adverse events of procedural sedation and analgesia in a pediatric emergency department. *Ann Emerg Med* 1999;34:483-91.
41. Coté CJ, Karl HW, Notterman DA, Weinberg JA, McCloskey C. Adverse sedation events in pediatrics: Analysis of medications used for sedation. *Pediatrics* 2000; 106:633-44.
43. Hoffman GM, Nowakowski R, Troshynski TJ, Berens RJ, Weisman SJ. Risk reduction in pediatric procedural sedation by application of an American Academy of Pediatrics/American Society of Anesthesiologists process model. *Pediatrics* 2002;109:236-43.
44. Nahata MC, Clotz MA, Krogg EA. Adverse effects of meperidine, promethazine, and chlorpromazine for sedation in pediatric patients. *Clin Pediatr* 1985;24:558-60.
45. Brown ET, Corbett SW, Green SM. Iatrogenic cardiopulmonary arrest during pediatric sedation with meperidine, promethazine, and chlorpromazine. *Pediatr Emerg Care* 2001;17:351-3.
46. Benusis KP, Kapaun D, Furnam LJ. Respiratory depression in a child following meperidine, promethazine, and chlorpromazine premedication: Report of case. *J Dent Child* 1979;46:50-3.
47. Garriott JC, Di Maio VJ. Death in the dental chair: Three drug fatalities in dental patients. *J Toxicol Clin Toxicol* 1982;19:987-95.
48. Goodson JM, Moore PA. Life-threatening reactions after pedodontic sedation: An assessment of narcotic, local anesthetic, and antiemetic drug interaction. *J Am Dent Assoc* 1983;107:239-45.
49. Jastak JT, Pallasch T. Death after chloral hydrate sedation: Report of case. *J Am Dent Assoc* 1988;116:345-8.
50. Jastak JT, Peskin RM. Major morbidity or mortality from office anesthetic procedures: A closed-claim analysis of 13 cases. *Anesth Prog* 1991; 38: 39-44.
51. Kaufman E, Jastak JT. Sedation for outpatient dental procedures. *Compend Contin Educ Dent* 1995;16:462, 464, 466.
52. Wilson S. Pharmacological management of the pediatric dental patient. *Pediatr Dent* 2004;26:131-6.
53. Sams DR, Thornton JB, Wright JT. The assessment of two oral sedation drug regimens in pediatric dental patients. *J Dent Child* 1992;59:306-12.
54. Malviya S, Voepel-Lewis T, Tait AR. Adverse events and risk factors associated with the sedation of children by non-anesthesiologists. *Anesth Analg* 1997;85:1207-13.
55. Green S. Ketamine sedation for paediatric procedures; Part 1, a prospective series. *Ann Emerg Med* 1990; 19:1024-32.
56. Green SM, Johnson NE. Ketamine sedation for pediatric procedures; Part 2, review and implications. *Ann Emerg Med* 1990; 19:1033-46.
57. Vara darajulu S, Elobeidi MA, Tamhane A, Wilcox CM. Prospective randomized trial evaluating ketamine for advanced endoscopic procedures in difficult to sedate patients. *Alimentary Pharmacology & Therapeutics* 2007; 25(8): 987-997.
58. Drummond GB. Comparison of sedation with midazolam and ketamine: effects on airway muscle activity. *Br. J Anaesth* 1996; 76:663-7.
59. Hazma J, Ecoffey C, Gross JB. Ventilatory response to CO_2 following intravenous ketamine in children. *Anaesthesiology* 1989; 70:422-5.
60. Petrack EM, Marx CM, wright MS. Intramuscular ketamine is superior to meperidine, promethazine and chlorpromazine for paediatric emergency department sedation. *Arch Pediatr Adolesc Med* 1996; 150:676-81.
61. Gooden CK, Dilos B, Anaesthesia for magnetic resonance imaging, *Int Anesthesiol Clin.* 2003 Spring 41 (2) 29: 37.
62. Malviya S, Voepel-Lewis T, Eldsik OP, et al. Sedation and general anaesthesia in children undergoing MRI and CT : adverse events and outcomes. *Br J Anaesth* 2000; 84; 743-748.