

Postoperative nausea and vomiting

A comparison of sufentanil, nitrous oxide, and isoflurane

ERIC BLOOMFIELD, MD; MARK HILBERMAN, MD; PHILLIP BROWN, MB, BS; HILDA NOE, CRNA; JOHN GAYDOS, MS; GAIL MILLER, BS; DUANE SHERRILL, MS; GEORGE WILLIAMS, PHD

• Postoperative nausea and vomiting have been associated with the use of nitrous oxide and, in some studies, with isoflurane. Sufentanil, a new synthetic narcotic with a duration as long as fentanyl, was studied with regard to postoperative nausea and vomiting. A total of 63 patients undergoing extra-abdominal procedures (excluding thoracotomies and intracranial, ophthalmologic, and middle-ear surgery) was studied and randomly divided into four groups: Group A, sufentanil/N₂O/O₂ with 0.25% isoflurane; Group B, O₂/N₂O/isoflurane; Group C, O₂/isoflurane/sufentanil; Group D, O₂/isoflurane. Patients with a history of postoperative nausea and vomiting were excluded. The incidence of postoperative nausea and vomiting was observed in the recovery room. The overall incidence of nausea was 25% and of vomiting 9.5%; differences between techniques were not statistically significant.

□ INDEX TERMS: ANESTHETICS; NAUSEA; VOMITING □ CLEVE CLIN J MED 1988; 55:549–552

AUSEA AND VOMITING postoperatively can be a distressing sequel to anesthesia and surgery, especially when early recovery is prejudiced in outpatients. Many factors have been implicated, including anesthetic agents, narcotics, age, gender, weight, type of surgical procedures, and pain.¹ Recently, Alexander et al² have suggested that N_2O can significantly increase the incidence of postoperative nausea and vomiting. We undertook a pilot study to compare the incidence of postoperative nausea and vomiting when using sufentanil, nitrous oxide, and isoflurane.

MATERIALS AND METHODS

Sixty-three patients were studied, all of whom had signed consent forms as required by the Human Subjects Committee of the University of Colorado Health Science Center. Their age range was 18-60 years and their ASA Classification I-III. They were allocated to one of four groups, using random number tables, and received one of the following anesthetics: Group A (n=16), suferial with nitrous oxide/oxygen and 0.25% isoflurane; Group B (n=16), isoflurane with nitrous oxide/oxygen; Group C (n=15), suferial with oxygen and 0.5% isoflurane; Group D (n=16), isoflurane and oxygen. Because of the decreased potency of nitrous oxide in Denver, isoflurane 0.25% was added in Group A to assure amnesia. No premedication was used because of possible gastrointestinal effects. Patients taking antacids or histamine-type 2 beta blockers were excluded from the study, as were morbidly obese patients and those with a history of postoperative nausea and vomiting.

From the Division of Anesthesia, The Cleveland Clinic Foundation, and the Department of Anesthesiology, University of Colorado Health Sciences Center. Submitted June 1987; accepted Aug 1987.

Address reprint requests to E.B., Division of Anesthesiology, The Cleveland Clinic Foundation, One Clinic Center, 9500 Euclid Avenue, Cleveland, Ohio 44195.

