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output results. As the common causes of intraoperative bradycardia are hypoxia or vagal activity, such therapy should consist of ventilation with oxygen and iv atropine. To obtain a rapid response, iv atropine should be administered before marked bradycardia is present.

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Effect of Droperidol Pretreatment on Postanesthetic Vomiting in Children Undergoing Strabismus Surgery

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Vomiting is a frequent and often disturbing complication following strabismus surgery, particularly in children.1-5 Droperidol, a potent antiemetic, reduces the incidence of vomiting after many types of surgery, including strabismus surgery. 6-14 Abramowitz et al. reported that intravenous droperidol 0.075 mg·kg-1 reduces the incidence of vomiting in children undergoing strabismus repair from 85% (control group) to 43% when administered after manipulation of the eye.1 However, Meyers and Tomeldan observed that intramuscular droperidol 0.10 mg·kg-1 reduces the incidence of vomiting after eye surgery in adults to 10% when administered preoperatively and before manipulation of the eye. 15 The lower incidence of vomiting in the latter study may be attributed, in part, to the more effective antiemetic action of droperidol when administered before manipulation of the eye rather than afterward. We speculated that if intravenous droperidol were administered before manipulation of the eye, the incidence of vomiting after strabismus repair in children might be reduced to a similar extent as in adults.15 Therefore, we determined the incidence of

vomiting after strabismus repair in children who received intravenous droperidol pretreatment, or one of the routine postanesthetic analgesics used in our institution: rectal acetaminophen or intramuscular codeine.

METHODS

With approval from the Committee on Human Research, a prospective, randomized study was undertaken. Informed written consent was obtained from the parents of 100 children scheduled for elective strabismus repair. The children were ASA Physical Status 1 or 2, fasting, unpremedicated, and older than 2 yr of age. Children with a history of motion sickness or vomiting after previous strabismus surgery were not excluded from the study.

The children were randomly assigned to one of three treatment groups: intravenous droperidol (0.075 mg·kg⁻¹) (n = 31); rectal acetaminophen (10 mg·kg⁻¹) (n = 35); or intramuscular codeine (1.5 mg·kg⁻¹) (n = 34). Droperidol was given at induction of anesthesia immediately after succinylcholine. Acetaminophen and codeine were given in the postanesthetic room (PAR) when the children were arousable. Acetaminophen was administered instead of a placebo to alleviate any possible discomfort after surgery. Intramuscular codeine was included as the third treatment after it was incriminated as a possible cause of vomiting after strabismus repair by one of our ophthalmologists.§

After a precordial stethoscope, electrocardiogram, blood pressure cuff, and Doppler probe were applied, general anesthesia was induced with intravenous thiopental (5 mg·kg⁻¹), atropine (0.02 mg·kg⁻¹), and succinyl-

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choline (1.5 mg·kg⁻¹). The lungs were ventilated with 100% oxygen by mask after administration of succinylcholine. Care was taken to avoid inflation of the stomach. The trachea was then intubated and the lungs ventilated mechanically with a mixture of halothane (0.5-1.25%), nitrous oxide (70%), and oxygen (30%). Nondepolarizing muscle relaxants were not used. Intravenous fluids were administered to replace the estimated fluid deficit and the ongoing maintenance fluid requirements. Gastric contents were not aspirated. At the completion of surgery, the children were extubated when their gag reflex had returned. The children returned to the ward after recovering in the PAR. Intramuscular dimenhydrinate (1.0-1.5 mg·kg⁻¹) was administered to any child with Grade 2 or 3 vomiting (table 1). Acetaminophen (10 mg·kg⁻¹ per rectum) was administered to any child in the droperidol group who experienced pain after surgery. Clear fluids were offered to the children as soon as they were awake enough to drink. Children were discharged on the evening of surgery, providing they fulfilled three discharge criteria: 1) stable vital signs; 2) clear sensorium; and 3) tolerating fluids.

We recorded the demographic data for the three groups of children as follows: age; number of extraocular muscles repaired; duration of anesthesia; incidence of delayed extubation; and, the incidence of overnight admissions. The nurses in the PAR and on the ward recorded the time and severity (table 1) of each episode of vomiting and the time at which oral fluids were tolerated. Retching was included in the incidence of vomiting. The nurses also recorded the PAR recovery scores, ¹⁶ the duration of stay in the PAR, the post anesthetic pain scores, ¶ and the time to discharge from the hospital. The parents of each child were contacted 72–96 h after discharge from the hospital by one of the authors to determine the incidence and severity of vomiting after discharge.

