



TECHNOLOGY STATUS EVALUATION REPORT
**PROPOFOL USE DURING
GASTROINTESTINAL ENDOSCOPY**

NOVEMBER, 2000

INTRODUCTION

In order to promote the appropriate use of new or emerging endoscopic technologies, the ASGE Technology Committee has developed a series of status evaluation papers. By this process relevant information about these technologies may be presented to practicing physicians for the education and care of their patients. In many cases, data from randomized controlled trials is lacking and only preliminary clinical studies are available. Practitioners should continue to monitor the medical literature for subsequent data about the efficacy, safety and socioeconomic aspects of the technologies.

BACKGROUND

Propofol, or 2-6 diisopropylphenol (AstraZeneca, Wilmington, DL; Baxter Pharmaceutical Products, Inc., New Providence, NJ) is an ultrashort acting sedative hypnotic agent that has received increased attention for use during endoscopy.¹⁻⁷ The distinctive sedative properties, pharmacokinetics, and pharmacodynamics form the basis of this report.

PHARMACOLOGIC PROPERTIES

Propofol is an alkyl phenol derivative that possesses sedative, amnestic, and hypnotic properties but provides minimal analgesia.^{8,9} The drug is lipophilic and is prepared as an oil/water emulsion consisting of 1% propofol, 10% soybean oil, 2.25% glycerol, and 1.2% egg lecithin.^{10,11} Propofol is contraindicated in patients with hypersensitivity to egg or soybean. In addition, a generic formulation contains sodium metabisulfite and is contraindicated in patients with sulfite allergies.¹²

Propofol has a rapid onset and a short duration of action. Hypnosis is induced within 30-60 seconds of intravenous administration, essentially the time of one arm-brain circulatory pass.^{8,9} The half-life of propofol is 1.8-4.1 minutes. After cessation of infusion, blood concentrations rapidly decline due to rapid tissue distribution and high metabolic clearance.¹¹ Clinically this accounts for rapid recovery within 10-30 minutes in most patients after discontinuation of the drug.⁹

Propofol is 98% plasma-protein bound, and is metabolized primarily in the liver. Propofol potentiates the effects of benzodiazepines, barbiturates, and opioids.^{9,10,13} The pharmacokinetic properties do not significantly change in patients with moderate chronic liver disease or renal failure.⁸⁻¹⁰ However, dose reductions are indicated in the elderly and in patients with diminished cardiac output due to decreased clearance of the drug.¹⁴

EFFICACY

EGD

A randomized study of 40 patients receiving either propofol or midazolam titrated to an equivalent level of sedation prior to endoscopy reported that propofol provided more rapid recovery compared with midazolam, but was associated with pain on injection, a shorter amnesia span, and reduced patient acceptance.¹⁵ In contrast, another study randomized 90 patients to receive either midazolam or propofol administered both before and during the procedure. Patients receiving propofol tolerated endoscopy better, reached a deeper level of sedation, and recovered more rapidly. There was a similar fre-

quency of amnesia for the procedure and perceived patient discomfort.³

Colonoscopy

An uncontrolled study of 60 patients evaluated different propofol infusion rates after a fixed loading dose during colonoscopy. Patients lost consciousness after a mean of 60.6 seconds and preservation of the hypnotic state was dependent on the infusion rate.¹⁶ A small study of 20 subjects using patient-controlled sedation (PCS) with propofol alone or in combination with alfentanil demonstrated feasibility but suggested that propofol alone did not provide adequate analgesia.¹⁷ A double-blinded study randomized 57 patients to one of three groups: diazepam/meperidine, midazolam/fentanyl, or propofol/fentanyl. There were no significant differences in sedation, analgesia, recovery rate or incidence of side-effects.¹⁸ Another randomized controlled trial compared sedation with pethidine and diazepam versus patient-controlled sedation with propofol and alfentanil. Patient controlled sedation provided significantly lighter sedation, less analgesia, and a faster recovery time (10 vs. 40 minutes). All patients were satisfied with their level of sedation.⁴ Another study randomly assigned 79 patients to receive either midazolam or midazolam plus propofol. The study results are difficult to interpret due to the concomitant administration of nalbuphine and ketamine.⁵