Postoperative Nausea \blacksquare Bloomfield and Associates

TABLE 1 TYPES OF CASES

| 1. | Acrominoplasty | 32. | Laminectomy |
|-----|--|-----|---|
| 2. | Parathyroid removal | 33. | Open reduction internal fixation, right ankle |
| 3. | Marshall-Matchetti-Krantz (cystourethroplexy), | 34. | Vaginal hysterectomy |
| | anteriorculporrhapy | 35. | Open reduction internal fixation, left ankle |
| 4. | Cone biopsy | 36. | Laser excision of anal warts |
| 5. | Laparoscopic tubal ligation | 37. | Drainage, sphenoid sinus |
| 6. | Cone biopsy | 38. | Excisional breast biopsy |
| 7. | Open reduction internal fixation, left ulna | 39. | Septoplasty |
| 8. | Inguinal hernia repair | 40. | Open urethral lithotomy |
| 9. | Parotidectomy | 41. | Right mastectomy |
| 10. | Open reduction internal fixation, left hand | 42. | Tonsillectomy |
| 11. | Laser excision of anal warts | 43. | Right inguinal hernia repair |
| 12. | Hemorrhoidectomy | 44. | Hemorrhoidectomy |
| 13. | Inguinal hernia repair | 45. | Chest wall reconstruction |
| 14. | Laparotomy | 46. | Vaginal hysterectomy |
| 15. | Breast removal | 47. | Insertion of penile prosthesis |
| 16. | Fusion of talus | 48. | Cone biopsy |
| 17. | Vein ligation, left leg | 49. | Lumbar sympathectomy |
| 18. | Breast reduction | 50. | Split-thickness skin graft, forearm |
| 19. | Left subclavian vein bypass | 51. | Nerve repair, right ulna |
| 20. | Right hip pinning | 52. | Removal of hardware, left elbow |
| 21. | Laparoscopy | 53. | Incision and drainage, right leg |
| 22. | Inguinal hernia repair | 54. | Incision and drainage, right leg |
| 23. | Revision of medi-port catheter | 55. | Release contracture, right finger |
| 24. | Right inguinal hernia repair | 56. | Split thickness skin graft, left thigh |
| 25. | Benegraft, right mandible | 57. | Right mastectomy |
| 26. | Right temporomandibular joint replacement | 58. | Ventral hernia repair |
| 27. | Tonsillectomy | 59. | Removal of Harrington rod |
| 28. | Resection (submandibular) of hand | 60. | Laser excision of condylomata |
| 29. | Septoplasty | 61. | Transphenoidal tumor excision |
| 30. | Radical hysterectomy | 62. | Rectovaginal fistula repair |
| 31. | Tonsillectomy | 63. | Cone biopsy |
| _ | | | |

TABLE 2 SEX

| TABLE 3 |
|-------------|
| AGE (YEARS) |

| Group | Male | Female | Group | n | Mean | S.D | |
|-------|------|--------|-------|----|------|-----|--|
| A | 9 | 7 | A | 16 | 32 | 11 | |
| В | 8 | 8 | В | 16 | 37 | 13 | |
| Ĉ | 7 | 8 | С | 15 | 39 | 11 | |
| D | 8 | 8 | D | 16 | 37 | 12 | |

The operations all lasted more than 30 minutes and were extra-abdominal in all but two patients (two laparoscopies). Patients undergoing intracranial, ophthalmic, middle-ear, and intrathoracic procedures were excluded. The duration of anesthesia extended over similar ranges for all four groups, and the median durations did not differ significantly: Group A, 30–250 minutes, median 78 minutes; Group B, 25–230 minutes, median 72 minutes; Group C, 50–445 minutes, median 115 minutes; Group D, 35–295 minutes, median 118 minutes. Patients were monitored by arterial pressure cuff, ECG, neuromuscular blockade monitor, and end-tidal CO_2 . After placement of intravenous and monitoring devices, the patients were given oxygen, and anesthesia was induced as follows: vecuronium (0.015

mg/kg, administered intravenously) as a priming dose and sufentanil (0.6–1.2 μ g/kg) with Surital (thiamylal sodium)(2–3 mg/kg in the groups that received narcotics [A and C] and 3–5 mg/kg in the others). Vecuronium 0.06 mg/kg was given to all patients for full muscular relaxation.

Following tracheal intubation, an orogastric tube was placed and suctioned for stomach decompression. Anesthesia was then continued according to group protocol. Patients in Groups A and C received sufentanil (0.1 to 0.3 μ g/kg/hr); those in Groups B and D were given isoflurane to maintain vital signs within 20% of baseline values. Patients in Groups A and C received 0.25% and 0.5% isoflurane, respectively, whereas those in Groups B and D received generally 1.5% or more.