Statistical significance (*P* < 0.05) was determined using the Fisher exact test, Chi-square analysis with the Yates correction, and the Mann-Whitney U-test with the Bonferroni correction for the incidence of vomiting, delayed extubations, number of muscles repaired, the PAR recovery and pain scores; and ANOVA and the Student-Neuman-Keuls test for the patients' ages, duration of anesthesia, duration of stay in the PAR, time to discharge from hospital, and the time interval from admission to PAR to tolerating fluids.

RESULTS

The mean age, number of extraocular muscles repaired, duration of anesthesia, and incidence of delayed extubation did not differ significantly among the three treatment groups (table 2). Delayed extubation (greater than 10 min after discontinuation of the anesthetic) was

Emesis Score	
Grade 1	Isolated episode(s) of vomiting in which antiemetic
	medication was not required and/or hospitalization

was not prolonged.

Grade 2 Episodes of vomiting that require antiemetic medication. This was arbitrarily defined as three episodes of vomiting/h or four consecutive h with one or more episode of vomiting/h.

Grade 3 Uncontrolled vomiting, despite antiemetic medication. Patient required hospitalization overnight.

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observed in one child who received droperidol and one who received rectal acetaminophen.

When the incidence of vomiting predischarge, post-discharge, and overall were compared for the three treatments, significantly fewer children treated with droperidol vomited predischarge and overall compared with children treated with acetaminophen and codeine (P < 0.05) (table 2). The incidence of vomiting predischarge was only 10%

TABLE 2. Summary of Demographic Data and Incidence of Vomiting in Three Treatment Groups

	Droperidol	Acetaminophen	Codeine
Number of			25
children	31	35	34
Age (yr)	6.3 ± 3.3	6.5 ± 3.2	5.9 ± 17.5
Number of muscles	2 (1-4)	2 (1-3)	2 (1-4)
repaired* Duration of anesthesia	2 (1-4)		
(min)	54.2 ± 39.0	48.9 ± 20.4	54.3 ± 3.4
Incidence of delayed extubation*	1	1	0
Incidence of overnight admissions	1 [3]	2 [6]	4 [12]
Incidence of vomiting Predischarge	3 [10]	16 [46]†	16 [47]†
Postdischarge only	2 [6]	5 [14]	7 [21]
Total	5 [16]	21 [60]†	23 [68]†

Mean ± SD.

Numbers in brackets indicate the per cent of children.

[¶] Modified pain score excluded posture: maximum 10 points. 17

^{*} Median (range)

[†] Significantly greater than droperidol (P < 0.05).

TABLE 3. Postanesthetic Scores and Times

	Droperidol	Acetaminophen	Codeine
PAR scores*			
Admit	2 (0-6)	2 (0-4)	3 (0-6)
5 min	3 (1-6)	3 (1-6)	4 (1-6)
15 min	4 (1-6)	4 (1-6)	6‡ (1-6)
30 min	6 (3-6)	5 (3-6)	6 (1-6)
Discharge	6 (3-6)	6 (5-6)	6 (4-6)
Time in PAR			1
(min)	71 ± 38.7	63 ± 26.1	65 ± 20.8
Number of patients discharged			
same day Time to discharge	30	33	30
from hospital (h)†	6.41 ± 1.6	7.14 ± 1.7	6.68 ± 2.1

Mean ± SD.

* Median (range)

† Time interval from admission to postanesthetic room (PAR) to discharge from hospital for patients discharged the same day.

 \pm Significantly different from droperidol (P < 0.05).

with droperidol compared with 46% with acetaminophen and 47% with codeine (P < 0.05) (table 2). None of the children who were given droperidol required hospitalization overnight because of vomiting, whereas two of those given acetaminophen and four of those given codeine remained overnight (Grade 3 vomiting) (table 2). One child who was given droperidol required hospitalization overnight because of drowsiness and refusal to drink. None of the children in any treatment group experienced retching without vomiting. To determine the incidence of vomiting postdischarge, 95% of the parents were contacted by telephone. The incidence of vomiting postdischarge with droperidol (6%), did not differ significantly from that with acetaminophen (14%), or codeine (21%) (table 2). None of the children who vomited after discharge from hospital required readmission for vomiting.