ERCP

Two studies comparing midazolam to propofol during ERCP have been reported.^{6,7} A randomized, controlled, unblinded study of 80 patients found that adequate sedation was possible in 80% of patients with midazolam alone and 97.5% of patients receiving propofol ($p < 0.01$). Recovery times were significantly shorter and sedation was judged by physicians and patients to be significantly better with propofol.⁶ In the second randomized controlled trial involving 198 patients, propofol provided more rapid sedation and significantly better patient cooperation. Recovery time was also significantly shorter with propofol (19 vs. 29 minutes).⁷

Combined studies

Propofol has been evaluated in three studies comprising 545 patients undergoing EGD, colonoscopy and ERCP. The authors concluded that sedation with propofol was comparable to that achieved with conventional agents, while providing for faster recovery time.^{2,19,20}

Pediatric Use

Propofol is not approved for use in children less than 3 years of age.²¹ There is limited published experience

on the use of propofol for endoscopic sedation in the pediatric population.²²⁻²⁴ A retrospective review published in abstract form reported on the successful use of propofol in 115 pediatric patients (mean age 6.4; range 10 days to 20.8 years) undergoing a variety of procedures including endoscopy in an ICU.²²

SAFETY

Propofol is a respiratory depressant with effects including a reduction in minute ventilation, tidal volume, and functional residual capacity.^{25,26} Three studies involving a total of 300 patients receiving propofol for endoscopic sedation each reported an episode of severe respiratory depression.^{6,7,20} In a small study using propofol for endoscopic sedation, apnea was detected by end-tidal capnography in 6 of 10 patients. This enabled a timely decrease in the propofol infusion avoiding significant oxygen desaturation.²⁷

The predominant cardiovascular effect of propofol is a reduction in the systemic vascular resistance, which may induce hypotension.²⁸ When used for general anesthesia, hypotension (systolic blood pressure under 90mmHg) occurred in 15.7% and bradycardia (heart rate below 50) in 4.8% of patients.²⁹

Infections have been reported with the use of contaminated propofol.³⁰⁻³² Due to the rapid growth of organisms in this lipid based medium at room temperature, techniques to minimize contamination are critical. These include adherence to aseptic techniques, avoidance of reusing a syringe, use of propofol within 6 hours of original withdrawal from an ampule, and refrigeration.^{33,34}

Intravenous propofol given by peripheral vein has been reported to cause pain on injection in 30-90% of patients. Reported techniques to minimize this effect include warming the drug to body temperature, dilution, use of lidocaine, or concomitant administration of select sedatives.³⁵⁻⁴⁰

Twenty five cases of pancreatitis associated with propofol use were reported to the food and drug administration by 1996.⁴¹ The mechanism of pancreatitis with propofol has not been established but a causality link is regarded as probable.⁴²

COSTS

The direct cost of medication is increased with propofol compared to opioid and benzodiazepine sedation.^{4,26} The additional cost of monitoring and personnel for sedation has not been weighed against the shortened recovery time or other indirect patient costs.

Cost of Sedative Drugs²⁶

Drug	Amount	Cost
Morphine	10mg	\$0.51
Meperidine	100mg	\$0.41
Fentanyl	100µg	\$0.24
Midazolam	5mg	\$9.33
Propofol	200mg	\$10.20

SUMMARY

Propofol provides effective sedation during gastrointestinal endoscopy with shorter recovery times than other commonly used sedative agents. However, the therapeutic window is narrow and further research on the optimal use of propofol during endoscopy is needed.

