PROSPECTIVE OBSERVATIONAL STUDY OF INTRA OPERATIVE AWARENESS AND RECALL DURING GENERAL ANAESTHESIA

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ABSTRACT

BACKGROUND

Awareness under anaesthesia is a rare but extremely unpleasant phenomenon. There are very few studies in the developing world and none from rural areas where incidence of intraoperative awareness may be higher due to increased patient load, limited patient knowledge and lack of trained hospital staff, reliance on older, cheaper but less effective drugs, and lack of proper equipment both for providing anaesthesia, as well as monitoring the patient.

MATERIALS AND METHODS

This hospital based cross sectional study was performed in M.G.M. Medical College and Hospital, Jamshedpur. Patients undergoing elective surgical procedures in various specialties (including general surgery, orthopedics, ear, nose and throat surgery, urology, and plastic surgery) under general anaesthesia from October 1, 2016 to September 30, 2017, i.e., over a period of 1 year were considered for this study.

RESULTS

A total of 896 patients completed the questionnaire. Postoperatively, in response to the questionnaire, seven patients reported to have remembered something under anaesthesia. Out of these, three patients described events that were confirmed by operation theatre staff to have occurred whereas they were under anaesthesia.

CONCLUSION

The incidence of awareness with recall during general anaesthesia is low. The intraoperative monitoring including clinical analysis of anaesthetized patient as well as measurement of end-tidal concentration of volatile anaesthetic agent seems to be sufficient for prevention of episodes of awareness during general anaesthesia.

KEY WORDS

General Anaesthesia, Awareness, Anaesthesia, Monitoring.
anaesthesia from October 1, 2016, to September 30, 2017, i.e., over a period of 1 year were considered for this study.

**Inclusion Criteria**

Patients aged 18 years or more, American Society of Anesthesiologist (ASA) Physical Status III or less, with normal neurological status.

Written informed consent was obtained from all patients in their own language before including them in this study.

The technique and drugs used for anaesthesia varied according to patient’s preoperative condition, surgical procedure planned, and choice of anesthesiologist; however, all patients considered for this study underwent general anaesthesia with endotracheal intubation and positive pressure ventilation. Induction agent used was either thiopentone sodium or propofol given intravenously followed by muscle relaxant which was either vecuronium or atracurium. After intubation maintenance of anaesthesia was done using 50% oxygen, 50% nitrous oxide, and volatile anesthetic (Isoflurane or sevoflurane). During the procedure and throughout the postoperative period, vital signs (including heart rate, oxygen saturation, electrocardiography, and noninvasive blood pressure) of the patients were monitored. Concentration of volatile anesthetic was adjusted according to patient’s vital signs, as well as clinical parameters such as pupillary response, sweating, and tearing. Instrumental monitoring of cerebral electrical activity and volatile anesthetic concentration was not carried out, as the same was not available in our institute. None of the patients included in this study underwent surgical procedure under mask ventilation or laryngeal mask airway insertion. Furthermore, total intravenous anaesthesia was not used in any patient. A consultant anesthesiologist (With sufficient experience of general anaesthesia) who was unaware about the patients being included in this study anesthetized all patients. Noise levels in Operation Theater were kept to a minimum, with only minor chat surgeons, anesthesiologist, and operation theater staff. After the completion of surgical procedure, anaesthesia was reversed, extubated, and shifted to postanaesthesia care unit (PACU) after adequate return of consciousness.

Approximately after 1 h of arrival in PACU, anesthesiologist (Not involved in administering anaesthesia) assessed intraoperative awareness. Anesthesiologist visited the patient and asked questions in his/her own language. First general information such as age, sex, ASA status, anaesthesia technique used, history of chronic drug intake or substance abuse, and any previous history of awareness was obtained. The second part of the questionnaire was a modified form of Brice questionnaire, used by similar studies designed to assess intraoperative awareness, in the past:

- What is the last thing you remember before going to sleep?
- What is the first thing you remember after waking up?
- Do you remember anything between going to sleep and waking up?
- Did you dream during your procedure?
- What was the worst thing about your operation?

After the questionnaire was completed, it was analyzed, and patients were categorized into either having definite awareness, possible awareness, and no awareness. If the event recalled was confirmed by attending personnel present in Operation Theater, or investigators were convinced that memory was real, patients were categorized under definite awareness. If the patient was unable to recall any event definitely indicative of awareness, but memories could have been related to intraoperative events, he/she was categorized as possible awareness. No awareness was defined as a patient with no reported awareness or if the recalled events had a high probability of occurring in immediate pre- or postoperative period. This classification is similar to one used in various other studies assessing intraoperative awareness. Patients having intraoperative dreaming were categorized separately.

Patients having awareness or possible awareness were offered to follow up in anaesthesia outpatient department.

**Statistical Analysis**

Data were collected on a Microsoft Excel® sheet and analyzed using Statistical Package for the Social Sciences® version 23 (SPSS Inc, Chicago, IL, USA) for windows. Data were expressed as ratio, percentages and proportions. Fisher’s extract test was used for comparison. P < 0.05 was considered statistically significant.

