

Awake craniotomy anesthetic management using dexmedetomidine, propofol, and remifentanyl

Andrea Prontera¹
Stefano Baroni²
Andrea Marudi²
Franco Valzania³
Alberto Feletti¹
Francesca Benuzzi⁴
Elisabetta Bertellini²
Giacomo Pavesi¹

¹Department of Neurosurgery, Nuovo Ospedale Civile SAgostino-Estense,

²Department of Anesthesiology, Nuovo Ospedale Civile SAgostino-Estense, ³Department of Neurology, Nuovo Ospedale Civile S Agostino-Estense, ⁴Department of Neuroscience, University of Modena and Reggio Emilia, Modena, Italy

Introduction: Awake craniotomy allows continuous monitoring of patients' neurological functions during open surgery. Anesthesiologists have to sedate patients in a way so that they are compliant throughout the whole surgical procedure, nevertheless maintaining adequate analgesia and anxiolysis. Currently, the use of α_2 -receptor agonist dexmedetomidine as the primary hypnotic–sedative medication is increasing.

Methods: Nine patients undergoing awake craniotomy were treated with refined monitored anesthesia care (MAC) protocol consisting of a combination of local anesthesia without scalp block, low-dose infusion of dexmedetomidine, propofol, and remifentanyl, without the need of airways management.

Results: The anesthetic protocol applied in our study has the advantage of decreasing the dose of each drug and thus reducing the occurrence of side effects. All patients had smooth and rapid awakenings. The brain remained relaxed during the entire procedure.

Conclusion: In our experience, this protocol is safe and effective during awake brain surgery. Nevertheless, prospective randomized trials are necessary to confirm the optimal anesthetic technique to be used.

Keywords: dexmedetomidine, awake surgery, anesthesia

Abbreviations

bpm, beats per minute; DEX, dexmedetomidine; EcoG, electrocorticography; HR, heart rate; LMA, laryngeal mask airway; MAC, monitored anesthesia care; MAP, mean arterial pressure; MEPs, motor evoked potentials; PRO, propofol; REM, remifentanyl; SEPs, somatosensory evoked potentials.

Introduction

Awake craniotomy enables neurosurgeons to continuously monitor the patient's neurological functions during open surgery.

Anesthesiologists are required to maintain the patient's cooperation along with adequate sedation, analgesia, and anxiolysis throughout the whole procedure. Although awake craniotomy procedures have become common in neurosurgical practice, there is still no consensus on the optimal pharmacological regimen during the intervention.

The use of α_2 -receptor agonist DEX as primary hypnotic–sedative medication is common nowadays, especially in intensive care unit settings, because it provides effective and safe conscious sedation.

DEX rapidly acts on the pontine locus coeruleus, providing subcortical sedation characterized by a comfortable and quick arousal, with minimal effect on the respiratory drive. Bradycardia and moderate hypotension are the most common hemodynamic effects.

Correspondence: Alberto Feletti
Department of Neurosurgery, Nuovo Ospedale Civile S Agostino-Estense
Modena Hospital, via Pietro Giardini,
1355, Modena 41126, Italy
Tel +39 059 396 1471
Fax +39 059 396 2408
Email alberto.feletti@gmail.com

We describe the anesthetic management of patients who underwent awake craniotomies at our institution (NOCSAE Modena Hospital, Modena, Italy). A combination of DEX, PRO, and REM was used without any need for airway instrumentation.

Methods

Population

We retrospectively reviewed all awake craniotomies performed with MAC protocol at our institution from May 2013 to December 2014. We identified 9 patients (5 males and 4 females) meeting the inclusion criteria. They underwent craniotomy to remove 7 gliomas, 1 meningioma, and 1 cavernoma. Unfavorable anatomical features such as short neck or overweight were not considered exclusion criteria. Data regarding anesthetic technique, vital parameters, and patients' outcome were collected.

The most involved locations were the middle frontal gyrus, the middle temporal gyrus, the precentral gyrus, the inferior frontal gyrus, the angular gyrus, and the superior temporal gyrus.

Patients' demographics and pathological characteristics are shown in Table 1.

A neuropsychologist instructed the patients to perform the neurological examination during surgery.

The official institutional board (Modena Local Health Authority – Internal Review Board) approved the study. Patient consent was not required by the institutional board to review medical records.

Anesthetic and surgical procedures

All patients underwent awake craniotomy using the MAC protocol.

Ranitidine 50 mg and metoclopramide 10 mg were administered 2 h before induction of anesthesia to prevent nausea and vomiting.