REFERENCES

1. Bell GD. Premedication, preparation, and surveillance. *Endoscopy* 2000;32:92-100.
2. Koshy G, Nair S, Norkus EP, Hertan HI, Pitchumoni CS. Propofol versus midazolam and meperidine for conscious sedation in GI endoscopy. *Am J Gastroenterol* 2000;95:1476-9.
3. Carlsson U, Grattidge P. Sedation for upper gastrointestinal endoscopy: a comparative study of propofol and midazolam. *Endoscopy* 1995;27:240-3.
4. Roseveare C, Seavell C, Patel P, Criswell J, Kimble J, Jones C, et al. Patient-controlled sedation and analgesia, using propofol and alfentanil, during colonoscopy: a prospective randomized controlled trial. *Endoscopy* 1998;30:768-73.
5. Reimann FM, Samson U, Derad I, Fuchs M, Schiefer B, Stange EF. Synergistic sedation with low-dose midazolam and propofol for colonoscopies. *Endoscopy* 2000;32:239-44.
6. Jung M, Hofmann C, Kiesslich R, Brakertz A. Improved sedation in diagnostic and therapeutic ERCP: propofol is an alternative to midazolam. *Endoscopy* 2000;32:233-8.
7. Wehrmann T, Kokabpick S, Lembcke B, Caspary WF, Seifert H. Efficacy and safety of intravenous propofol sedation during routine ERCP: a prospective, controlled study. *Gastrointest Endosc* 1999;49:677-83.
8. Smith I, White PF, Nathanson M, Gouldson R. Propofol: an update on its clinical use. *Anesthesiology* 1994;81:1005-43.
9. Marinella JA. Propofol for sedation in the intensive care unit: essentials for the clinician. *Respiratory Medicine* 1997;91:505-10.
10. Bryson HM, Fulton BR, Faulds D. Propofol: an update of its use in anaesthesia and conscious sedation. *Drugs* 1995; 50:513-9.
11. Diprivan 1%. Astra-Zeneca, Wilmington, DL, 2000. (package insert).
12. Propofol. Baxter Pharmaceutical Products, Inc., New Providence, New Jersey, 2000. (package insert).
13. Vuyk J. Pharmacokinetics and pharmacodynamic interactions between opioids and propofol. *J Clin Anesth* 1997;9:23S-26S.
14. Kirkpatrick T, Cockshott ID, Douglas EJ, Nimmo WS. Pharmacokinetics of propofol (Diprivan) in elderly patients. *Br J Anaesth* 1988;60:146-50.
15. Patterson KW, Casey PB, Murray JP, O'Boyle CA, Cunningham AJ. Propofol sedation for outpatient upper gastrointestinal endoscopy: comparison with midazolam. *Br J Anaesth* 1991;67: 108-11.
16. Gepts E, Claeys MA, Camu F, Smekens L. Infusion of propofol ('Diprivan') as sedative technique for colonoscopies. *Postgrad Med J* 1985;61(Suppl. 3):120-6.
17. Heiman DR, Tolliver BA, Weis FR, O'Brien BL, DiPalma JA. Patient-controlled anesthesia for colonoscopy using propofol: results of a pilot study. *Southern Med J* 1998;91:560-4.
18. Kostash MA, Johnston R, Bailey RJ, Konopad EM, Guthrie LP. Sedation for colonoscopy: a double-blind comparison of diazepam/meperidine, midazolam/fentanyl and propofol/fentanyl combinations. *Can J Gastroenterol* 1994;8:27-31.
19. Dubois A, Balatoni E, Peeters JP, Baudoux M. Use of propofol for sedation during gastrointestinal endoscopies. *Anaesthesia* 1988;43(Suppl.):75-80.
20. Tellan G, Fegiz A, Iannarone C, Baumgartner I, Navarra M, Fantera A. The use of di-hydroxypropylphenol (propofol) in endoscopic procedures. *Eur Rev Med Pharmacol Sci* 1998;3-4:147-50.
21. Susla GM. Propofol toxicity in critically ill pediatric patients: show us the proof. *Crit Care Med* 1998;26:1959-60.
22. Hertzog JH, Campbell JK, Dalton HJ, Hauser GJ. Propofol anesthesia for invasive procedures in ambulatory and hospitalized children: experience in the pediatric intensive care unit [abstract]. *Pediatrics* 1999;103:657.
23. Rich JB, Yaster M, Brandt J. Anterograde and retrograde memory in children anesthetized with propofol. *Journal of Clinical and Experimental Neuropsychology* 1999;21:535-46.
24. Lowrie L, Weiss AH, Lacombe C. The pediatric sedation unit: a mechanism for pediatric sedation [abstract]. *Pediatrics* 1998: 102:627.
25. Mirenda J, Broyles G. Propofol as used for sedation in the ICU. *Chest* 1995;108:539-48.
26. Graber RG. Propofol in the endoscopy suite: an anesthesiologist's perspective. *Gastrointest Endosc* 1999;49:803-6.
27. Vargo JJ, Zuccaro G, Dumot JA, Shay SS, Conwell DL, et al. Gastroenterologist-administered propofol for therapeutic upper endoscopy with graphic assessment of respiratory activity. *Gastrointest Endosc* 2000;52:250-5.
28. Turner RJ, Gatt SP, Kam A, Ramzan I, Daley M. Administration of a crystalloid fluid preload does not prevent the decrease in arterial blood pressure after induction of anaesthesia with propofol and fentanyl. *Br J Anaesth* 1998;80: 737-41.
29. Hug CC, McLeskey CH, Nahrwold ML, Roizen MF, Stanley TH, Thisted RA, et al. Hemodynamic effects of propofol: data from over 25,000 patients. *Anesth Analg* 1993;77:S21-9.
30. Anonymous. Postsurgical infections associated with an extrinsically contaminated intravenous anesthetic agent in California, Illinois, Maine, and Michigan. *MMWR* 1990;39: 426-7.
31. Bennett SN, McNeil MM, Bland LA, Arduino MJ, Villarino ME, Perotta DM, et al. Post-operative infections traced to contamination of an intravenous anesthetic propofol. *N Engl J Med* 1995;333:147-54.
32. Veber B, Gachot B, Bedos JP, Wolff M. Severe sepsis after intravenous injection of contaminated propofol. *Anesthesiology* 1994;80:712-3.
33. Patterson JS, Hopkins KJ, Albanese R. Propofol handling techniques. *Acta Anaesthesiol Scand* 1991;35:370
34. Bach A, Motsch J. Infectious risk associated with the use of propofol. *Acta Anaesthesiol Scand* 1996;40:1189-96.
35. Fletcher GC, Gillespie JA, Davidson JAH. The effect of temperature upon pain during injection of propofol. *Anesthesia* 1996;51:498-9.
36. Smith AJ, Power I. The effect of pretreatment with ketorolac on