The median PAR recovery scores, time in the PAR, and time to discharge from the hospital (not including overnight admissions) did not differ significantly among the three treatment groups (table 3), with the exception of the PAR scores for codeine and droperidol at 15 min. The postoperative pain scores did not differ significantly among the three groups at any time. Only two children in the droperidol group required acetaminophen for postoperative pain. The time from admission to PAR to drinking fluids did not differ significantly among the three groups.

The incidence of vomiting did not differ significantly among three age groups⁷: 2-5 yr, 6-10 yr, and older than 10 yr of age, for any treatment.

DISCUSSION

Although droperidol has proven to be an effective antiemetic for many surgical procedures in children, ⁷⁻¹³ the incidence of vomiting after strabismus repair remains un-

acceptably high.¹⁻³ We found that the incidence of vomiting predischarge in children who received intravenous droperidol 0.075 mg·kg⁻¹ before manipulation of the eye is simlar to that reported by Meyers and Tomeldan,¹⁵ but significantly less than that reported by Abramowitz et al.¹ In the latter study, droperidol was given after manipulation of the eye. The reduced effectiveness of droperidol in the latter study, may be attributed in part to the administration of droperidol after the stimulus to vomit was already present. In order to reduce the incidence of vomiting after strabismus repair in children to a clinically acceptable level, intravenous droperidol should be given at induction of anesthesia (approximately 10 minutes before manipulation of the eye).¹⁸

The incidence of vomiting in the acetaminophen group is similar to that in control groups undergoing strabismus repair in some studies^{9,19} but is significantly less (85%) than that in the study by Abramowitz et al. This latter difference may be attributed to several factors. First, the administration of intravenous atropine may have reduced the incidence of vomiting in this study. 20 Second, we administered acetaminophen to the control group to provide postoperative analgesia. Although acetaminophen itself does not prevent vomiting, it may have contributed to a reduction in the incidence of vomiting indirectly by alleviating postoperative eye pain.21 Third, although the stomach was not decompressed, we avoided gastric inflation by carefully ventilating the lungs only after administration of succinylcholine and when the child was apneic. Fourth, the effect of nitrous oxide on the gas volume in the stomach was minimized by the brief duration of surgery (table 2).22

The potential side effects of droperidol may reduce its use in children undergoing elective strabismus repair. 1,3,23-26 Prolonged sedation, hypotension, restlessness, and extrapyramidal side effects have been associated with the use of droperidol in children.24 Despite the risk of prolonged sedation, 25 the recovery room scores and times to discharge from the hospital in the droperidol group did not differ significantly from the acetaminophen group (table 3). It was noted that those children who received droperidol had to be aroused more vigorously to encourage them to drink compared with children in the other two treatment groups, although there was no objective evidence of prolonged sedation in the droperidol group. Furthermore, the time interval from admission to PAR until the children were tolerating fluids did not differ significantly among the three groups. Anticholinergic drugs have been recommended to prevent many of the side effects caused by droperidol, including anxiety and restlessness, and to attenuate the oculocardiac reflex. Neither anxiety nor restlessness was observed in this study. This may be attributed, in part, to the administration of intravenous atropine coincidental with droperidol at induction of anesthesia.

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If the incidence of vomiting postdischarge had been ignored in this study, then the adjusted incidence of vomiting in the acetaminophen and codeine groups would have been 46% and 47%, respectively (the predischarge incidences, table 2). These incidences are significantly less than the incidence of vomiting in the control group (85%) in the study by Abramowitz et al., but are similar to that in the study by Hardy et al. (62.5%). 19 The high incidence of vomiting after discharge from the hospital underscores the necessity for close follow-up of all patients after surgery and discharge from the hospital in order to monitor accurately the incidence of postanesthetic vomiting. Failure to do so may result in a falsely low incidence of postoperative complications.

Although it is well known that narcotics increase the incidence of postanesthetic vomiting, 5,27,28 codeine did not significantly increase the incidence of vomiting above that for acetaminophen in this study. The incidence of vomiting predischarge, postdischarge, and in total did not differ significantly between the codeine and acetaminophen groups. The authors conclude that in comparison to acetaminophen, intramuscular codeine does not increase the incidence of vomiting after strabismus repair.

In summary we have shown that droperidol 0.075 $mg \cdot kg^{-1}$ significantly reduces the incidence of vomiting in children undergoing strabismus repair. Recovery will be comfortable and uncomplicated, and the need for narcotic analgesia will be precluded in the postoperative period. In this way, strabismus surgery will be a more positive experience for our patients.

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