After electrocardiography, pulse oximetry, noninvasive blood pressure monitoring, and double peripheral venous

cannulation, we started the simultaneous infusion of DEX, PRO, and REM.

Radial artery-line, Foley catheter, and an additional peripheral intravenous catheter were placed after induction of anesthesia.

Initial DEX dose of 0.7 µg/kg/h was followed by 0.2–1.4 µg/kg/h maintenance infusion. No loading dose was used, and infusion rate was progressively increased until adequate sedation was obtained. We used Ramsay sedation score to assess the level of sedation every 90 s. A Ramsey sedation score between 4 and 6 was considered adequate. It was reached by slowly increasing the DEX infusion to 0.2 µg/kg, unless the HR decreased to 50 bpm or the MAP decreased to 60 mmHg. In case of persistently inadequate sedation despite these adjustments, we increased the concentration of the PRO infusion.

All patients had spontaneous ventilation with supplemental oxygen (2 L/min) through a capnomask (Mediplus, Bucks, UK). No laryngeal mask or other devices were used.

A 3-pin head-holder was placed on the patient under local anesthesia with subcutaneous injection of bupivacaine with epinephrine (1:200,000) and ropivacaine 7.5 mg/mL, without nerve block.

Comfortable positioning of the patient was achieved, keeping the head slightly elevated in order not to compress jugular venous flow. Such positioning prevents airway occlusion during the asleep period, especially in patients with anatomically unfavorable neck.

Patients underwent EcoG using a 32-channel Nicolet alliance portable unit (Nicolet BMSI, Inc, Madison, WI, USA). Standard MEPs and SEPs were recorded.

Surgical incision was planned using neuronavigation in order to expose both the tumor and the surrounding cortical eloquent areas. A solution of bupivacaine with epinephrine (1:200,000) and ropivacaine 7.5 mg/mL was used for local anesthesia.

Draping was adjusted to keep eyes and airways easily accessible.

Table 1 Patient demographics and clinicopathological characteristics

Number	Age, years	Sex	Previous history	Pathology	Grade	Location (gyrus)
1	48	M	Glaucoma	Glioma	IV	Posterior T2–T3
2	57	F	Hypertension	Meningioma	I	Precentral–F3
3	56	F	Hypertension, obesity, diabetes mellitus type 2	Glioma	III	Posterior F1
4	62	F	Dyslipidemia	Glioma	II	F2
5	30	M	None	Glioma	II	Precentral
6	31	F	None	Cavernoma		F2
7	41	M	None	Glioma	II	T2–angular

Abbreviations: F, female; M, male.

The patient was maintained asleep during scalp incision, bone flap removal, and dural opening. Dura mater was soaked with lidocaine 10% before incision.

After brain exposure, all drugs were stopped, and the patient progressively awakened. The surgeon gave ongoing feedback to the anesthesiologist regarding the brain appearance, in order to maintain the brain relaxed.

Cortical mapping was achieved using Ojemann cortical stimulator (Radionics Corp, Burlington, MA, USA) with a monopolar probe, delivering stimuli with a single 1 ms pulse with a 60 Hz frequency. Skill tests were performed during surgical resection.

Analgesics (acetaminophen 1 g) were added when needed.

When resection adjacent to functionally relevant areas was completed, anesthesia was reinduced with DEX, PRO, and REM infusions titrated to patient sedation level.

Patients spontaneously breathed throughout the whole procedure without the aid of a laryngeal mask.

After wound closure, all anesthetics were stopped, allowing the patient to awaken from anesthesia in the operative theater. All patients were admitted to the subintensive care unit.

Results

Table 2 summarizes the drug infusion doses, HR, arterial blood pressure, MAP, awakening periods, and length of anesthesia.

From induction to awakening, DEX, PRO, and REM infusion doses ranged between 0.28 and 0.90 $\mu\text{g}/\text{kg}/\text{h}$, 0.51 and 1.26 $\text{mg}/\text{kg}/\text{h}$, and 0.02 and 0.05 $\mu\text{g}/\text{kg}/\text{min}$, respectively.

We observed HR and MAP decrease as expected; however, their values were never less than 50 bpm and 60 mmHg, respectively.

EcoG confirmed appropriate patient sedation.

After stopping drug infusion, the patients were able to collaborate for an average period of 10 min. The awakening was always smooth, resembling natural awakening. Moreover, as we avoided the use of a laryngeal mask and other airway devices, we did not observe any coughing.