pain during intravenous injection of propofol. *Anaesthesia* 1996;51:883-5.

37. Angst MS, Mackey SC, Zupfer GH, Tataru CD, Brock-Utne JG. Reduction of propofol injection pain with a double lumen IV set. *J Clin Anesth* 1997;9:462-6.
38. Pang WW, Mok MS, Huang S, Hwang MH. The analgesic effect of fentanyl, morphine, meperidine, and lidocaine in the peripheral veins: a comparative study. *Anesth Analg* 1998;86:382-6.
39. Ho CM, Tsou MY, Sun MS, Chu CC, Lee TY. The optimal effective concentration of lidocaine to reduce pain on injection of propofol. *J Clin Anesth* 1999;11:296-300.
40. Sadler PJ, Thompson HM, Maslowski P, Liddle A, Rowbotham DJ. Iontophoretically applied lidocaine reduces pain on propofol injection. *Br J Anaesth* 1999;82:432-4.
41. Leisure GS, O'Flaherty J, Green L, Jones DR. Propofol and post-operative pancreatitis. *Anesthesiology* 1996;84:224-7.
42. Kumar AN, Schwartz DE, Lim KG. Propofol-induced pancreatitis: recurrence of pancreatitis after rechallenge. *Chest* 1999; 115:1198-9.

**Prepared by:
TECHNOLOGY COMMITTEE**

Douglas B. Nelson, MD, Chair
Alan N. Barkun, MD
Kevin P. Block, MD
J. Steven Burdick, MD
Gregory G. Ginsberg, MD
David A. Greenwald, MD
Peter B. Kelsey, MD
Naomi L. Nakao, MD
Adam Slivka, MD
Paulette Smith, BS, RN, CGRN
Nimish Vakil, MD

**American Society for Gastrointestinal Endoscopy,
13 Elm Street, Manchester, MA 01944-1314
www.asge.org, asge@shore.net**