Postoperative Nausea \blacksquare Bloomfield and Associates

TADIES

| INCIDENCE OF NAUSEA | | | INCIDENCE OF | INCIDENCE OF VOMITING | | | |
|---------------------|--------|----------|--------------|-----------------------|--------|----------|------|
| Group | Yes | No | Р | Group | Yes | No | Р |
| A C | 5 2 | 11 13 | 0.39 | A C | 1 2 | 15 13 | 0.60 |
| A B | 5 6 | 11 10 | 1.00 | A B | 1 2 | 15 14 | 1.00 |
| B D | 6 3 | 10 13 | 0.43 | B D | 2 1 | 14 15 | 1.00 |
| C D | 23 | 13 13 | 1.00 | C D | 2 | 13 15 | 0.60 |

| TABLE 6 | | |
|-------------------------|----------------|--------------------|
| 95% CONFIDENCE INTERVAL | S FOR DIFFEREN | ICES IN INCIDENCES |
| BETWEEN GROUPS | | |

TADIEA

| | Groups | Lower | Upper |
|----------|--------|-------|-------|
| Vomiting | A-C | -28 | 14 |
| 9 | B-D | -18 | 30 |
| | A–B | -30 | 18 |
| | C-D | -14 | 28 |
| Nausea | A–C | -10 | 46 |
| | B-D | -12 | 50 |
| | A-B | -38 | 28 |
| | C-D | -39 | 27 |

At the end of surgery, the orogastric tubes were aspirated and removed; neuromuscular blockade was reversed with glycopyrrolate (0.02 mg/kg) and edrophonium (0.7 mg/kg). Extubation was performed before transfer to the recovery room. No sufentanil was given during the last 30 minutes of anesthesia.

In the recovery room, nausea and vomiting were recorded by an observer unaware of the anesthetic technique used. Chi-square tests or Fisher's exact tests, as appropriate, were used to compare the frequency of nausea and vomiting between different pairs of groups. Nausea and vomiting were measured on a two-point scale. The distribution of operative time was compared among treatment groups by means of a Kruskal-Wallis test. Analysis of variance was used to compare patient age and time from extubation to discharge from the recovery room among the groups.

There were 15 or 16 patients in each set of two groups. With a two-tailed significance level of 0.05 (and no adjustment for multiple comparisons), differences in incidence rates of approximately 45% (e.g., 10% v 55%) could be detected with a statistical power of 80%. Hence, the trial was not designed to detect differences of

 TABLE 7

 TIME FROM EXTUBATION TO RECOVERY ROOM

 DISCHARGE (MINUTES)

| Group | n | Mean | S.D. |
|-------|----|------|------|
| A | 16 | 88 | 33 |
| В | 16 | 96 | 31 |
| С | 15 | 92 | 37 |
| D | 16 | 100 | 24 |
| 1 | | | |

less than 45% in incidence rates between groups. To aid in the interpretation of differences in incidence rates, 95% confidence intervals are also provided.

RESULTS

A listing of the types of cases evaluated for this study is shown in *Table 1*. There was an equal sex distribution of patients in the four groups (*Table 2*). Mean ages, ranging from 32 to 39 years, were not significantly different (*Table 3*).

Table 4 shows the incidence of nausea in all four groups. Group A had an incidence of 31.25%; Group B, 37.5%; Group C, 13.3%; and Group D, 18.75%. The incidence of nausea varied insignificantly among all groups.

Table 5 shows the incidence of vomiting in all four groups. Group A had an incidence of 6.25%; Group B, 12.5%; Group C, 13.3%; and Group D, 6.6%. All incidences of vomiting varied insignificantly among all groups.

Table 6 shows the 95% confidence limits for all groups.

Table 7 shows the time from extubation in the operating room until discharge from the recovery room. The mean times varied from 88 to 100 minutes.

POSTOPERATIVE NAUSEA BLOOMFIELD AND ASSOCIATES

DISCUSSION

These results confirm that nausea and vomiting remain common sequels to modern anesthetic techniques. An overall incidence of 25% for nausea and 10% for vomiting represents obvious room for improvement. No difference could be shown, moreover, between methods based mainly on inhalation anesthesia with isoflurane, and intravenously administered analgesia with sufentanil. To our knowledge, no one has studied this problem in regards to sufentanil.