The awakening period lasted on average 77 min (range: 40–120 min). Three patients required additional analgesia (acetaminophen 1 g).

Drugs infusion doses during the final part of the operation ranged between 0.23 and 0.91 $\mu\text{g}/\text{kg}/\text{h}$ for DEX, 0.2 and 1.45 $\text{mg}/\text{kg}/\text{h}$ for PRO, and 0.04 and 0.08 $\mu\text{g}/\text{kg}/\text{min}$ for REM. Similar to the previous asleep period, we did not observe critical levels of HR and MAP.

After surgery, all patients reached an Aldrete score of 10 after a mean period of 11 min (range: 4–22 min).

No significant respiratory depression was noticed. Intra-operative arterial blood gas analysis confirmed normoxia and normocapnia.

Discussion

In the past few years, awake craniotomy has allowed wider and safer surgical resections of lesions located in the vicinity of functionally relevant areas.

The possibility to easily wake the patients up and neurologically monitor them during surgery is the main advantage of this procedure.

However, there is still no consensus or any established protocol for the best anesthetic management of such peri-operative sedation.¹

Recently, DEX, a selective alpha-2 receptor agonist, has been proposed as the primary sedative–hypnotic drug, as it does not depress respiratory drive and it does not interfere with electrophysiological testing. Its sedative effects are mediated through decreased firing of the locus coeruleus, the main noradrenergic nucleus located on the floor of the fourth ventricle, and subsequent decrease in noradrenaline release in the sympathetic nerves. Such characteristics encourage the use of DEX during awake neurosurgical procedures.

However, DEX has well-known adverse effects, such as bradycardia and hypotension. These cardiovascular effects are dose dependent: with lower infusion rates, the central effect predominates, leading to a decrease in HR and blood pressure. With higher doses, peripheral vasoconstriction prevails, leading to an increase in systemic vascular resistance and blood pressure, while the bradycardic effect is further emphasized. Cardiac arrest and respiratory depression have also been reported.^{2,3} Several studies^{4,5} have extensively shown the role of DEX during functional neurosurgery. We reviewed the pertinent literature about the use of DEX during awake surgery in functionally relevant brain areas.

de Almeida et al⁶ operated on 3 epileptic patients with cavernous angiomas in the speech area using DEX as the primary hypnotic drug, without the need to protect airway patency. They administered boluses of fentanyl, midazolam, PRO, and droperidol during the procedure when judged necessary. In their series, they did not report either significant hemodynamic instability or respiratory depression.

Shen et al⁷ prospectively compared DEX and PRO as primary hypnotic drugs during awake craniotomy, performed with asleep–awake–asleep protocol. They concluded that both DEX and PRO can be effectively and safely administered for conscious sedation. However, they found that DEX has a shorter arousal time.

Table 2 Table summarizing drug infusion doses during asleep, awake, and second asleep periods*

S no	Preinduction			Asleep			Awake			Asleep			Postsurgery						
	HR, bpm range (mean)	AP, mmHg (MAP)	DEX, PRO, REM, µg/kg/h	AP (MAP)	HR range (mean)	Time, min	Drugs	Length, min	DEX, PRO, REM, µg/kg/h (mean)	AP (MAP)	HR range (mean)	Time, min	Time, min	Length of awakening anesthesia, min					
1	67–76 (70)	140/80 (100)	0.47	0.51	0.05	90/50 (63)	60–66 (62)	12	75–84 (82)	140/90 (107)	–	120	0.37	0.20	0.04	110/70 (83)	65–42 (61)	18	410
2	70–92 (82)	140/80 (100)	0.73	1.18	0.05	100/60 (73)	55–62 (57)	8	65–70 (67)	140/90 (107)	–	50	0.91	1.45	0.08	80/60 (67)	65–60 (62)	8	360
3	90–111 (104)	160/90 (113)	0.48	0.6	0.06	140/80 (100)	67–77 (71)	14	100–125 (117)	160/90 (113)	–	60	0.81	0.37	0.04	140/80 (100)	80–90 (87)	22	380
4	62–81 (78)	115/55 (75)	0.29	1.07	0.05	80/50 (60)	47–60 (56)	11	67–75 (70)	120/80 (93)	–	90	0.29	0.95	0.05	110/80 (90)	60–70 (64)	10	400
5	70–89 (80)	140/80 (100)	0.28	0.76	0.02	110/60 (76)	48–68 (60)	8	62–70 (64)	130/70 (90)	Paracetamol	40	0.23	0.58	0.04	100/60 (73)	60–70 (66)	8	340
6	66–87 (71)	130/80 (96)	0.86	1	0.04	110/60 (76)	55–68 (62)	9	60–74 (64)	120/80 (93)	–	70	0.62	0.84	0.05	110/60 (77)	60–65 (61)	12	360
7	65–78 (74)	140/70 (93)	0.7	0.56	0.04	90/45 (60)	62–71 (65)	11	75–87 (84)	165/95 (118)	–	90	0.5	0.37	0.05	130/60 (83)	70–75 (72)	10	420
8	65–70 (67)	120/80 (93)	0.67	1.26	0.05	90/60 (70)	55–67 (59)	4	75–93 (80)	140/80 (100)	Paracetamol	60	0.58	0.84	0.04	135/70 (92)	60–75 (65)	7	365
9	72–90 (80)	160/85 (110)	0.90	1.5	0.05	110/60 (77)	50–65 (55)	11	75–90 (80)	145/80 (101)	Paracetamol	114	0.65	1	0.03	130/65 (87)	42–50 (45)	4	390

