

LETTER TO THE EDITOR

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A patient with intraoperative awareness history requiring high propofol effect-site concentrations for general anesthesia

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To the Editor,

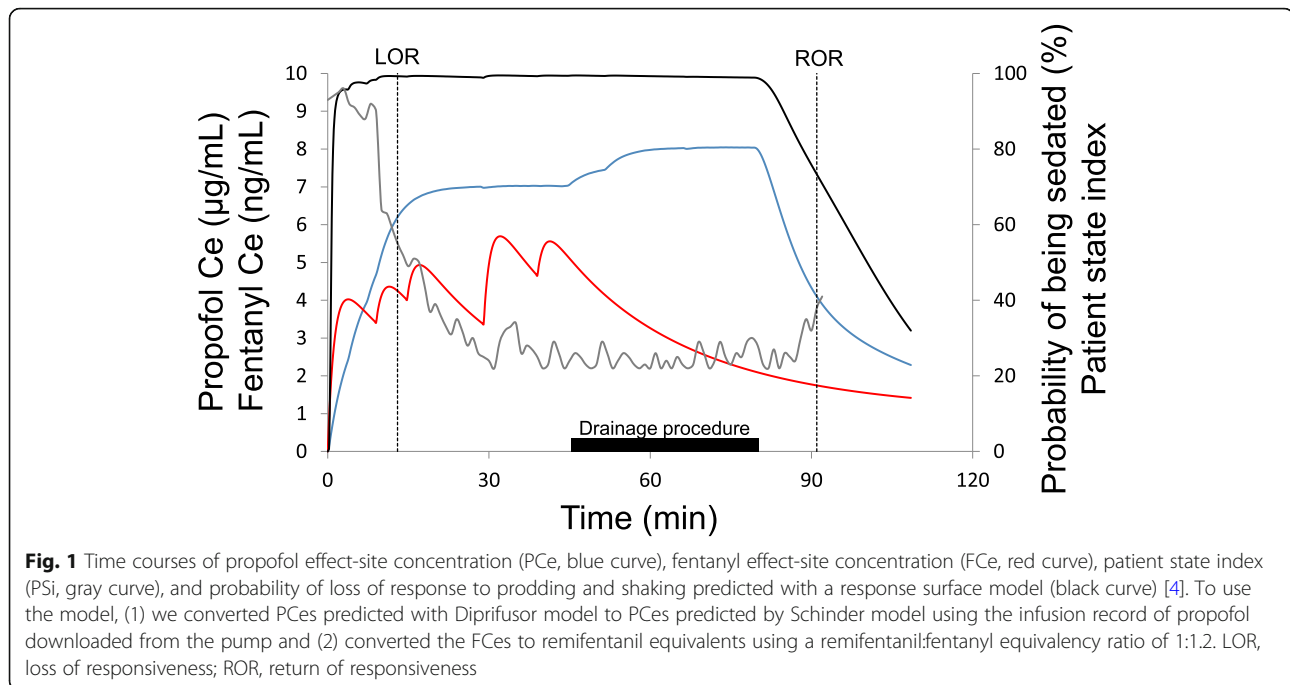
Intraoperative awareness (IA) occurs in up to 0.12% of patients under general anesthesia [1]. Herein, we describe the case of patient with IA history requiring high propofol effect-site concentrations (PCe) for anesthesia. We obtained a written informed consent from the patient for publication. The patient was a 34-year-old woman (158 cm, 46 kg) who underwent a semi-emergency transcutaneous drainage of an intrapelvic tumor under computed tomography (CT) guidance. She was being administered celecoxib, loxoprofen, and pregabalin for chronic pain. She had undergone multiple operations for neurilemmomas, had experienced IA with unclear details, and desired a complete loss of consciousness for her upcoming operation. In the CT room, we induced anesthesia with fentanyl (150 µg) and propofol via target-controlled infusion (TCI, Diprifusor model, TE-371, Terumo, Tokyo, Japan). We initiated the target plasma concentration of propofol at 4 µg/mL and increased it gradually (Fig. 1). When the patient's responses to voice, mild prodding, and shaking were lost, the PCe was 6.4 µg/mL and the patient state index (PSi) derived with Sed-Line® (Masimo, USA) was 79 (PSi 25–50 is recommended during anesthesia). At this stage, fast waves in electroencephalogram were still observed but were not predominant. Spectral edge frequency (SEF), which can be used as a surrogate measure to observe the change in the amount of fast waves, was 12.9 Hz. Following the tracheal intubation using rocuronium (40 mg), we moved the patient to a prone position

and installed a mechanical ventilator. We titrated target propofol concentrations in accordance with PSi values and with the patient's vital signs. The drainage was uneventfully completed. The maximum PCe was 8.0 µg/mL and the total fentanyl dose was 400 µg. Ten minutes after PCe reached 8.0 µg/mL, PSi and SEF were 22 and 10.8 Hz, respectively. The anesthesia duration was 100 min. The patient regained responsiveness to voice when the PCe decreased to 4.0 µg/mL. We did not observe any postoperative complications and the patient was satisfied with the clinical course without any IA. Although no consensus exists on IA, using inhalation anesthetics may produce less IA [2, 3]. However, the CT room was not equipped with the excess anesthesia gas extraction system and this prevented us from using inhaled anesthetics with an anesthesia machine, which is why we chose total intravenous anesthesia. Our patient required much higher PCes than usual (TCIs are 3.0–6.0 µg/mL and 2.0–5.0 µg/mL for anesthesia induction and maintenance, respectively; according to the propofol package insert; Diprivan®, Aspen Japan, Tokyo). The patient was apparently awake, although the probability of loss of response to prodding and shaking right before the patient lost responsiveness was predicted at 99.3% with the response surface model [4]. Therefore, we think she was relatively insensitive to propofol due to pharmacokinetic and/or pharmacodynamic variability, and this may have explained her previous IA. Although pregabalin reportedly reduces propofol requirements [5], our patient required a large propofol dose. Our case highlights the need to

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be aware of potential propofol insensitivities, especially in patients with IA history.

Abbreviations

CT: Computed tomography; FCE: Fentanyl effect-site concentration; IA: Intraoperative awareness; PCE: Propofol effect-site concentration; PSi: Patient state index; TCI: Target-controlled infusion

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None

Authors' contributions

SO treated the patient and wrote the manuscript. YN and RO treated the patient and revised the manuscript. TH treated the patient. CH and MM helped to design the case report. All authors reviewed and approved the final draft.

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Availability of data and materials

Not applicable

Ethics approval and consent to participate

In our institution, IRB approval is not required for a case report.

Consent for publication

Written informed consent was obtained from the patient for publication of this case report.

Competing interests

The authors declare that they have no competing interests.

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