Alexander et al² found a significant difference in postoperative nausea and vomiting when comparing oxygen-fentanyl-nitrous oxide with oxygen-fentanylisoflurane. White et al³ found a surprisingly high incidence of nausea and vomiting when nitrous oxide was used with alfentanil or fentanyl for outpatients, raising a question whether nitrous oxide significantly increased the incidence of postoperative nausea and vomiting when associated with narcotics. The data of Alexander et al² would indicate that this is true with fentanyl. Others have attributed emetic properties to nitrous oxide, and our study tends to support this, although the differences were not statistically significant.^{2,4} However, in a large series of 780 patients, Muir et al¹ found no association between postoperative nausea and vomiting and the use of nitrous oxide in combination with enflurane or isoflurane, which would tend to argue against the emetic properties of nitrous oxide.

The emetic properties of sufentanil have not, to our knowledge, been examined in any great detail. Sufentanil, a narcotic five to 10 times more potent than fentanyl, has just as rapid an awakening time when used in equipotent dosages. When used for cardiac surgery with 100% oxygen, both fentanyl and sufentanil tend to have very low incidences of nausea and vomiting. We wondered whether the addition of nitrous oxide would make a significant difference.

Our study shows the inherent difficulty of comparing different anesthetic techniques. Supplementation of sufentanil with inhalational anesthetics was necessary in this study since suppression of awareness is not intrinsic to this drug. Also, the reduced potency of nitrous oxide at the high altitude of Denver deserves emphasis. The treatment received by Group A was planned as a strict nitrous oxide/narcotic technique. However, we have found a higher incidence of intraoperative recall with this technique because of the decreased partial pressure of nitrous oxide in Denver. Therefore, we added 0.25% isoflurane in Group A to insure total amnesia.

There are several design limitations in this study. The first is the possible cross-contamination of inhalational agents with narcotics. The combination becomes necessary to avoid postoperative respiratory depression and to insure intraoperative amnesia. The second is the small number of cases. A much larger series will be required to gain accurate data in regard to "potency equivalents" for inhalation and intravenous agents. A third factor involves postoperative evaluation. Muir et al¹ evaluated their patients in the recovery room and at a 24-hour follow-up visit. It is hard to know how long postoperatively the emetic effects of intravenous and inhalational agents last. Finally, other miscellaneous factors, such as age, gender, length of the surgery, weight, pain, medical history, and the type of surgical procedures influence postoperative nausea and vomiting.^{5,6} Very old and very young patients were excluded from our study. No statistical differences were found in any of the groups regarding sex or length of surgery. Morbidly obese patients and patients with a history of nausea and vomiting were also excluded. All surgical procedures were extra-abdominal, except for two laparoscopies.

CONCLUSION

In this pilot study, we were unable to demonstrate significant differences in postoperative nausea and vomiting with sufentanil in combinations with and without nitrous oxide and isoflurane. A larger series to enhance the lack of statistical power caused by this small sample will be needed.

REFERENCES

- 1. Alexander GD, Skupski JN, Brown EM. The role of nitrous oxide in postoperative nausea and vomiting (abst). Anesth Analg 1984; 63:175.
- Muir JJ, Warner MA, Offord KP, Buck CF, Harper JV, Kunkel SE. Role of nitrous oxide and other factors in postoperative nausea and vomiting: a randomized and blinded prospective study. Anesthesiology 1987; 66:513–518.
- 3. White PF, Coe V, Shafer A, Sung M-L: Comparison of alfentanil with fentanyl for outpatient anesthesia. Anesthesiology 1986; **64**:99–106.
- Buffington CW. Reflex actions during isoflurane anesthesia. Can Anaesth Soc J 1982; 29:S35–S43.
- 5. Palazzo MG, Strunin L. Anesthesia and emesis. I. Etiology. Can Anaesth Soc J 1984; 31:178–187.
- 6. Purkis IE: Factors that influence postoperative vomiting. Can Anaesth Soc J 1964; 11:335–353.