Notes: *HR, AP, and MAP during preinduction, asleep, awake, and second asleep periods; awakening periods, and total length of anesthesia are presented. **Abbreviations:** AP, arterial pressure; DEX, dexmedetomidine; HR, heart rate; MAP, mean arterial pressure; PRO, propofol; REM, remifentanyl.

Mohd Nazaruddin et al⁸ successfully treated 4 patients with a combination of DEX and REM intravenous infusions and scalp block with the MAC protocol. They delivered oxygen to patients only through nasal prongs. Only 1 patient required supplementary infusion of PRO to achieve adequate sedation.

Kallapur and Bhosale⁹ reported a case of tumor removal with awake craniotomy, using scalp block without 3-pin holder, with simultaneous infusion of DEX and PRO and oxygen delivery through a simple mask. They concluded that DEX is a useful adjuvant during awake craniotomy for tumor resection.

Chung et al¹⁰ performed 6 different anesthetic managements during awake tumor removal in 6 different patients. They simultaneously infused DEX, PRO, and REM in 2 patients. In 1 case, without airway protection, they did not obtain adequate sedation despite a high dose of sedatives. In the second patient, they reported good sedation without perioperative events, although with the aid of LMA.

Mack et al¹¹ presented a retrospective review of 10 patients who received DEX for awake craniotomy. Five of them were managed with MAC protocol. They concluded that DEX appears to be a useful sedative for awake craniotomy.

In our study, we applied a refined MAC protocol to perform awake craniotomy. We obtained adequate sedation and analgesia with the simultaneous infusion of DEX, PRO, and REM without any loading dose. Using a combination of drugs rather than a single drug presents several advantages. First, this regimen allows the use of lower PRO doses, thus reducing the risk of respiratory depression. Second, DEX doses can also be reduced, thus decreasing the risk of cardiovascular side effects. Third, this analgesedative regimen allows spontaneous airway patency without the need for tracheal intubation, LMA, or nasal airway, therefore reducing the risk of cough during the awake period. All patients were able to breathe spontaneously.

We did not perform scalp block, despite the use of a 3-pin head holder. We provided proper local anesthesia by injecting only the 3 sites around the pins and the skin incision. Actually, avoiding scalp block can reduce the risk of eventual transient bacteremia due to multiple needle-stick injuries, at the same time ensuring a comfortable surgical experience to the patient.

After stopping the infusions, all patients experienced smooth and rapid awakening. The brain remained relaxed throughout the entire procedure. Similarly, we also observed stability of hemodynamic parameters during the period of neurological monitoring.

Only 3 patients complained of mild pain, successfully treated with acetaminophen. Two patients experienced dry mouth.

We also observed that a small increase of PRO dose during simultaneous infusion of DEX leads to a higher-than-expected increase of respiratory frequency.

The main limitations of our study are the limited number of included patients, as well as the retrospective design of the study.

Conclusion

Awake craniotomy entails specific anesthetic management. DEX is a unique sedative agent that does not directly cause respiratory depression. At the moment, the optimal dose regimen of DEX during awake craniotomy is unknown. Within the limits of a retrospective study conducted on a limited number of patients, we have refined a MAC protocol combining local anesthesia without scalp block and low-dose infusion of DEX, PRO, and REM, with no need for airway support. This protocol proved safe and effective for this surgery. Nevertheless, prospective randomized trials are necessary to confirm the optimal anesthetic technique in this setting.

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Disclosure

The authors report no conflicts of interest in this work.

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