**RESULTS**

A total of 1068 adult patients underwent elective surgical procedures under general anaesthesia. Out of these, 37 patients were ASA Grade IV, so were excluded from the study, 121 patients refused to take part in the study. Of the remaining, 12 had to be shifted to Intensive Care Unit intubated and 2 were disoriented so were thus excluded from the study. There was no disruption of anaesthesia or circuit failure in any case. Hence, a total of 896 patients completed the questionnaire.

Of these 896 patients, 25.89% of patients were <30 years of age, 32.48% were between 30 and 44 years, 25.78% were between 45 and 59 years, and remaining 15.85% were 60 years and above. Nearly 32.59% of participants were males and the rest were females. As far as education level was concerned, 194 (21.65%) patients were illiterate, 258 (28.79%) patients studied up to primary standard, 189 (21.09%) patients till secondary level, 211 (23.55%) were graduates, and remaining 44 (4.91%) were postgraduates. Based on socioeconomic status, roughly half of the patients were poor (51.01%), 44.31% belonged to middle class, and 42 (4.61%) patients were rich. According to ASA grading, 62.95% of patients were ASA Grade I, 23.32% were ASA Grade II, and 13.73% were ASA Grade III. All the patients underwent elective surgical procedures under general anaesthesia with endotracheal intubation using muscle relaxants. A maximum number of patients participating in the present study underwent general surgical procedures (77.01%), followed by ear, nose, and throat surgery (11.38%), orthopedic surgery (5.13%), urology surgery (4.24%), and plastic surgery (2.23%). Among the patients undergoing general surgical procedures, 439 patients (65.72%) underwent laparoscopic cholecystectomy, 58 (8.68%) underwent laparoscopic hernioplasty, 14 (2.09%)
laparoscopic appendectomy, 25 (3.74%) open cholecystectomy, 30 (4.49%) modified radical mastectomy, 16 (2.39%) laparotomy, and remaining 86 patients (12.87%) had various other elective surgical procedures.

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Demographic Variables</th>
<th>N (%)</th>
<th>Awareness N (%)</th>
<th>Dreaming N (%)</th>
<th>No Awareness N (%)</th>
<th>P Value</th>
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<td>&lt;30</td>
<td>237 (26.32)</td>
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<td>30-44</td>
<td>286 (31.20)</td>
<td>2 (0.33)</td>
<td>64 (7.5)</td>
<td>220 (24.27)</td>
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<td></td>
<td>45-59</td>
<td>226(27.93)</td>
<td>1 (0.112)</td>
<td>40 (4.66)</td>
<td>190 (20.25)</td>
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Table 1. Demographic Variables of Patients Included in the Study

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<th>Attempted Awareness N (%)</th>
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DISCUSSION
Incidence of awareness under anaesthesia with postoperative recall was found to be 0.33% in our study. However, if patients with possible awareness were also included, incidence of awareness under anaesthesia would become 0.78%. This occurrence is higher than reported by most studies conducted in the Western world which estimated the incidence to be between 0.1% and 0.2%. However, in a study by Errando et al done on over 4000 patients, incidence of awareness with recall was found to be 1.0% among all patients operated under anaesthesia and 0.8% after removing high-risk cases.

These awareness cases were followed retrospectively to study the intraoperative events and vitals, anaesthetic technique and preoperative status, including demographic data. We could not find any significant predictor of possible awareness in these cases of awareness. All the cases were monitored intraoperatively by using entropy from induction to emergence at fixed intervals of time and in these 2 cases of awareness entropy was found consistently above 60 but there was no association between increased entropy readings and the incidence of awareness as many patients had entropy above 60 but did not report awareness in our study.

Awareness is caused by the administration of general anaesthesia that is inadequate to maintain unconsciousness and to prevent recall during surgical stimulation. Common causes include large anaesthetic requirements, equipment misuse or failure and smaller doses of anaesthetic drugs.

Various studies found an increased risk of awareness with sicker patients undergoing major surgery, this finding may reflect the use of smaller anaesthetic doses and light anaesthetic techniques in sicker patients.

In many cases, awareness during anaesthesia is a potentially avoidable adverse anaesthetic outcome. In light of follow up studies suggesting that such victims of awareness may exhibit significant psychological after effects such as PTSD, attempts to further reduce its incidence are warranted.

Because awareness occurred despite the usual clinical monitoring of anaesthetic depth (BP, HR and end tidal anaesthetic monitoring), a monitor of cerebral function and depth of anaesthesia may be of theoretical benefit.

The limitation of our study is that it did not assess the long-term psychological sequelae of intraoperative awareness and recall among the victims.

CONCLUSION
The incidence of awareness with recall during general anaesthesia is low. Intraoperative monitoring including clinical analysis of anaesthetized patient as well as measurement of end-tidal concentration of volatile anaesthetic agent seems to be sufficient for prevention of episodes of awareness during general anaesthesia.

REFERENCES


